Jla wbqšY KwgwUi 23 Rby 2016 Zwi‡L Abyô‡Z 245 Zg mfvi KvhweeiYx

¯'v¯'¨ I cwievi Kj¨vY gšYvj‡qi mwPe Rbve %mq` gbRyij Bmjvg Gi mfvcwZ‡Z Jla wbqšY KwgwUi 245 Zg mfv weMZ 23 Rby 2016 ZwiL ‡ejv 12:00 NwUKvq gšYvj‡qi mfv K‡¶ Abyô‡Z nq|

mfvq KwgwUi নিম্নবর্ণিত সদস্যগণ উপস্থিত ছিলেন (জ্যেষ্ঠতার ক্রমানুসারে নয়) ঃ

1| Rbve mKgvi iÄb †Nvl, cwZwbwa, evsjv‡`k dvgwmDwUK¨vjm B‡¤úvUvm G‡mvwm‡qkb, XvKv|
2| Aa¨vcK †gvt kvidzwÏb Avn‡¤§`, cwZwbwa, e½eÜz kL gyRe †gwW‡Kj wekwe`¨vjq, XvKv|
3| ‡gRi †Rbv‡ij †gvt Ave`yi Avjx wgqv, Kbmvj‡U>U wdwRwkqvb ‡Rbv‡ij, evsjv‡`k AvgW †dv‡mm †gwW‡Kj mwf©mm|
4| Wvt gÄyi Avn‡g`, mnKvix cwiPvjK, nvmcvZvj-2, c‡ÿ gnvcwiPvjK, ¯'v¯'¨ Awa`ßi, gnvLvjx, XvKv|
5| Wvt ‡gvt Avãyi nwjg, Pxd †f‡Uwibvix Awdmvi, †m›Uvj †f‡Uwibvix nmwcUvj, c‡¶ gnvcwiPvjK, cwY m¤ú` Awa`ßi, XvKv|
6| cvifxb AvKZvi, hyM¥ mwPe, Rb¯'v¯'¨, ¯'v¯'¨ I cwievi Kj¨vY gšYvjq|
7| Aa¨vcK Wvt †gvt BmgvBj Lvb, Wxb wPwKrmv Abyl`, XvKv wekwe`¨vjq I cwZwbwa, dvgv‡KvjwR wefvM, XvKv †gwW‡Kj K‡jR|
8| Aa¨vcK ‡gvt mvBdzj Bmjvg, Wxb, dv‡gmx Abyl`, XvKv wekwe`¨vjq|
9| Aa¨vcK dwi`v †eMg, wKwbK¨vj dv‡gmx I dvgv‡KvjwR wefvM, XvKv wekwe`¨vjq|
10| Aa¨vcK ‡gvt AwRRyi Kvnnvi, †gwWwmb wefvM, XvKv †gwW‡Kj K‡jR|
11| Aa¨vcK Wvt RwKi †nvmvBb Mwje, Pg I †hŠb †ivM we‡klÁ, m¨vi mwjgyjøvn †gwW‡Kj K‡jR, XvKv|
12| ডাঃ মালা বনিক, সহকারী অধ্যাপক, গাইনী ও অবস্টেট্রিক্স বিভাগ, স্যার সলিমুলণ্ডাহ্ মেডিকেল কলেজ, ঢাকা।
13| Rbve Ave`yi †gv³vw`i, mv‡eK gnvmwPe, evsjv‡`k Jla wkí mwgwZ Ges e¨e¯'vcbv cwiPvjK, Bb‡mÞv dvgv wjt|
14| Rbve Gg †gvQv‡ÏK †nv‡mb, cwZwbwa, evsjv‡`k dv‡gmx KvDwÝj, XvKv|
15| nvKxg †gvt BDmyd nvi“b fBqv, BDbvbx we‡klÁ, evsjv‡`k BDbvbx I Avqy‡e©w`K †evW©|
16| ‡gRi †Rbv‡ij †gvt ‡gv¯—wdRyi ingvb, gnvcwiPvjK, Jla ckvmb Awa`ßi, XvKv|

mfvq Av‡jvP¨ welq mgn-wbgœi|c t

1| weMZ 25-05-2016 Zwi‡L Abyô‡Z Jla wbqšY KwgwUi ‡UKwbK¨vj mve-KwgwUi mfvq Av‡jwPZ I mycwikKZ.wb¤œewYZ wel‡q Av‡jvPbv I wm×vš MnY t

K| ¯'vbxqfv‡e Drcv`‡bi Rb¨ wnDg¨vb Jl‡ai †iwR‡÷k‡bi wbwg‡Ë `wLjKZ.335 wU Jl‡ai g‡a¨ †UKwbK¨vj mve-KwgwU KZK.Abyjgv`‡bi Rb¨ mycwikKZ.174wU Ges Av‡e`b bv-gÄyii mycwikKZ.161 wU Av‡e`b Gi Dci gZvgZ I wm×vš MnY cm‡½|

L| Avg`vbxi Rb¨ wnDg¨vb Jl‡ai †iwR‡÷k‡bi wbwg‡Ë `wLjKZ.43wU Jl‡ai g‡a¨ †UKwbK¨vj mve-KwgwU KZK.Abyjgv`‡bi Rb¨ mycwikKZ.31 wU Ges Av‡e`b bv-gÄyii mycwikKZ.12 wU Av‡e`b Gi Dci gZvgZ I wm×vš MnY cm‡½|

M| ¯'vbxq Drcv`‡bi Rb¨ Av‡ew`Z ‡f‡Uwibvix Jl‡ai †iwR‡÷k‡bi wbwg‡Ë Av‡ew`Z 27wU Jl‡ai g‡a¨ †UKwbK¨vj mve-KwgwU KZK.Abyjgv`‡bi Rb¨ mycwikKZ.21 wU Ges ‡idv‡iÝmn ‡UKwbK¨vj mve-KwgwUi mfvq cieZx‡Z Dc¯'vc‡bi Rb¨ mycwikKZ.6 wU Av‡e`b Gi Dci gZvgZ I wm×vš MnY cm‡½|

N| Avg`vbxi Rb¨ ‡f‡U wibvix Jl‡ai †iwR‡÷k‡bi wbwg‡Ë Av‡e`Z 64wU Jl‡ai g‡a¨ †UKwbK¨vj mve-KwgwU KZK Abyôgv`‡bi Rb¨ mycwikKZ.61 wU Ges Av‡e`b bv-gÄyi mycwikKZ.2 wU Ges ‡idv‡iÝmn ‡UKwbK¨vj mve-KwgwUi mfvq cieZx‡Z Dc¯vc‡bi Rb¨ mycwikKZ.1 wU Av‡e`b Gi Dci gZvgZ I wm×vš MnY cm‡½|

O| gvbyl i †`‡n e¨enwh Avg`vbxKZ.Jl‡ai †iwR‡÷kb c`v‡bi †¶‡Î DbZ 07wU †`‡ki mv‡_ EMA-Gi wd †mj mwUwd‡KU Mn‡Yi wel‡q Av‡jvPbv I wm×vš MnY|

P| ‡iwR‡÷kb mvgwqK ewZjKZ.45wU Jl‡ai †iwR‡÷kb ewZ‡ji wel‡q Av‡jvPbv I wm×vš MnY|

Q| Flupentixol + Melitracen Combination RvZxq c`wUi ‡iwR‡÷kb ewZ‡ji wel‡q ADRAC KwgwUi PZz mfvi mycwik Abyôvqx Jla wbqšY KwgwUi †UKwbK¨vj mve-KwgwUi mycwik m¤ú‡K Av‡jvPbv I wm×vš MnY|

2| MZ 25 †g 2016 Ges 15 Ryb 2016 Zwi‡L Abyôv‡Z ‡gwW‡Kj wWfvBs মূল্যায়ণ সংক্রান্ত ‡UKwbK¨vj mve-KwgwUi mycwik Abyôvqx ¯vbxq Drcv`‡bi Rb¨ 2 wU Ges Avg`vbxi Rb¨ 257 wU †gwW‡Kj wWfvB‡mm Gi †iwR‡÷k‡bi wbwg‡Ë `wLjKZ.Av‡e`b Gi Dci gZvgZ c`vb cm‡½|

3| MZ 13/06/2016 Zwi‡L Abyôv‡Z nvevj Jla GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwU)-Gi mfvi mycwikKZ.129 wU nvevj Jl‡ai g‡a¨ Abyôgv`‡bi mycwikKZ.97 wU Ges bv-gÄyi mycwikKZ.32 wU nvevj Jl‡ai †iwR‡÷kb c`vb m¤ú‡K Av‡jvPbv I wm×vš MnY cm‡½|

<u>mfvi Av‡jvPbv I wm×vš t</u>

সভাপতি মহোদয় উপস্থিত সকলকে স্বাগত জানিয়ে সভার কার্যক্রম শুরু করেন। আলোচ্যসূচী অনুযায়ী বিষয়সমূহ Dc¯vcb Kiv nq|

1| wecMZ 25-05-2016 Zwi‡L Abyôv‡Z Jla wbqšY KwgwUi ‡UKwbK¨vj mve-KwgwUi mfvq Av‡jvPZ I mycwikKZ.wb¤ewYZ wel‡q Av‡jvPbv I wm×vš MnY t

K| ¯vbxqfv‡e Drcv`‡bi Rb¨ wnDg¨vb Jl‡ai †iwR‡÷k‡bi wbwg‡Ë `wLjKZ.335 wU c` ‡UKwbK¨vj mve-KwgwUi mfvq Dc¯vcb Kiv n‡j m`m¨MY Jla¸wji Safety, Efficacy and Usefulness we‡ePbv K‡i D³ Jla¸‡jvi g‡a¨ 174wU Jl‡ai Abyôgv`‡bi mycwik Ges Awkó 161 wU Jla bvgÄyi Kiv mycwik K‡ib (Annex-A)|

wm×vš t mfvq me-সম্মতিক্রমে টেকনিক্যাল সব-KwgwUi mycwik Abyôgv`b Kiv nq|

L| Avg`vbxi Rb¨ wnDg¨vb Jl‡ai †iwR‡÷k‡bi wbwg‡Ë `wLjKZ.43wU Jla mfvq Dc¯vcb Kiv n‡j ‡UKwbK¨vj mve-KwgwUi mদস্যগণ বিস্তারিত আলোচনাক্রমে উ³ Jlamg‡ni g‡a¨ 31wU Jl‡ai Abyôgv`‡bi mycwik Ges 12wU Jl‡ai †iwR‡÷k‡bi Av‡e`b c‡qvRb ‡bB weavq bvgÄyi Kiv mycwik K‡ib (Annex-B)|

<u>wm×vš t</u> mfvq me-সম্মতিক্রমে টেকনিক্যাল সব-KwgwUi mycwik MnY Kiv nq|

M| ¯vbxq Drcv`‡bi Rb¨ Av‡ew`Z ‡f‡Uwibvwix Jl‡ai †iwR‡÷k‡bi wbwg‡Ë `wLjKZ.27wU Jl‡ai g‡a¨ ‡UKwbK¨vj mve-KwgwUi m`m¨MY we¯wiZ Av‡jvPbv K‡i 21wU Jl‡ai ‡iwR‡÷kb Abyôgv`‡bi mycwik K‡ib Ges 6wU Jl‡ai Av‡e`b c‡qvRbxq ‡idv‡iÝmn ‡UKwbK¨vj mve-KwgwUi mfvq cieZx‡Z Dc¯vc‡bi mycwik K‡ib (Annex-C)|

wm×vš‘ t mfvq me-সম্মতিক্রমে টেকনিক্যাল সব-KwgwUi mycwik MnY Kiv nq|

N| Avg`vbxi Rb¨ ‡f‡Uwibvwix Jl‡ai †iwR‡÷k‡bi wbwg‡Ë `wLjKZ. 64wU Jl‡ai wel‡q mfvq †UKwbK¨vj mve-KwgwUi সদস্যগণ বিস্তারিত আলোচনাক্রমে ৬১টি ঔষধ আমদানী রেজিস্ট্রেশনের সুপwik K‡ib, 1wU Jla c‡qvRbxq †idv‡iÝmn cÖKvwiq Dc¯vc‡bi Rb¨ Ges 2wU Jla c‡qvRb ‡bB weavq Av‡e`b bvgÄyL Kivi mycwik K‡ib (Annex-D)|

wm×vš‘ t mfvq me-সম্মতিক্রমে টেকনিক্যাল সব-KwgwUi mycwik MnY Kiv nq|

‡f‡Uwibvwix dgj-vix cYqb সংক্রান্ত আলোচনা t

mfvq †ভেটেরিনারী ঔষধের ফর্মূলারী প্রণয়নের বিষয়ে আলোচনাক্রমে সদস্যগণ মত প্রকাশ করেণ ভেটেরিনারী Jla gj¨vq‡bi Rb¨ †Kvb †idv‡iÝ eB/Z¨ m`m¨‡`i Kv‡Q _v‡K bv †m‡ÿ‡Î ‡f‡Uwibvwix Jla gj¨vq‡Yi †ÿ‡Î †Kvb gvb`Û AbymiY Kiv hvq bv| ‡f‡Uwibvwix Jl‡ai h_vh_ gj¨vq‡bi Rb¨ GKwU †f‡Uwibvwix Jl‡ai dgj-vix cYq‡bi e¨e¯v MnY Kiv †h‡Z cv‡i|

wm×vš‘ t Jla ckvmb Awa`ßi †f‡Uwibvwix Jl‡ai dgj-vix cYq‡bi Rb¨ gš¿Yvj‡q GKwU Kwgwu MV‡bi c¯vebv †ck Ki‡eb|

0| gvbyl‡i †`‡n e¨envwi Avg`vbxKZ.Jl‡ai †iwR‡÷kb c`v‡bi †¶‡Î DbZ 07wU †`‡ki mv‡_ EMA-Gi wd †mj mwUwd‡KU Mn‡Yi wel‡q Av‡jvPbv I wm×vš‘ MnY cm‡½|

‡UKwbK¨vj mve-Kwgwui Av‡jvPbv I mycwik t

‡UKwbK¨vj mve-Kwgwui m`m¨ mwPe Rbve †gvt †Mvjvg wKewiqv, cwiPvjK, Jla ckvmb Awa`ßi Avg`vbxKZ. Jl‡ai †iwR‡÷kb c`v‡bi †¶‡Î DbZ 7 (mvZ) wU †`k h_v- hy³ivó, hy³ivR¨, mybRvij¨vÛ, Rvgvbx, dvÝ, Rvcvb I A‡÷wjqv-Gi mv‡_ EMA-Gi wd †mj mwUwd‡KU Ašfy³ Kivi c¯ve Av‡jvPbv Kivi Rb¨ mfvq Dc¯vb K‡ib|

G cm‡½ mfvi mfvcwZ ‡gRi †Rbv‡ij †gvt †gv¯wdRyi ingvb e‡jb †h, 7(mvZ)wU DbZ †`‡ki g‡a¨ hy³ivR¨, dvÝ Ges Rvgvb BD‡iwcqvb BDwbqbfz³ nIqvq EMA-KZK.c`Ë mwUwd‡KU Jl‡ai Avg`vbx †iwR‡÷k‡bi †ÿ‡Î MnY Kiv nq| G cm‡½ wZwb D‡jL K‡ib †h, BD‡iwcqvb BDwbq‡bi GKwU cwZwbwa `j Zui mv‡_ mvÿvrKv‡j EMA-c`Ë mwUwd‡KU-‡K AvBbx ‰eaZv c`v‡bi Rb¨ c¯ve K‡i‡Qb| G cm‡½ wZwb Av‡ivI e‡jb †h, weMZ 12 †g, 2016 Zwi‡L gvbbxq ewYR¨ gšxi †bZ‡Z ewYR¨ gšvj‡q BD‡ivcxq BDwbqbf-³ 11 (GMvi) wU †`‡ki ivó`Z‡`i mwnZ Abyôv‡Z EU-Bangladesh Business Climate Dialogue- G ivó`ZMY Jla Avg`vbxi †ÿ‡Î EMA-mwUwd‡KU-‡K wjMvj WKzg‡›U wn‡m‡e AvBbx KvVv‡gvi AvIZvq Avbvi c¯ve K‡ib| c¯weZ Jla bwZ‡Z Avg`vbxKZ.Jl‡ai †iwR‡÷kb c`v‡bi †ÿ‡Î DbZ 7 (mvZ) wU †`‡ki wd-‡mj mwUwd‡KU Mn‡Yi c¯ve Kiv n‡q‡Q| D‡jwLZ Jla bwZ‡Z ewbZ 7 (mvZ) wU DbZ †`‡ki wd-‡mj mwUwd‡KU-Gi mv‡_ EMA-mwUwd‡KU Ašf-³ Kiv †h‡Z cv‡i|

এ প্রসঙ্গে বিস্তারিত আলোচনাক্রমে সদস্যগণ মত প্রকাশ করেন যে, বিষয়টি সম্পর্কে সিদ্ধান্ত গ্রহণের জন্য ড্রাগ *K‡›Uvj KwgwU (wWwmwm) †Z Dc¯vcb Kiv ‡h‡Z cv‡i|*

mfvi Av‡jvPbv t

mfvi m`m¨ mwPe ‡gRi †Rbv‡ij †gvt †gv¯wdRyi ingvb mfvq Dc¯vcb K‡ib †h, Jl‡ai Avg`vbx †iwR‡÷k‡bi †ÿ‡Î DbZ 07wU †`k h_v- hy³ ivó, hy³ ivR¨, mgvRvij¨vÛ, Rvgvbx, dvÝ, Rvcvb I A‡÷wjqv-Gi mv‡_* EMA *I* Canada *Gi wd †mj mwUwd‡KU Ašfy³ Kiv wel‡q Av‡jvPbv Kiv c¯ve K‡ib| G cm‡½ KwgwUi AwaKvsk m`m¨ EMA (European Medicines Agency) *KZK.c`Ë mwUwd‡KU-‡K MnY Kiv †h‡Z cv‡i e‡j gZ cKvk K‡ib Ges KvbWv-‡K GB gyÝ‡Z Ašfz³ bv Kiv c‡ÿ AwfgZ e¨³ K‡ib|*

G cm‡½ KwgwUi m`m¨ Rbve mykgvi iÄb †NvI, Ggwc c¯ve K‡ib †h* EMA (European Medicines Agency) *f³ mKj iv‡÷i Zv‡`i Regulatory Agency *KZK.c`Ë mwUwd‡KU-‡K Jl‡ai Avg`vbx †iwR‡÷kb c`v‡bi †ÿ‡Î MnY Kiv †h‡Z cv‡i|*

mvgMxK welqwU chv‡jvPbv K‡i mfvcwZ gZvgZ e¨³ K‡ib †h, EMA (European Medicines Agency) *f³ 28 wU iv‡÷i mwUwd‡KU-‡K Jla Avg`vbx †iwR‡÷k‡bi Rb¨ MnY bv K‡i ïaygvÎ* EMA Certificate-*†K MnY Kiv †h‡Z cv‡i| mfvq Ab¨vb¨ m`m¨MY Zvnvi c¯ve‡K mg_b K‡ib|*

mfvi wm×vš t

Jl‡ai Avg`vbx †iwR‡÷kb c`v‡bi †ÿ‡Î DbZ 07 (mvZ) wU †`‡ki mv‡_ EMA Certificate-*†K MnY Kiv n‡e|*

P| *‡iwR‡÷kb mvgwqK ewZjKZ.45wU Jl‡ai †iwR‡÷kb ewZj cm‡½ t*

gvb-ewnfz nIqvi Kvi‡Y jvB‡mwÝs A_wiwU WvMm KZK.mvgwqK ewZjKZ.ewfb †Kv¤úvbxi Drcw`Z 45wU ev‡Ûi Jl‡ai †iwR‡÷ikb ewZ‡ji Rb¨ mfvq D¯vcb Kiv nq| GZ`wel‡q Jla ckvmb Awa`ß‡ii gnvcwiPvjK †gRi †Rbv‡ij †gvt †gv¯wdRyi ingvb e‡jb AvBbx eva¨evaKZvi Kvi‡Y c`¸wji †iwR‡÷kb ewZ‡ji Rb¨ KwgwUi Abygv`b c‡qvRb|

mfvi wm×vš t *সভায় বিস্তারিত আলোচনাক্রমে mvgwqK ewZjKZ.45wU Jl‡ai †iwR‡÷kb (*Annex-E) *ewZj Kiv wm×vš Mp,xZ nq|*

Q| Flupentixol + Melitracen Combination *RvZxq c`wUi ‡iwR‡÷kb ewZ‡ji wel‡q* ADRAC *KwgwUi PZz mfvi mycwik Abyhvqx Jla wbqšY KwgwUi †UKwbK¨vj mve-KwgwUi mycwik m¤ú‡K Av‡jvPbv I wm×vš t*

‡UKwbK¨vj mve-KwgwUi m`m¨ mwPe* Flupentixol + Melitracen *Gi* Combination *JlawU ewZ‡ji wel‡q* ADRAC (Adverse Drug Reaction Advisory Committee) *Gi weMZ 14-01-2015 Zwi‡Li mfvi mycwik †UKwbK¨vj mve KwgwUi mfvq Dc¯vcb K‡ib| wZwb mfv‡K AewnZ K‡ib,* ADRAC *Gi wØZxq mfvi wm×vš Abyhvqx Ab¨vb¨ K‡qKwU c‡`i mwnZ Flupentixol + Melitracen Combination *c`wUi ewZ‡ji mycwik Jla wbqšY KwgwU (wWwmwm)-Gi 243 mfvq Dc¯vcb Kiv n‡j, cwß Z_¨-DcvË WK‡qg›Um AwaKZi chv‡jvPbv K‡i* ADRAC *KwgwU‡Z cybivq wm×vš Mn‡Yi Rb¨ †diZ cvVv‡bv nq|*

ADRAC Kwgwi ZZxq mfvq Flupentixol + Melitracen Combination c`wi wel‡q wm×vš Mn‡Yi Rb¨ 6 (Qq) m`m¨ wewkó Dc-Kwgw MVb Kiv nq| D³ Dc-Kwgwi 03-12-2014 wLt Zwi‡Li mfvi wm×v‡ši Av‡jv‡K ADRAC-Gi 14-01-2015 wLt Zwi‡Li 4_ mfvq Flupentixol + Melitracen Combination c`wi ewZ‡ji mgvwik Kiv n‡q‡Q hv AvR‡Ki mfvq Dc¯vcb Kiv nj|

Dchy³ wel‡q †UKwbK¨vj mve Kwgwi wewfb m`m¨MY Av‡jvPbvq AskMnY K‡ib|

G wel‡q Kwgwi m`m¨ Rbve Avãyj gy³w`i mv‡eK mvaviY m¤úv`K evsjv‡`k Jla wkí mwgwZ I e¨e¯vcbv cwiPvjK, Bb‡mÞv dvgwmDwUK¨vjm wjt e‡jb †h, Jla wbqšY Kwgwi 243 mfv †_‡K c`wi wel‡q wm×vš Mn‡Yi Rb¨ ADRAC Kwgw‡Z †diZ cvVv‡bvi Kv‡iY wQj, †`‡k Jlaw‡U m¤ú‡K কোন বিরূপ প্রতিক্রিয়া রিপোর্ট পাওয়া গেলে তা পর্যালোচনা করা। দেশে Flupentixol + Melitracen c`wi কোন বিরূপ প্রতিক্রিয়া পাওয়া গেছে ADRAC-Gi wi‡cv‡U Gi|c †Kvb Z_¨ †bB| Dciš ÿwZKi cvk প্রতিক্রিয়ার কারণে ঔষধটি কোন দে‡k wbwl× †Nwl‡Z nqwb|

G cm‡½ Rbve bvRgyj nvmvb Gg,wc, Ges mfvcwZ evsjv‡`k Jla wkí mwgwZ e‡jb BD‡iv‡c Switzerland, Spain, Belgium, Austria, Luxembourg, Bulgaria, Malta ‡Z Jlaw‡U cPwjZ Av‡Q Ges Gwkqv‡Z Singapore, Thailand, Hong Kong, China, Vietnam and Taiwan G Jlaw‡U cPwjZ Av‡Q| ADRAC Kwgwi Awbw`ó Z_¨ wbfi mgvwik Abjyvqx c`w wbwl× Kivi †Kvb KviY †bB|

G cm‡½ Av‡jvPbvq Ask MnY K‡i Aa¨vcK †mwjg †iRv Rvb‡Z Pvb AZx‡Z evsjv‡`‡k cPwjZ wQj, Ggb Jla wbwl× Kivi †Kvb D`vniY Av‡Q wK bv|

G cm‡½ Kwgwi m`m¨ Rbve Avãyj gy³ াদির বলেন, ক্ষতিকর পার্শ্ব প্রতিক্রিয়ার কারণে USFDA ev DbZ we‡ki Ab¨ Regulatory Agency KZK wbwl×KZ.ev Awi‡R‡bUi †Kv¤úvwbi ¯-D‡`¨‡M cZ¨vnvi KZ. wKQy Jla Z_¨ m‡Î wWmwm KZK. evsjv‡`‡k wbwl× Kiv n‡q‡Q| Flupentixol + Melitracen Combination c`wi USFDA ev MHRA KZK. wbewÜZ nqwb weavq H mg¯ †`‡k c`w †iwR‡÷kb ewZj Kiv nq wb|

mwPeK we‡ePbvq c`wi †iwR‡÷kb envj ivLv †h‡Z cv‡i, Kwgwi Ab¨vb¨ m`m¨MY G wel‡q GKgZ †cvlY K‡ib|

†UKwbK¨vj mve Kwgwi mgvwik t Flupentixol + Melitracen Combination RvZxq c`wi ‡iwR‡÷kb envj ivLv †h‡Z cv‡i|

mfvi wm×vš t সভায় বিস্তারিত আলোচনাক্রমে টেকনিক্যাল সাব-Kwgwi mgvwik Abjyg`b Kiv nq|

2| MZ 25 †g 2016 Ges 15 Rby 2016 Zwi‡L Abyôv‡Z ‡gvW‡Kj wWfvB‡mi †iwR‡÷k‡bi Av‡e`b gj¨vqv** সংক্রান্ত কমিটির **mfvq mgvwikKZ.¯vbxq Drcv`‡bi Rb¨ 2 wU Ges Avg`vbxi Rb¨ 257 wU †gvW‡Kj wWfvB‡mm Gi †iwR‡÷k‡bi wbwg‡Ë `wLjKZ.Av‡e`b Gi Dci gZvgZ c`vb cm‡½|

¯vbxq Drcv`‡bi Rb¨ 2 (`k) wU Ges Avg`vbxi Rb¨ 257 wU †gvW‡Kj wWfvBm-Gi †iwR‡÷kb Abjyg`v‡bi wel‡q mfvq Dc¯vcb Kiv n‡j m`m¨MY we¯wiZ Av‡jvPbv ceK gZvgZ c`vb K‡ib th,

Av‡e`Z †gvW‡Kj wWfvBmmgn- evsjv‡`‡k cPwjZ nIqv c‡qvRb| †m Abyhvqx D‡jwLZ †gvW‡Kj wWfvBmmgn-†iwR‡÷‡kb c`v‡bi mycwik Kiv nq (Annex-F)|

15-06-২০১৬ তারিখে অনুষ্ঠিত মেডিকেল ডিভাইস সংক্রান্ত টেকনিক্যাল সাব-KwgwUi mfvq wewea Av‡jvPbv I wm×vš t

K| W. gyw¤§` ZwiK Avivdv‡Z A‡_v‡cwWK RvZxq †gvW‡Kj wWfvBm-Gi Component Kvhc‡Î D‡jL Kiv‡i c¯‘e K‡ib|

L| Wvt †gv¯‘vK Avn‡¤§` KwWIjwRi Stent-mg‡n-i wel‡q NICVD-Gi we‡klÁ‡`i gZvgZ MnY Kiv c‡qvRb, †h‡nZzZvi Stent-mgn- c‡hvR¨ †ivMx‡`i Rb¨ wbqwgZ e¨envi K‡i _v‡Kb| G cm‡½ mfvi mfvcwZ e‡jb †h, KwgwUi cieZx mfvq †gvW‡Kj wWfvBm h_vh_fv‡e gj¨vq‡bi wbwg‡Ë A‡_v‡cwWK, KwWIjwR, BGbwU Ges BD‡ivjwR we‡klÁ‡`i Aš—f³KiY ev Zv‡`i gZvgZ MnY Kiv n‡e|

‡UKwbK¨vj mve-KwgwUi mycwik t

K) A‡_v‡cwWK RvZxq †gvW‡Kj wWfvBm-Gi Component Kvhc‡Î D‡jL Ki‡Z n‡e|
L) †gvW‡Kj wWfvBm h_vh_fv‡e gj¨vq‡bi wbwg‡Ë A‡_v‡cwWK, KwWIjwR, BGbwU Ges BD‡ivjwR we‡klÁ‡`i Aš—f³KiY ev Zv‡`i gZvgZ MnY Ki‡Z n‡e|

mfvi wm×vš t ‡UKwbK¨vj mve-KwgwUi mycwik Abyhvg`b Kiv nq|

3| MZ 13/06/2016 Zwi‡L Abyôv‡Z nvevj Jla †UKwbK¨vj mve KwgwUi mfvi Dc¯wcZ 129 wU nvevj Jl‡ai g‡a¨ Abyhvg`‡bi mycwikKZ. 97 wU Ges bv-gÄyi mycwikKZ. 32 wU nvevj Jl‡ai †iwR‡÷kb c`vb m¤ú‡K Av‡jvPbv I wm×vš MnY cm‡½|

nvevj Jla †UKwbK¨vj mve KwgwUi mfvq 129 wU bZzb Jl‡ai †iwR‡ókb m¤ú‡K we¯wiZ Av‡jvPbv †k‡l 97 wU nvevj Jl‡ai Abyhvg`‡bi mycwik Kiv nq Ges 32 wU Jla bv-gÄy Kiv nq (Annex-G)|

mfvi wm×vš t mfvq me-সম্মতিক্রমে হার্বাল টেকনিক্যাল সাব-KwgwUi wm×vš Abyhvg`b Kiv nq|

nvevj Jla †UKwbK¨vj mve KwgwUi mfvq wewea Av‡jvPbv I wm×vš t

K| nvevj Jl‡ai gvb wbwðZK‡í nvem Gi Marker Component Gi Identity I Assay Kwievi j‡ÿ Jla c¯ZKvix cwZôvb hvnv‡Z c‡qvRbxq e¨e¯v MnY K‡i ‡mB wel‡q we¯wiZ Av‡jvPbv Kiv nq Ges welqwU hvnv‡Z Jla c¯ZKvix cwZôvbmgn- hvnv‡Z cwZcvjb K‡i Zvnv wbwðZ Kwievi j‡ÿ Jla ckvmb Awa`ßi‡K Abyiva Kiv nq| Bnv Qvov nvem msM‡ni mwVK Drm m¤ú‡KI Av‡jvPbv nq|

wm×vš t nve‡mi Source Valid Kiv mn Marker Component Gi Identity I Assay wbwðZ Kwi‡Z nB‡e|

L| nvevj Jla GWfvBRix KwgwU Gi 12 (evi) Zg mfvq 15(c‡bi)wU Reference eB wnmv‡e wbaviY Kiv nq| D³ 15(c‡bi)wU Reference eB Gi cvkvcwk British National

Formulary (BNF) ‡KI Reference eB wnmv‡e ms‡hvRb Kwievi Rb¨ Dcw¯Z m`m¨e„`c¯ve K‡ib|

wm×vš t nvevj Jl‡ai Rb¨ c‡e wbawiZ 15(c‡bi)wU eB Gi cvkvcwk British National Formulary (BNF) ‡KI Reference eB wnmv‡e wbaviY Kiv nBj| D³ Reference eB mgn¨nvevj Jla Drcv`b, gvb-wbqšY I †ivM wb‡`kbvi Z_¨w`i Rb¨ Abyw„Z n‡e|

M| c¥MwVZ nvevj Jl‡ai †UKwbK¨vj mve-কমিটিতে ক্রমিক নং- 13 G ewYZ evsjv‡`k nvevj †gwWwmb g¨vbyd¨vKPvivm G‡mwm‡qk‡bi cwZwbwa wnmv‡e Aš—f©³ Av‡Q, wKš D³ G‡mwm‡qk‡bi ewYR¨ gšYvj‡qi †Kvb †iwR‡÷kb †bB, A_vr ewYR¨ gšYvj‡qi KZ„K Abyhvgw`Z bq, weavq ewYwR¨K msMVb wnmv‡e D³ G‡mwm‡qkb A‰ea| d‡j Zv‡`i cwZwbwaË nvevj Jl‡ai †UKwbK¨vj mve-KwgwU‡Z _vK‡Z cv‡i bv g‡g evsjv‡`k nvevj †cvWv± g¨vbyd¨vKPvivm G‡mwm‡qk‡bi cwZwbwa mfv‡K AewnZ K‡ib| Bnv QvovI D³ nvevj Jla GWfvBRix KwgwU‡Z XvKv wekwe`¨vj‡qi Dw™` weÁvb wefv‡Mi Plant Taxonomist †K cwZwbwa wn‡m‡e ivLvi Rb¨ K‡qKRb m`m¨ gZ cKvk K‡ib|

wm×vš t nvevj Jl‡ai †UKwbK¨vj mve-KwgwU‡Z evsjv‡`k nvevj †gwWwmb g¨vbyd¨vKPvivm G‡mwm‡qk‡bi cwZwbwa‡K ev` †`qv Ges GKRb Plant Taxonomist †K Aš—f©³ K‡i KwgwU c¥MVb Kivi wbwg‡Ë gšYvjq eivei Jla ckvmb Awa`ßi n‡Z c¯ve †ciY Kiv n‡e|

N| nvevj Jla GWfvBRix KwgwUi wØZxq mfvq Abyhvgw`Z nvevj Jl‡ai wbeÜb bxwZgvjv nvjbvMv` Kivi wel‡q we¯wiZ Av‡jvPbv nq|

wm×vš t nvevj Jl‡ai wbeÜb bxwZgvjvi Kwc nvevj Jla GWfvBRix KwgwUi mKj m`m¨ বরাবরে হালনাগাদকরন সংক্রান্ত বিষয়ে মতামত প্রদানের জন্য ঔষধ প্রশাসন অধিদপ্তর হতে প্রেরণ Kiv n‡e| gZvgZ cwß ¯v‡c‡ÿ Pvš bxwZgvjv cYqb Kiv n‡e|

mfvi wm×vš t mfvq me-সম্মতিক্রমে হার্বাল টেকনিক্যাল সাব-KwgwUi wm×vš Abyhvgv`b Kiv nq|

Ab¨ †Kvb Av‡jv‡¯¨ welq bv _vKvq mfvcwZ g‡nv`q Dcw¯Z mKj‡K ab¨ev` Ávcb K‡i mfvi mgwß †NvlYv K‡ib|

‡gRi †Rbv‡ij †gvt ‡gv¯wdRyi ingvb
gnvcwiPvjK
Jla ckvmb Awa`ßi
I
m`m¨-mwPe
Jla wbqšY KwgwU|

‰mq` gbRyj Bmjvg
mwPe
¯v¯¨ I cwievi Kj¨vY gšYvjq
I
mfvcwZ
Jla wbqšY KwgwU|

Annex-A

Proposed Product for Locally Manufacture (Human)

bs	c¯ZKvi‡Ki bvg	JI‡ai bvg I †RvbvwiK bvg	wb‡`kbv	Contra-indication & Side-effect	Status (New Molecule/ Existing)	Av‡e`bKvix c`Ë USFDA or MHRA Ref.	‡UKwbK¨vj mve-KwgwUi 64 Zg mfvi wm×vš	mfvi wm×vš
1.	Beximco Pharmaceuticals Ltd., Tongi ,Gazipur	L-Ornithine L-Aspartate 3.00gm/Sachet Granules for Oral Solution L-Ornithine L-Aspartate INN 3.00gm/Sachet **Ammonia detoxifying Agent**	It is indicated in the treatment of hyperammonemia due to acute or chronic liver diseases eg liver cirrhosis, fatty liver, hepatitis; for the treatment of incipient disturbances of consciousness (pre-coma) or neurological complications.	Contraindications: Renal dysfunction, Side effects: The most common side effect is nausea and vomiting etc.	L-Ornithin-L-Aspartate 150mg+ Pancreatin 100mg Enteric Coated Tablet		Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
2.	Libra Infusions Limited	2% Propofol Injectable Emulsion; 50ml Vial Propofol BP 20mg/ml; 50ml Vial **Anaesthetics**	Propofol belongs to a group of medicines called general anaesthetics. General anaesthetics are used to cause unconsciousness so that surgical operations or other procedures can be performed. Safety, effectiveness and dosing guidelines for Propofol Injectable Emulsion have not been established for MAC Sedation in the pediatric population; therefore, it is not recommended for this use. Propofol Injectable Emulsion is not recommended for induction of anesthesia below the age of 3 years or for maintenance of anesthesia below the age of 2 months because its safety and effectiveness have not been established in those populations.	Contra-indications: Propofol injectable emulsion is contraindicated in patients with a known hypersensitivity to Propofol injectable emulsion or any of its components. Propofol injectable emulsion is contraindicated in patients with allergies to eggs, egg products, soybeans or soy products. Side effects: General Adverse event information is derived from controlled clinical trials and worldwide marketing experience. In the description below, rates of the more common events represent US/Canadian clinical study results. Less frequent events are also derived from publications and marketing experience in over 8 million patients; there are insufficient data to support an accurate estimate of their incidence rates. These studies were conducted using a variety of premedicants, varying lengths of surgical/diagnostic procedures, and various other anesthetic/sedative agents. Most	1% Injection	MHRA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

				adverse events were mild and transient. Anesthesia and MAC Sedation in Adults The following estimates of adverse events for Propofol Injectable Emulsion include data from clinical trials in general anesthesia/MAC sedation (N=2889 adult patients). The adverse events listed below as probably causally related are those events in which the actual incidence rate in patients treated with Injectable Emulsion was greater than the comparator incidence rate in these trials. Therefore, incidence rates for anesthesia and MAC sedation in adults generally represent estimates of the percentage of clinical trial patients which appeared to have probable causal relationship. The adverse experience profile from reports of 150 patients in the MAC sedation clinical trials is similar to the profile established with Propofol Injectable Emulsion.				
3.	ACI Ltd., Narayanganj	Ibuprofen 100 mg/5 ml Oral Suspension in Sachet Ibuprofen BP 100 mg/5ml Analgesic	Children aged 3 months to 12 years: Mild to moderate pain due to sore throat, teething pain, toothache, rheumatic or muscular pain, headache, minor aches and pains, symptoms of cold and influenza, post-immunisation pyrexia and reduction of fever.	**Contraindications:** Hypersensitivity to ibuprofen or any of the excipients in the product. Patients who have previously shown hypersensitivity reactions (e.g. asthma, rhinitis, angioedema or urticaria) in response to aspirin or other non-steroidal anti-inflammatory drugs. Active or history of recurrent peptic ulcer / hemorrhage (two or more distinct episodes of proven ulceration or bleeding). History of gastrointestinal bleeding or perforation, related to previous NSAIDs therapy. Severe heart failure, renal failure or hepatic failure Last trimester of pregnancy. **Side Effects:** Hypersensitivity reactions have been reported and these may consist of: (a) Non-specific allergic reactions and anaphylaxis (b) Respiratory tract reactivity, e.g. asthma, aggravated asthma, bronchospasm-or dyspnoea.	New	MHRA BNF-70 Page-929	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

				(c) Various skin reactions, e.g. pruritis, urticaria, angioedema and more rarely exfoliative and bullous dermatoses (including epidermal necrolysis and erythema multiforme). Other Side effects are abdominal pain, acid sour stomach, diarrhea, oedema, headache, hypertension, cardiac failure etc.				
4.	Square Pharmaceuticals Ltd., Pabna Unit, Salgaria, Pabna	Nabumetone 500mg Tablet Nabumetone USP 500mg **Analgesic**	Nabumetone is indicated for relief of signs and symptoms of osteoarthritis and rheumatoid arthritis.	Contraindications: Use in patients with active, or a history of recurrent peptic ulcer/ GI haemorrhage, perforation or peptic disease (two or more distinct episodes). Use in patients hypersensitive to Nabumetone or to any of the excipients. Severe heart failure, hepatic failure and renal failure. Use in patients who have shown previous hypersensitivity reactions (e.g. asthma, rhinitis, angiodema or urticaria) in response to ibuprofen, aspirin or other non-steroidal anti-inflammatory drugs. Severe, rarely fatal, anaphylactic like reactions to NSAIDs have been reported in such patients. Use in patients with a history of gastrointestinal bleeding or perforation, related to previous NSAIDs therapy. During the last trimester of pregnancy and in nursing mothers. Patients with current cerebrovascular or other haemorrhage. Side effects: Undesirable effects may be minimized by using the lowest effective dose for the shortest duration necessary to control symptoms. The use of Nabumetone with concomitant NSAIDs, including cyclooxygenase-2 selective inhibitors should be avoided.	New	USFDA	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

5.	Beacon Pharmaceutical Ltd., Square Pharmaceuticals Ltd., Pabna Unit, Salgaria, Pabna	Nabumetone 750mg Tablet Nabumetone USP 750mg **Analgesic**	Nabumetone is indicated for relief of signs and symptoms of osteoarthritis and rheumatoid arthritis.	Contraindications: Use in patients with active, or a history of recurrent peptic ulcer/ GI haemorrhage, perforation or peptic disease (two or more distinct episodes). Use in patients hypersensitive to Nabumetone or to any of the excipients. Severe heart failure, hepatic failure and renal failure. Use in patients who have shown previous hypersensitivity reactions (e.g. asthma, rhinitis, angiodema or urticaria) in response to ibuprofen, aspirin or other non-steroidal anti-inflammatory drugs. Severe, rarely fatal, anaphylactic like reactions to NSAIDs have been reported in such patients. Use in patients with a history of gastrointestinal bleeding or perforation, related to previous NSAIDs therapy. During the last trimester of pregnancy and in nursing mothers. Patients with current cerebrovascular or other haemorrhage. Side effects: Undesirable effects may be minimized by using the lowest effective dose for the shortest duration necessary to control symptoms. The use of Nabumetone with concomitant NSAIDs, including cyclooxygenase-2 selective inhibitors should be avoided.	New	USFDA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
6.	Incepta Pharmaceuticals Ltd.	Nalbuphine Hydrochloride 20mg/ml Injection Nalbuphine Hydrochloride INN 20 mg/ml Analgesic	Nalbuphine hydrochloride is indicated for the relief of moderate to severe pain. Nalbuphine hydrochloride can also be used as a supplement to balanced anesthesia, for preoperative and postoperative analgesia, and for obstetrical analgesia during labor and delivery.	Contraindication: Nalbuphine hydrochloride injection should not be administered to patients who are hypersensitive to nalbuphine hydrochloride, or to any of the other ingredients in nalbuphine hydrochloride injection. Side effect: The most frequent adverse reaction in 1066 patients treated with nalbuphine hydrochloride injection was sedation 381 (36%). Less frequent reactions were: sweaty/clammy 99 (9%), nausea/ vomiting 68 (6%), dizziness/vertigo 58 (5%), dry mouth 44 (4%), and headache 27 (3%).	10mg/ml Injection 20mg/2ml Injection	USFDA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
7.	Eskayef Bangladesh	Phenazopyridine Hydrochloride	It is indicated for the symptomatic	Contraindications:	New		c‡qvRbxq †id v‡i Y bv_vKq	c‡qvRbxq †id v‡i Y bv_vKq

	Limited	100mg FC Tablet Phenazopyridine Hydrochloride USP 100mg Analgesic	relief of pain, burning, urgency, frequency, and other discomforts resulting from irritation of the mucosa of the lower urinary tract caused by infection, trauma, surgery, endoscopic procedures, or the passage of sounds or catheters. It is compatible with antimicrobial therapy and can help relieve pain and discomfort during the interval before an antimicrobial therapy controls the infection.	• Patients who are hypersensitive to the drug or its ingredients. • Patients with renal insufficiency or any liver disease. • Side effects: • Gastrointestinal: nausea, vomiting and diarrhea. • Nervous System: headache, aseptic meningitis. • Integumentary: rash, pruritus, discoloration, jaundice, Renal: renal toxicity usually associated with overdose, renal calculi.			*Av‡e`b bvgÄyi Kiv †h‡Z cv‡i |*	*Av‡e`b bvgÄyi Kiv nj |*
8.	Eskayef Bangladesh Limited.	Phenazopyridine Hydrochloride 200mg FC Tablet Phenazopyridine Hydrochloride USP 200mg Analgesic	Do	Contraindications: • Patients who are hypersensitive to the drug or its ingredients. • Patients with renal insufficiency or any liver disease. • Side effects: • Gastrointestinal: nausea, vomiting and diarrhea. • Nervous System: headache, aseptic meningitis. • Integumentary: rash, pruritus, discoloration, jaundice, Renal: renal toxicity usually associated with overdose, renal calculi.	New		*c‡qvRbxq †idv‡iÝ bv_vKq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i |*	*c‡qvRbxq †idv‡iÝ bv_vKq Av‡e`b bvgÄyi Kiv nj |*
9.	Incepta Pharmaceuticals Ltd.	Remifentanil 2.0mg/Vial Lyophilized Powder for Injection Remifentanil Hydrochloride INN 2.194mg eq.to Remifentanil 2.0mg /Vial Analgesic	Remifentanil is indicated for IV administration: As an analgesic agent for use during the induction and maintenance of general anesthesia for inpatient and outpatient procedures. For continuation as an analgesic into the immediate postoperative period in adult patients under the direct supervision of an anesthesia practitioner in a postoperative anesthesia care unit or intensive care setting. As an analgesic component of	Contraindication: Due to the presence of glycine in the formulation, Remifentanil is contraindicated for epidural or intrathecal administration. Remifentanil is also contraindicated in patients with known hypersensitivity to fentanyl analogs. Side effect: Remifentanil produces adverse events that are characteristic of μ-opioids, such as respiratory depression, bradycardia, hypotension, and skeletal muscle rigidity. These adverse events dissipate within minutes of discontinuing or decreasing the infusion rate of	1mg/ml Injection	USFDA	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

			monitored anesthesia care in adult patients.	Remifentanil.				
10.	Incepta Pharmaceuticals Ltd.	Remifentanil 5 mg/Vial Lyophilized Powder for Injection Remifentanil Hydrochloride INN 5.485mg eq.to Remifentanil 5 mg /Vial Analgesic	Remifentanil is indicated for IV administration: As an analgesic agent for use during the induction and maintenance of general anesthesia for inpatient and outpatient procedures. For continuation as an analgesic into the immediate postoperative period in adult patients under the direct supervision of an anesthesia practitioner in a postoperative anesthesia care unit or intensive care setting. As an analgesic component of monitored anesthesia care in adult patients.	Contraindication: Due to the presence of glycine in the formulation, Remifentanil is contraindicated for epidural or intrathecal administration. Remifentanil is also contraindicated in patients with known hypersensitivity to fentanyl analogs. Side effect: Remifentanil produces adverse events that are characteristic of μ-opioids, such as respiratory depression, bradycardia, hypotension, and skeletal muscle rigidity. These adverse events dissipate within minutes of discontinuing or decreasing the infusion rate of Remifentanil.	1mg/ml Injection	USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
11.	UniMed & UniHealth Mfg. Ltd., Gazipur Healthcare Pharmaceuticals Ltd., Gazipur	Tapentadol 150 Modified Release Tablet Tapentadol HCl INN 174.75mg eq. to 150mg Tapentadol Analgesic (Opioid)	Moderate to severe acute pain which can be managed only with opioid analgesic	Contra-indications: Opioid analgesics should be avoide in patients with acute respiratory depression and when there is a risk of paralytic ileus. They are also contra-indicated in condition s associated with raised intracranial pressure and in head injury (opioid analgesic interfere with papillary responses vital for neurological assessment). Comatose patients should not be treated with opioid analgesics. Side effects: Opioid analgesics share many sideeffects, although qualitative and quantitative differences exist. The most common side-effects include nausea and vomiting (particularly in initial stages), constipation, dry mouth, and biliary spasm; larger doses produce muscle rigidity, hypotension, and respiratory depression. Other common side-effects of opioid analgesics include bradycardia, tachycardia, palpitation, oedema, postural hypotension, hallucinations, vertigo, euphoria, dysphoria, mood changes, dependence, dizziness, confusion, drowsiness, sleep disturbances,	100mg MR Tablet	BNF-70 Page-372	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

				headache, sexual dysfunction, difficulty with micturition, urinary retention, ureteric spasm, miosis, visual disturbances, sweating, flushing, rash, urticaria, and pruritus. also decreased appetite, diarrhoea, dyspepsia, abdominal discomfort, weight loss, anxiety, tremor, ataxia, dysarthria, hypoaesthesia, paraesthesia, seizures, malaise, muscle spasms.				
12.	UniMed & UniHealth Mfg. Ltd., Gazipur	Tapentadol 50mg Modified Release Tablet Tapentadol HCl INN 58.25mg eq. to 50mg Tapentadol Analgesic (Opioid)	Do	Contra-indications: Opioid analgesics should be avoide in patients with acute respiratory depression and when there is a risk of paralytic ileus. They are also contra-indicated in condition s associated with raised intracranial pressure and in head injury (opioid analgesic interfere with papillary responses vital for neurological assessment). Comatose patients should not be treated with opioid analgesics. Side effects: Opioid analgesics share many sideeffects, although qualitative and quantitative differences exist. The most common side-effects include nausea and vomiting (particularly in initial stages), constipation, dry mouth, and biliary spasm; larger doses produce muscle rigidity, hypotension, and respiratory depression. Other common side-effects of opioid analgesics include bradycardia, tachycardia, palpitation, oedema, postural hypotension, hallucinations, vertigo, euphoria, dysphoria, mood changes, dependence, dizziness, confusion, drowsiness, sleep disturbances, headache, sexual dysfunction, difficulty with micturition, urinary retention, ureteric spasm, miosis, visual disturbances, sweating, flushing, rash, urticaria, and pruritus. also decreased appetite, diarrhoea, dyspepsia, abdominal discomfort, weight loss, anxiety, tremor, ataxia, dysarthria, hypoaesthesia, paraesthesia, seizures, malaise, muscle spasms.	100mg MR Tablet	BNF-70 Page-372	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj

13.	ACI Ltd., Narayanganj	Paracetamol 120 mg/5 ml Oral Suspension in Sachet Paracetamol BP 120mg/5ml Analgesic and Antipyretic	It is indicated for the treatment of mild to moderate pain and as an antipyretic. It can be used in many conditions including headache, toothache, earache, teething, sore throat, colds & influenza, aches and pains and post-immunisation fever.	Contraindications: Hypersensitivity to Paracetamol or any of the excipients. Side Effects: Adverse effects of paracetamol are rare but hypersensitivity including skin rash may occur. Very rare cases of serious skin reactions have been reported. Very rarely there have been reports of blood dyscrasias including thrombocytopenia and agranulocytosis, but these were not necessarily causally related to paracetamol. Most reports of adverse reactions to paracetamol related to overdosage and overusage of the drug	120 mg/ 5 ml Oral suspension	MHRA BNF-70 (page no.356)	c‡qvRb ‡bB wea‡q Av‡e`b bv gÄy Kiv †h‡Z cv‡i	c‡qvRb ‡bB wea‡q Av‡e`b bv gÄy Kiv nj
14.	ACI Ltd., Narayanganj	Paracetamol 250 mg/5 ml Oral Suspension in Sachet Paracetamol BP 250mg/5ml Analgesic and Antipyretic	It is indicated for the treatment of mild to moderate pain and as an antipyretic. It can be used in many conditions including headache, toothache, earache, teething, sore throat, colds & influenza, aches and pains and post-immunisation fever.	Contraindications: Hypersensitivity to Paracetamol or any of the excipients. Side Effects: Adverse effects of paracetamol are rare but hypersensitivity including skin rash may occur. Very rare cases of serious skin reactions have been reported. Very rarely there have been reports of blood dyscrasias including thrombocytopenia and agranulocytosis, but these were not necessarily causally related to paracetamol. Most reports of adverse reactions to paracetamol related to overdosage and overusage of the drug	120 mg/ 5 ml Oral suspension	MHRA BNF-70 (page no.356)	c‡qvRb ‡bB wea‡q Av‡e`b bv gÄy Kiv †h‡Z cv‡i	c‡qvRb ‡bB wea‡q Av‡e`b bv gÄy Kiv nj
15.	Ziska Pharmaceuticals Ltd.	Oxymorphone Hydrochloride 20.0 mg Extended Release Tablet Oxymorphone Hydrochloride USP 20.0mg Narcotic analgesic	It is an opioid agonist indicated for the management of pain severe enough to require daily, around-the-clock, long-term opioid treatment and for which alternative treatment options are inadequate. Limitations of Use: Because of the risks of addiction, abuse, and misuse with opioids, even at recommended doses, and because of the greater risks of overdose and death with extended-release opioid formulations, reserve this tablet for use in patients for whom alternative treatment options (e.g., non-opioid analgesics or immediate-release opioids) are ineffective, not tolerated, or would	Contraindications: Significant respiratory depression, Acute or severe bronchial asthma, Known or suspected paralytic ileus, Hypersensitivity to Oxymorphone, Moderate or severe hepatic impairment. Side effects: Adverse reactions in ≥2% of patients in placebo-controlled trials: nausea, constipation, dizziness, somnolence, vomiting, pruritus, headache, sweating increased, dry mouth, sedation, diarrhea, insomnia, fatigue, appetite decreased, and abdominal pain.	10mg IR Tablet	USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

			be otherwise inadequate to provide sufficient management of pain. It is not indicated as an as-needed (prn) analgesic					
16.	Mundipharma (Bangladesh) Private Limited., Gazipur.	Oxycodone Hydrochloride Controlled Release USP 10mg Tablet Oxycodone Hydrochloride Controlled Release USP 10mg Analgesic, Opoid	Indicated for pain severe enough to require daily, around-the-clock, long-term opioid treatment. .	Contraindication: • Patients who are hypersensitive to the active substance (oxycodone) or other opioid analgesics or to any ingredient in the formulation,acute appendicitis or pancreatitis, acute asthma or other obstructive airway, acute respiratory depression, elevated carbon dioxide levels in the blood, and cor pulmonale. Side effects: The most frequently observed adverse are asthenia, constipation, dizziness, dry mouth, headache, nausea, pruritus, somnolence, sweating and vomiting.	New	USFDA & BNF-70; Page: 370(MR)	Abyjgv`b Kiv †h‡Z cv‡i|	Abyjgv`b Kiv nj|
17.	Mundipharma (Bangladesh) Private Limited., Gazipur.	Oxycodone Hydrochloride Controlled Release USP 20mg Tablet Oxycodone Hydrochloride Controlled Release USP 20mg Analgesic, Opoid	Do	Contraindication: • Patients who are hypersensitive to the active substance (oxycodone) or other opioid analgesics or to any ingredient in the formulation,acute appendicitis or pancreatitis, acute asthma or other obstructive airway, acute respiratory depression, elevated carbon dioxide levels in the blood, and cor pulmonale. Side effects: The most frequently observed adverse are asthenia, constipation, dizziness, dry mouth, headache, nausea, pruritus, somnolence, sweating and vomiting.	New	USFDA & BNF-70; Page: 370(MR)	Abyjgv`b Kiv †h‡Z cv‡i|	Abyjgv`b Kiv nj|
18.	Eskayef Bangladesh Limited.	Glucosamine HCl BP 10.0gm/100ml Oral Solution Glucosamine HCl BP 10.0gm/100ml Analgesics	Relief of symptoms in mild to moderate osteoarthritis of the knee.	Contraindications: • Known hypersensitivity to glucosamine or to any of the Excipients. • It must not be given to patients who are allergic to shellfish. • It must not be given to patients who suffer from phenylketonuria.	500mg Tablet		c‡qvRbxq †idv‡iY bv_vKq Av‡e`b bvgÄyi Kiv †h‡Z cv‡i|	c‡qvRbxq †idv‡iY bv_vKq Av‡e`b bvgÄyi Kiv হল|

				Side effects: Nausea, abdominal pain, dyspepsia, headache, somnolence, rash, pruritus and erythema.				
19.	ACI Ltd., Narayanganj	Zileuton 600 mg ER Tablets Zileuton INN 600 mg Anti Asthamatic	Zileuton is a leukotriene synthesis inhibitor indicated for the prophylaxis and chronic treatment of asthma in adults and children 12 years of age and older. Do not use Zileuton to treat an acute asthma attack.	Contraindications: Active liver disease or persistent hepatic function enzyme elevations ≥3 times the upper limit of normal. History of allergic reaction to zileuton or any of the ingredients. Side Effects: Most common adverse reactions included: sinusitis, nausea, and pharyngolaryngeal pain.	New	USFDA	*c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i|*	*c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv* হল|
20.	Beacon Pharmaceutical Ltd.,	Zileuton 600 mg ER Tablets Zileuton INN 600 mg Antiasthmatic	Zileuton is a leukotriene synthesis inhibitor indicated for the prophylaxis and chronic treatment of asthma in adults and children 12 years of age and older. Do not use Zileuton to treat an acute asthma attack.	Contraindications: Active liver disease or persistent hepatic function enzyme elevations ≥3 times the upper limit of normal. History of allergic reaction to zileuton or any of the ingredients. Side Effects: Most common adverse reactions included: sinusitis, nausea, and pharyngolaryngeal pain.	New	USFDA (CR Tablet)	*c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i|*	*c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv* হল|
21.	Eskayef Bangladesh Limited	Metronidazole 1.3gm/100gm Vaginal Gel Metronidazole BP 1.3gm/100gm Antibacterial	Metronidazole vaginal gel 1.3% is indicated in the treatment of bacterial vaginosis in non-pregnant women.	Contraindications: • History of hypersensitivity to metronidazole, parabens, other ingredients of the formulation, or other nitroimidazole derivatives. • Concomitant use of disulfiram or within 2 weeks of disulfiram. • Concomitant use of alcohol. Side effects: The most common side effects observed in clinical studies (incidence ≥1%) were vulvovaginal candidiasis, headache, vulvovaginal pruritus, nausea, diarrhea, and dysmenorrhea.	0.75% Vaginal Gel	USFDA	*Aby튠gv`b Kiv †h‡Z cv‡i|*	*Abyটgv`b Kiv nj|*
22.	Globe Pharmaceuticals Ltd., Noakhali	Minocycline 100 mg FC Tablet Minocycline HCl BP 115.85 mg eq. to Minocycline 100 mg Antibacterial	Minocycline is a broad spectrum antibiotic used for treatment of infections caused by tetracycline sensitive organisms. Tetracyclines, including minocycline, are the treatment of choice for infections caused by:	Contraindications: Hypersensitivity to the active substance, or to any of the excipients listed in section • Known hypersensitivity to tetracyclines •Pregnancy and lactation • Systemic lupus erythematosus (SLE) • Complete renal failure	New	MHRA BNF 70 Page: 498	*Abyটgv`b Kiv †h‡Z cv‡i|*	*Abyটgv`b Kiv nj|*

			Chlamydia (trachoma, non-gonococcal urethritis, pelvic inflammatory disease, prostatitis, epididymitis, lymphogranuloma venereum and psittacosis). • Rickettsia (typhus, Q-Fever, Rocky mountain spotted fever). • Mycoplasma (respiratory and genital). • Brucella (normally in combination with other antibiotics such as streptomycin). • Borrelia burgdorferi (Lyme disease). Other uses include: • acute and chronic bronchitis, • acne and, in cases of penicillin allergy, • treatment of actinomycosis, nocardiosis, leptospirosis and syphilis. Minocycline is also used for meningococcal prophylaxis.	Paediatric population Not recommended for children under 12 years. Side effects: Rare- Acute renal failure, alopecia, anorexia, hyperaesthesia, impaired hearing, paraesthesia, pigmentation, tinnitus Very rare- Discoloration of conjunctiva, discoloration of sweat, discoloration of tears, systemic lupus. Frequent not known- Dizziness (more common in women), vertigo (more common in women).				
23.	Incepta Pharmaceuticals Ltd. Beacon Pharmaceutical Ltd.,	Ado-Trastuzumab Emtansine 100mg/Vial lyopphilized Powder for Injection Ado-Trastuzumab Emtansine Solution (Sterile) INN 2.0ml eq. to Ado-Trastuzumab Emtansine 100 mg/Vial Anticancer	Ado-Trastuzumab is a HER2-targeted antibody and microtubule inhibitor conjugate indicated, as a single agent, for the treatment of patients with HER2-positive, metastatic breast cancer who previously received trastuzumab and a taxane, separately or in combination. Patients should have either: • Received prior therapy for metastatic disease, or • Developed disease recurrence during or within six months of completing adjuvant therapy.	Contraindication: None Side effect: The most common adverse drug reactions (frequency > 25%) with Ado-Trastuzumab (n=884 treated patients) were fatigue, nausea, musculoskeletal pain, hemorrhage, thrombocytopenia, headache, increased transaminases, constipation and epistaxis.	New	USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
24.	Incepta Pharmaceuticals Ltd.	Ado-Trastuzumab Emtansine 160 mg/Vial lyopphilized Powder for Injection	Ado-Trastuzumab is a HER2-targeted antibody and microtubule inhibitor conjugate indicated, as a	Contraindication: None Side effect: The most common adverse	New	USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

		Ado-Trastuzumab Emtansine Solution (Sterile) INN 2.5ml eq. to Ado-Trastuzumab Emtansine 160 mg/Vial Anticancer	single agent, for the treatment of patients with HER2-positive, metastatic breast cancer who previously received trastuzumab and a taxane, separately or in combination. Patients should have either: • Received prior therapy for metastatic disease, or • Developed disease recurrence during or within six months of completing adjuvant therapy	drug reactions (frequency > 25%) with Ado-Trastuzumab (n=884 treated patients) were fatigue, nausea, musculoskeletal pain, hemorrhage, thrombocytopenia, headache, increased transaminases, constipation and epistaxis.				
25.	Beacon Pharmaceutical Ltd.,	Ceritinib 150 mg Capsule Ceritinib INN 150mg Anticancer	It is indicated for the treatment of patients with locally advanced or metastatic non-small cell lung cancer (NSCLC) that is anaplastic lymphoma kinase (ALK)-positive.	Contraindication: Hypersensitivity to either of the active substances or to any of the excipients. Side Effects: Anemia, decreased appetite, hyperglycemia, hypophosphatemia, vision disorder, pericarditis, bradycardia, diarrhea, nausea, vomiting, abdominal pain, constipation, esophageal disorder, rash, fatigue, liver laboratory test abnormalities, blood creatinine increased	New	USFDA	*Abÿgv`b Kiv †h‡Z cv‡i |*	*Abÿgv`b Kiv nj |*
26.	Beacon Pharmaceuticals Ltd.	Dasatinib 20mg Film coated Tablet Dasatinib Monohydrate INN 20.738mg eq. to Dasatinib 20mg Anti cancer	Dasatinib is a kinase inhibitor indicated for the treatment of • newly diagnosed adults with Philadelphia chromosome-positive (Ph+) chronic myeloid leukemia (CML) in chronic phase. •adults with chronic, accelerated, or myeloid or lymphoid blast phase Ph+ CML with resistance or intolerance to prior therapy including imatinib. •adults with Philadelphia chromosome-positive acute lymphoblastic leukemia (Ph+ ALL) with resistance or intolerance to prior therapy.	Contraindication: None, Warnings & Precautions: Myelosuppression and Bleeding Events: Severe thrombocytopenia ncutropenia and ancmia may occur Use caution if used concomitantly with medications that inhibit platelet function or anticoagulants Monitor complete blood counts regulary Transfuse and interrupt SPRYCEL (dasatinib) when indicated. Fluid Retention ; Fluid retention, sometimes severe including pleural effusions Manage with supportive care measures and/or dose modification Side-effect: Most common adverse reactions (≥15%) in patients with newly diagnosed chronic phase CML included myelosuppression, fluid retention, and diarrhea. Most common adverse reactions (≥15%) in patients with resistance or intolerance to prior imatinib therapy included myelosuppression, fluid retention events, diarrhea, headache, fatigue,	100mg Tablet	USFDA	*Abÿgv`b Kiv †h‡Z cv‡i |*	*Abÿgv`b Kiv nj |*

				dyspnea, skin rash nausea, hemorrhage and musculoskeletal pain.				
27.	Beacon Pharmaceuticals Ltd.	Dasatinib 50mg Film coated Tablet Dasatinib Monohydrate INN 51.845mg eq. to Dasatinib 50mg Anti cancer	Dasatinib is a kinase inhibitor indicated for the treatment of • newly diagnosed adults with Philadelphia chromosome-positive (Ph+) chronic myeloid leukemia (CML) in chronic phase. •adults with chronic, accelerated, or myeloid or lymphoid blast phase Ph+ CML with resistance or intolerance to prior therapy including imatinib. •adults with Philadelphia chromosome-positive acute lymphoblastic leukemia (Ph+ ALL) with resistance or intolerance to prior therapy.	Contraindication: None, Warnings & Precautions: Myelosuppression and Bleeding Events: Severe thrombocytopenia ncutropenia and ancmia may occur Use caution if used concomitantly with medications that inhibit platelet function or anticoagulants Monitor complete blood counts regulary Transfuse and interrupt SPRYCEL (dasatinib) when indicated. Fluid Retention ; Fluid retention, sometimes severe including pleural effusions Manage with supportive care measures and/or dose modification Side-effect: Most common adverse reactions (≥15%) in patients with newly diagnosed chronic phase CML included myelosuppression, fluid retention, and diarrhea. Most common adverse reactions (≥15%) in patients with resistance or intolerance to prior imatinib therapy included myelosuppression, fluid retention events, diarrhea, headache, fatigue, dyspnea, skin rash nausea, hemorrhage and musculoskeletal pain.	100mg Tablet	USFDA	Abjygv`b Kiv †h‡Z cv‡i	Abjygv`b Kiv nj
28.	Incepta Pharmaceuticals LTd (Dhamrai Unit)	Enzalutamide 40 mg Soft Gelatin Capsule Enzalutamide INN 40 mg Anti Cancer	Enzalutamide is indicated for the treatment of patients with metastatic castration-resistant prostate cancer.	Contraindication: Pregnancy Side effect: The most common adverse reactions (≥ 10%) are asthenia/fatigue, back pain, decreased appetite, constipation, arthralgia, diarrhea, hot flush, upper respiratory tract infection, peripheral edema, dyspnea, musculoskeletal pain, weight decreased, headache, hypertension, and dizziness/vertigo.	New	USFDA	Abjygv`b Kiv †h‡Z cv‡i	Abjygv`b Kiv nj
29.	Beacon Pharmaceutical Ltd., Incepta Pharmaceuticals Ltd.	Ibrutinib 140 mg Capsule Ibrutinib INN 140 mg Anticancer	Ibrutinib is indicated for the treatment of patients with : • mantle cell lymphoma (MCL) who have received at least one prior therapy.Accelerated approval was granted for this indication based on overall response rate. Continued approval for this indication may be contingent upon verification of	Contraindication: None Side effect: The most common adverse reactions (≥25%) in patients with B-cell malignancies (MCL, CLL, WM) were thrombocytopenia, neutropenia, diarrhea, anemia, fatigue, musculoskeletal pain, bruising, nausea, upper respiratory tract infection, and rash.	New	USFDA	Abjygv`b Kiv †h‡Z cv‡i	Abjygv`b Kiv nj

			clinical benefit in confirmatory trials. • Chronic Lymphocytic Leukemia Chronic Lymphocytic Leukemia With 17p deletion • Ibrutinib is indicated for the treatment of patients with chronic lymphocytic leukemia (CLL) with 17p deletion • Ibrutinib is indicated for the treatment of patients with Waldenström's macroglobulinemia (WM)					
30.	Incepta Pharmaceuticals Ltd.	Idelalisib 100 mg Tablet Idelalisib INN 100 mg Anticancer	Idelalisib is a kinase inhibitor indicated for the treatment of patients with: •Relapsed chronic lymphocytic leukemia (CLL), in combination with rituximab, in patients for whom rituximab alone would be considered appropriate therapy due to other co-morbidities. •Relapsed follicular B-cell non-Hodgkin lymphoma (FL) in patients who have received at least two prior systemic therapies. •Relapsed small lymphocytic lymphoma (SLL) in patients who have received at least two prior systemic therapies. Accelerated approval was granted for FL and SLL based on overall response rate. Improvement in patient survival or disease related symptoms has not been established. Continued approval for these indications may be contingent upon verification of clinical benefit in confirmatory trials.	Contraindication: History of serious allergic reactions including anaphylaxis and toxic epidermal necrolysis Side effect: The most common adverse reactions (incidence ≥20%) are diarrhea, pyrexia, fatigue, nausea, cough, pneumonia, abdominal pain, chills, and rash. The most common laboratory abnormalities (incidence ≥30%) are neutropenia, hypertriglyceridemia, hyperglycemia, ALT elevations, and AST elevations.	New	USFDA	Abÿgv`b Kiv †h‡Z cv‡i	Abÿgv`b Kiv nj
31.	Beacon Pharmaceutical Ltd., Incepta Pharmaceuticals Ltd.	Idelalisib 150 mg Tablet Idelalisib INN 150 mg Anticancer	Idelalisib is a kinase inhibitor indicated for the treatment of patients with: •Relapsed chronic lymphocytic leukemia (CLL), in combination with rituximab, in patients for whom	Contraindication: History of serious allergic reactions including anaphylaxis and toxic epidermal necrolysis Side effect: The most common adverse reactions (incidence ≥20%) are diarrhea, pyrexia, fatigue, nausea, cough,	New	USFDA	Abÿgv`b Kiv †h‡Z cv‡i	Abÿgv`b Kiv nj

			rituximab alone would be considered appropriate therapy due to other co-morbidities. •Relapsed follicular B-cell non-Hodgkin lymphoma (FL) in patients who have received at least two prior systemic therapies. •Relapsed small lymphocytic lymphoma (SLL) in patients who have received at least two prior systemic therapies. Accelerated approval was granted for FL and SLL based on overall response rate. Improvement in patient survival or disease related symptoms has not been established. Continued approval for these indications may be contingent upon verification of clinical benefit in confirmatory trials.	pneumonia, abdominal pain, chills, and rash. The most common laboratory abnormalities (incidence ≥30%) are neutropenia, hypertriglyceridemia, hyperglycemia, ALT elevations, and AST elevations.				
32.	Ziska Pharmaceuticals Ltd.	Megestrol Acetate Micronised 160 mg Tablet Megestrol Acetate Micronised USP 160 mg Anticancer	It is indicated for the palliative treatment of advanced carcinoma of the breast (i.e. recurrent, inoperable or metastatic diseases). It should not be used in lieu of currently accepted procedures such as surgery, radiation or chemotherapy.	Contraindications: History of hypersensitivity to megestrol acetate or any component of the formulation ,Known or suspected pregnancy. Side-effects: The most common adverse events occurring in > 5% of all patients receiving 800mg/20mL of megestrol acetate oral suspension in the two clinical efficacy trials were nausea, diarrhea, impotence, rash, flatulence, hypertension, and asthenia	40 mg Tablet		*Abjgv`b Kiv †h‡Z cv‡i |*	*Abjgv`b Kiv nj |*
33.	Ziska Pharmaceuticals Ltd.	Megestrol Acetate Micronised 20 mg Tablet Megestrol Acetate Micronised USP 20 mg Anticancer	It is indicated for the palliative treatment of advanced carcinoma of the breast (i.e. recurrent, inoperable or metastatic diseases). It should not be used in lieu of currently accepted procedures such as surgery, radiation or chemotherapy.	Contraindications: History of hypersensitivity to megestrol acetate or any component of the formulation ,Known or suspected pregnancy. Side-effects: The most common adverse events occurring in > 5% of all patients receiving 800mg/20mL of megestrol acetate oral suspension in the two clinical efficacy trials were nausea, diarrhea, impotence, rash, flatulence, hypertension, and asthenia	40 mg Tablet	USFDA	*Abjgv`b Kiv †h‡Z cv‡i |*	*Abjgv`b Kiv nj |*
34.	Ziska Pharmaceuticals Ltd.	Megestrol Acetate Micronised 40 mg/ml oral Suspension Megestrol Acetate Micronised	This oral suspension is a progestin indicated for the treatment of anorexia, cachexia, or an unexplained significant weight loss	Contraindications: History of hypersensitivity to megestrol acetate or any component of the formulation ,Known or suspected pregnancy.	40 mg Tablet	USFDA	*Abjgv`b Kiv †h‡Z cv‡i |*	*Abjgv`b Kiv nj |*

		USP 40mg/ml Anticancer	in patients with a diagnosis of acquired immunodeficiency syndrome (AIDS)	Side-effects: The most common adverse events occurring in > 5% of all patients receiving 800mg/20mL of megestrol acetate oral suspension in the two clinical efficacy trials were nausea, diarrhea, impotence, rash, flatulence, hypertension, and asthenia				
35.	Beacon Pharmaceutical Ltd.,	Osimertinib 40 mg Tablet Osimertinib Mesylate INN 47.700mg eq to Osimertinib 40mg Tablet Anticancer	Osimertinib is a kinase inhibitor indicated for the treatment of patients with metastatic epidermal growth factor receptor (EGFR) T790M mutation-positive non-small cell lung cancer (NSCLC), as detected by an FDA-approved test, who have progressed on or after EGFR TKI therapy. This indication is approved under accelerated approval based on tumor response rate and duration of response. Continued approval for this indication may be contingent upon verification and description of clinical benefit in confirmatory trials.	Contraindication: None Side effect: Most common adverse reactions (≥25%) were diarrhea, rash, dry skin, and nail toxicity.	New	USFDA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
36.	Incepta Pharmaceuticals Ltd. Beacon Pharmaceutical Ltd.,	Osimertinib 80 mg Tablet Osimertinib Mesylate INN 95.40mg eq. to Osimertinib 80mg Anticancer	Osimertinib is a kinase inhibitor indicated for the treatment of patients with metastatic epidermal growth factor receptor (EGFR) T790M mutation-positive non-small cell lung cancer (NSCLC), as detected by an FDA-approved test, who have progressed on or after EGFR TKI therapy. This indication is approved under accelerated approval based on tumor response rate and duration of response. Continued approval for this indication may be contingent upon verification and description of clinical benefit in confirmatory trials.	Contraindication: None Side effect: Most common adverse reactions (≥25%) were diarrhea, rash, dry skin, and nail toxicity.	New	USFDA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
37.	Incepta Pharmaceuticals Ltd.	Pembrolizumab 25mg/Vial Injection	Pembrolizumab is a programmed death receptor-1 (PD-1)-blocking antibody indicated for the treatment	Contraindication: None Side effect: Most common adverse reactions (reported in ≥20% of patients)	New		Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj

		Pembrolizumab (Ready to fill sterile solution) INN 1.0ml containg Pembrolizumab 25mg/Vial Antineoplastic Agent	of: patients with unresectable or metastatic melanoma. patients with metastatic NSCLC whose tumors express PD-L1 as determined by an FDA-approved test and who have disease progression on or after platinum-containing chemotherapy. Patients with EGFR or ALK genomic tumor aberrations should have disease progression on FDA-approved therapy for these aberrations prior to receiving Pembrolizumab. This indication is approved under accelerated approval based on tumor response rate and durability of response. An improvement in survival or disease-related symptoms has not yet been established. Continued approval for this indication may be contingent upon verification and description of clinical benefit in the confirmatory trials.	with: melanoma included fatigue, pruritus, rash, constipation, diarrhea, nausea, and decreased appetite. NSCLC included fatigue, decreased appetite, dyspnea and cough.				
38.	Incepta Pharmaceuticals Ltd.	Pembrolizumab 50 mg/Vial Injection Pembrolizumab (Ready to fill sterile solution) INN 2.0ml containg Pembrolizumab 50mg/Vial Anticancer	Pembrolizumab is a programmed death receptor-1 (PD-1)-blocking antibody indicated for the treatment of: •patients with unresectable or metastatic melanoma. •patients with metastatic NSCLC whose tumors express PD-L1 as determined by an FDA-approved test and who have disease progression on or after platinum-containing chemotherapy. Patients with EGFR or ALK genomic tumor aberrations should have disease progression on FDA-approved therapy for these aberrations prior to receiving Pembrolizumab. This indication is approved under	Contraindication: None Side effect: Most common adverse reactions (reported in ≥20% of patients) with: melanoma included fatigue, pruritus, rash, constipation, diarrhea, nausea, and decreased appetite. NSCLC included fatigue, decreased appetite, dyspnea and cough.	New	USFDA	*Abjgv` b Kiv †h‡Z cv‡i |*	*Abjgv` b Kiv nj |*

			accelerated approval based on tumor response rate and durability of response. An improvement in survival or disease-related symptoms has not yet been established. Continued approval for this indication may be contingent upon verification and description of clinical benefit in the confirmatory trials.					
39.	Beacon Pharmaceutical Ltd.,	Pembrolizumab INN 50mg/2.0ml Vial Lyophilized Powder for Injection Pembrolizumab INN 50mg/ 2.0ml Anticancer	Pembrolizumab is a programmed death receptor-1 (PD-1)-blocking antibody indicated for the treatment of: •patients with unresectable or metastatic melanoma. •patients with metastatic NSCLC whose tumors express PD-L1 as determined by an FDA-approved test and who have disease progression on or after platinum-containing chemotherapy. Patients with EGFR or ALK genomic tumor aberrations should have disease progression on FDA-approved therapy for these aberrations prior to receiving Pembrolizumab. This indication is approved under accelerated approval based on tumor response rate and durability of response. An improvement in survival or disease-related symptoms has not yet been established. Continued approval for this indication may be contingent upon verification and description of clinical benefit in the confirmatory trials.	Contraindication: None Side effect: Most common adverse reactions (reported in ≥20% of patients) with: melanoma included fatigue, pruritus, rash, constipation, diarrhea, nausea, and decreased appetite. NSCLC included fatigue, decreased appetite, dyspnea and cough.	New	USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
40.	ACI Ltd., Narayanganj	Lacosamide 1.00 gm/100 ml oral solution Lacosamide INN 1.00gm/100 ml	Lacosamide is indicated as monotherapy or adjunctive therapy in patients with partial-onset seizures; Lacosamide Injection is indicated as short term replacement when oral administration is not	Contraindication: None Side-effects: Monotherapy: Most common adverse reactions are similar to those seen in adjunctive therapy studies	New	USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

		Anti convulsant	feasible	Adjunctive therapy: Most common adverse reactions (≥10% and greater than placebo) are diplopia, headache, dizziness, nausea				
41.	Square Pharmaceuticals Ltd., Pabna Unit, Salgaria, Pabna	Fluoxetine 90mg Delayed Release Capsule Fluoxetine Hydrochloride Enteric Coated Pellets (30% W/W) Ph. Grade 300.00mg eq. to 90mg Fluoxetine USP **Antidepressant**	It is indicated for ► acute and maintenance treatment of Major Depressive Disorde ► acute and maintenance treatment of obsessions and compulsions in patients with Obsessive Compulsive Disorder (OCD) ► Acute and maintenance treatment of binge-eating and vomiting behaviors in patients with moderate to severe Bulimia Nervosa ► Acute treatment of Panic Disorder, with or without agoraphobia	Fluoxetine is contraindicated in patients known to be hypersensitive to it. Monoamine oxidase inhibitors --There have been reports of serious, sometimes fatal, reactions (including hyperthermia, rigidity, myoclonus, autonomic instability with possible rapid fluctuations of vital signs, and mental status changes that include extreme agitation progressing to delirium and coma) in patients receiving fluoxetine in combination with a monoamine oxidase inhibitor (MAOI), and in patients who have recently discontinued fluoxetine and are then started on an MAOI. Some cases presented with features resembling neuroleptic malignant syndrome. Therefore, Fluoxetine should not be used in combination with an MAOI, or within a minimum of 14 days of discontinuing therapy with an MAOI. Since fluoxetine and its major metabolite have very long elimination half-lives, at least 5 weeks [perhaps longer, especially if fluoxetine has been prescribed chronically and/or at higher doses should be allowed after stopping Fluoxetine before starting an MAOI. Thioridazine -- Thioridazine should not be administered with Fluoxetine or within a minimum of 5 weeks after Fluoxetine has been discontinued	20mg Capsule	USFDA	*c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i|*	*c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv* হল|
42.	Square Pharmaceuticals Ltd., Pabna Unit, Salgaria, Pabna	Pizotifen 5mg/100ml Syrup Pizotifen Malate BP 7.270mg eq. to 5mg Pizotifen/100ml	Anorexia in underweight patients, mood elevation in the elderly, prophylactic (interval) treatment of migraine.	Contraindication: Known hypersensitivity to Pizotifen or any of the excipients. Side effects: Sedation, dizziness, dry mouth, constipation.	0.5mg Tablet & 1.5mg Tablet		*c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i|*	*c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv* হল|

		Antidepressant						
43.	Healthcare Pharmaceutical ltd., Rajendrapur, Gazipur	Canagliflozin 300mg Tablet Canagliflozin Hemihydrate INN 306.0mg eq. to Canagliflozin 300mg **Antidiabetic**	It is a sodium-glucose co-transporter 2 (SGLT2) inhibitor indicated as an adjunct to diet and exercise to improve glycemic control in adults with type 2 diabetes mellitus. Limitation of Use: • Not for treatment of type-1 diabetes mellitus or diabetic ketoacidosis	Contraindications: • History of a serious hypersensitivity reaction to canagliflozin • Patients having end stage renal disease or patients on dialysis Side effects: Most common adverse reactions associated with Canagliflozin (5% or greater incidence): female genital mycotic infections, urinary tract Infection and increased urination.	100mg Tablet	USFDA	ক্ষতিকর পার্শ্বপতিক্রিয়া থাকায় *c‡qvRb †bB wea|q Av‡e`b bv gÄy Ki| †h‡Z cv‡i|*	ক্ষতিকর পার্শ্বপতিক্রিয়া থাকায় *c‡qvRb †bB wea|q Av‡e`b bv gÄy Ki| হল|*
44.	Beximco Pharmaceuticals Ltd., Tongi ,Gazipur	Empagliflozin 10.00mg Tablet Empagliflozin INN 10.00mg **Antidiabetic**	Empagliflozin is indicated in the treatment of type 2 diabetes mellitus to improve glycaemic control.	Contraindications: Hypersensitivity to the active substance or to any of the excipients. Side effects: Hypotension, Impairment in Renal Function, Impaired Hepatic Function, Hypoglycemia with Concomitant Use with Insulin and Insulin Secretagoguese, UTI.	New	USFDA	ক্ষতিকর পার্শ্বপতিক্রিয়া থাকায় *c‡qvRb †bB wea|q Av‡e`b bv gÄy Ki| †h‡Z cv‡i|*	ক্ষতিকর পার্শ্বপতিক্রিয়া থাকায় *c‡qvRb †bB wea|q Av‡e`b bv gÄy Ki| হল|*
45.	ACI Ltd., Narayanganj	Exenatide 40 mg eqv. to Exenatide 2 mg /vial Extended Release Injectable Sterile Ready to fill powder for suspension Exenatide INN 40 mg eqv. to Exenatide INN 2 mg)/vial **Anti Diabetic**	Exenatide is a glucagon-like peptide-1 (GLP-1) receptor agonist indicated as an adjunct to diet and exercise to improve glycemic control in adults with type 2 diabetes mellitus Exenatide is an extended-release formulation of exenatide. Do not coadminister with Exenatide prefilled pen. . Important Limitations of Use Not recommended as first-line therapy for patients inadequately controlled on diet and exercise Should not be used to treat type 1 diabetes or diabetic ketoacidosis Use with insulin has not been studied and is not recommended Has not been studied in patients with a history of pancreatitis. Consider other antidiabetic	Contraindication: Exenatide is contraindicated in patients with personal or family history of medullary thyroid carcinoma or in patients with Multiple Endocrine Neoplasia syndrome type 2 Exenatide is contraindicated in patients with a prior serious hypersensitivity reaction to exenatide or any of the product components Side-effects: Most common (≥5%) and occurring more frequently than comparator in clinical trials: nausea, diarrhea, headache, vomiting, constipation, injectionsite pruritus, injection-site nodule, and dyspepsia. Warning: Risk of thyroid c-cell tumors • Exenatide extended-release causes thyroid C-cell tumors at clinically relevant exposures in rats. It is unknown whether It	New	USFDA	*Abyjgv`b Ki| ‡h‡Z cv‡i|*	*Abyjgv`b Ki| nj|*

			therapies in patients with a history of pancreatitis	causes thyroid C-cell tumors, including medullary thyroid carcinoma (MTC) in humans, as the human relevance of exenatide extended-release-induced rodent thyroid C-cell tumors has not been determined. • It is contraindicated in patients with a personal or family history of MTC or in patients with Multiple Endocrine Neoplasia syndrome type 2 (MEN 2). Counsel patients regarding the potential risk of MTC and the symptoms of thyroid tumors.				
46.	Beximco Pharmaceuticals Ltd., Tongi ,Gazipur Sun Pharmaceutical (Bangladesh) Ltd.	Voglibose 0.2mg Tablet Voglibose INN 0.2mg **Antidiabetic**	Voglibose is indicated as an adjustment to diet and exercise to improve glycaemic control in patients with NIDDM (non-insulin dependent diabetes mellitus). Also, to improve postprandial hyperglycemia in patient with type 2 Diabetes.	Contraindications: Hypersensitivity to Voglibose or to any of the excipients, Diabetic ketoacidosis, diabetic pre-coma, Severe infection, before and after operation or with serious trauma. Side effects: Diarrhoea, loose stools, abdominal pain, constipation, anorexia, nausea, vomiting etc.			*Abyjgv` b Kiv †h‡Z cv‡i |*	*Abyjgv` b Kiv nj |*
47.	Sun Pharmaceutical (Bangladesh) Ltd.	Voglibose 0.3 mg Tablet Voglibose INN 0.3mg **Antdiabetic Drug**	To improve postprandial hyperglycemia in patient with type 2 Diabetes.	Contraindications: Hypersensitivity to Voglibose or to any of the excipients, Diabetic ketoacidosis, diabetic pre-coma, Severe infection, before and after operation or with serious trauma. Side effects: Diarrhoea, loose stools, abdominal pain, constipation, anorexia, nausea, vomiting etc.	New		*Abyjgv` b Kiv †h‡Z cv‡i |*	*Abyjgv` b Kiv nj |*
48.	ACI Ltd., Narayanganj	Eluxadoline 100 mg Film Coated Tablet Eluxadoline INN 100 mg Anti diarrheal	Eluxadoline is indicated in adults for the treatment of irritable bowel syndrome with diarrhea (IBS-D).	Contraindication: Eluxadoline is contraindicated in patients with: Known or suspected biliary duct obstruction; or sphincter of Oddi disease or dysfunction. These patients are at increased risk for sphincter of Oddi spasm.	New	USFDA	*Abyjgv` b Kiv †h‡Z cv‡i |*	*Abyjgv` b Kiv nj |*

				Alcoholism, alcohol abuse or alcohol addiction, or in patients who drink more than 3 alcoholic beverages per day. These patients are at increased risk for acute pancreatitis. A history of pancreatitis; or structural diseases of the pancreas, including known or suspected pancreatic duct obstruction. These patients are at increased risk for acute pancreatitis. • Severe hepatic impairment (Child-Pugh Class C). These patients are at risk for significantly increased plasma concentrations of Eluxadoline. • A history of chronic or severe constipation or sequelae from constipation, or known or suspected mechanical gastrointestinal obstruction. These patients may be at risk for severe complications of bowel obstruction. Side effects: Most common adverse reactions are constipation, nausea and abdominal pain.				
49.	Acme Laboratories Ltd. Globe Pharmaceuticals Ltd., Noakhali Healthcare Pharmaceuticals Limited, Gazipur Ziska Pharmaceuticals Ltd. Beacon Pharmaceuticals Ltd. Julphar Bangladesh Ltd., Sreepur, Gazipur, Dhaka	Eluxadoline 100 mg Tablet Eluxadoline INN 100 mg Antidiarrheal	Eluxadoline is indicated in adults for the treatment of irritable bowel syndrome with diarrhea (IBS-D).	Contraindication: Eluxadoline is contraindicated in patients with: Known or suspected biliary duct obstruction; or sphincter of Oddi disease or dysfunction. These patients are at increased risk for sphincter of Oddi spasm. Alcoholism, alcohol abuse or alcohol addiction, or in patients who drink more than 3 alcoholic beverages per day. These patients are at increased risk for acute pancreatitis. A history of pancreatitis; or structural diseases of the pancreas, including known or suspected pancreatic duct obstruction. These patients are at increased risk for acute pancreatitis. • Severe hepatic impairment (Child-Pugh Class C). These patients are at risk for significantly increased plasma concentrations of Eluxadoline. • A history of chronic or severe constipation	New	USFDA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj

				or sequelae from constipation, or known or suspected mechanical gastrointestinal obstruction. These patients may be at risk for severe complications of bowel obstruction. Side effects: Most common adverse reactions are constipation, nausea and abdominal pain				
50.	ACI Ltd., Narayanganj Globe Pharmaceuticals Ltd., Noakhali Ziska Pharmaceuticals Ltd. Healthcare Pharmaceuticals Limited, Gazipur	Eluxadoline 75 mg FC Tablet Eluxadoline INN 75 mg Anti diarrheal	Eluxadoline is a mu-opioid receptor agonist, indicated in adults for the treatment of irritable bowel syndrome with diarrhea (IBS-D).	Contraindication: Eluxadoline is contraindicated in patients with: Known or suspected biliary duct obstruction; or sphincter of Oddi disease or dysfunction. These patients are at increased risk for sphincter of Oddi spasm. Alcoholism, alcohol abuse or alcohol addiction, or in patients who drink more than 3 alcoholic beverages per day. These patients are at increased risk for acute pancreatitis. A history of pancreatitis; or structural diseases of the pancreas, including known or suspected pancreatic duct obstruction. These patients are at increased risk for acute pancreatitis. • Severe hepatic impairment (Child-Pugh Class C). These patients are at risk for significantly increased plasma concentrations of Eluxadoline. • A history of chronic or severe constipation or sequelae from constipation, or known or suspected mechanical gastrointestinal obstruction. These patients may be at risk for severe complications of bowel obstruction. Side effects: Most common adverse reactions are constipation, nausea and abdominal pain	New	USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
51.	Incepta Pharmaceuticals Ltd.	Aprepitant 125 mg/Sachet Powder for Oral suspension Aprepitant INN 125mg/Sachet Antiemetic	Aprepitant for oral suspension is indicated • in combination with other antiemetic agents, in patients 12 years of age and older for prevention of: o acute and delayed nausea and vomiting associated with initial and	Contraindication: Known hypersensitivity to any component of this drug. Concurrent use with pimozide. Side effect: Most common adverse reactions are: Prevention of Chemotherapy Induced Nausea and Vomiting (CINV) Adults (≥3%): fatigue, diarrhea, asthenia, dyspepsia, abdominal pain, hiccups, white	40mg, 125mg Capsule	USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

			repeat courses of highly emetogenic cancer chemotherapy (HEC) including high-dose cisplatin o nausea and vomiting associated with initial and repeat courses of moderately emetogenic cancer chemotherapy (MEC) Limitations of Use: Aprepitant has not been studied for treatment of established nausea and vomiting. Chronic continuous administration of Aprepitant is not recommended.	blood cell count decreased, dehydration, and alanine aminotransferase increased. • Pediatrics (≥3%): neutropenia, headache, diarrhea, decreased appetite, cough, fatigue, hemoglobin decreased, dizziness, and hiccups. PONV Adults (≥3%): constipation and hypotension.				
52.	Incepta Pharmaceuticals Ltd.	Aprepitant 80 mg Capsule Aprepitant INN 80 mg Antiemetic	Aprepitant Capsule is indicated • in combination with other antiemetic agents, in patients 6 months of age and older for prevention of: ➢ acute and delayed nausea and vomiting associated with initial and repeat courses of highly emetogenic cancer chemotherapy (HEC) including high-dose cisplatin. ➢ nausea and vomiting associated with initial and repeat courses of moderately emetogenic cancer chemotherapy (MEC). • for prevention of postoperative nausea and vomiting (PONV) in adults Limitations of Use: Aprepitant has not been studied for treatment of established nausea and vomiting. Chronic continuous administration of Aprepitant is not recommended.	Contraindication: Known hypersensitivity to any component of this drug. Concurrent use with pimozide. Side effect: Most common adverse reactions are: Prevention of Chemotherapy Induced Nausea and Vomiting (CINV) Adults (≥3%): fatigue, diarrhea, asthenia, dyspepsia, abdominal pain, hiccups, white blood cell count decreased, dehydration, and alanine aminotransferase increased. • Pediatrics (≥3%): neutropenia, headache, diarrhea, decreased appetite, cough, fatigue, hemoglobin decreased, dizziness, and hiccups. PONV Adults (≥3%): constipation and hypotension.	40mg, 125mg Capsule	USFDA	Abyjgv`b Kiv †h‡Z cv‡i|	Abyjgv`b Kiv nj|
53.	Opsonin Pharma Limited, Bagura Road, Barisal	Domperidone 30mg Sustained Release Capsule Domperidone BP 30 mg	Stimulation of gut motility: Non–ulcer dyspepsia, Esophageal reflux and gastritis, Diabetic gastroparesis, functional dyspepsia, prevention	Contraindications: Domperidone is contraindicated to the patients who have hypersensitivity to this drug and in case of neonates.	10 mg tablet, 15 mg & 30 mg suppository, 60 ml suspension,		c‡qvRbxq †idv‡iY †bB wea|q Av‡e`b bv gÄyiK iv †h‡Z cv‡i|	c‡qvRbxq †idv‡iY bv_vKq Av‡e`b bvgÄyiK iv হল|

		Gastro-prokinetics	and symptomatic relief of acute nausea and vomiting from any cause but specifically cytotoxic therapy, radio therapy and anti-parkinsonism therapy, management of migraine	Side effects: Domperidone may produce hyperprolactinemia. This may result galactorrhoea, breast enlargement and soreness and reduced libido. Dry mouth, thirst headache, nervousness, drowsiness, diarrhea, skin rash and itching may occur during treatment with Domperidone. Extra-pyramidal reactions are seen in 0.055 of patients in clinical studies.	15 ml paediatric drop			
54.	Ziska Pharmaceuticals Ltd.	Fosaprepitant 115mg/Vial Sterile Lyophilized Powder for Injection Fosaprepitant Dimeglumine 188.0 mg eq. to Fosaprepitant INN 115 mg/Vial Adjunctive antiemetic	This injection is a substance P/neurokinin-1 (NK 1) receptor antagonist, indicated in adults, in combination with other antiemetic agents, for the prevention of acute and delayed nausea and vomiting associated with initial and repeat courses of highly emetogenic cancer chemotherapy (HEC) including high-dose cisplatin delayed nausea and vomiting associated with initial and repeat courses of moderately emetogenic cancer chemotherapy (MEC) Limitations of Use EMEND has not been studied for treatment of established nausea and vomiting.	Contraindications: Known hypersensitivity to any component of this drug, Concurrent use with pimozide. Side effects: Most common adverse reactions (≥2%) are: fatigue, diarrhea, neutropenia, asthenia, anemia, peripheral neuropathy, leukopenia, dyspepsia, urinary tract infection, pain in extremity.	Aprepitant 40 mg Capsule	USFDA	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
55.	Ziska Pharmaceuticals Ltd. Incepta Pharmaceuticals Ltd.	Lyophilized Powder for Injection Fosaprepitant Dimeglumine 245.30mg eq. to Fosaprepitant INN 150mg/Vial Adjunctive antiemetic	This injection is a substance P/neurokinin-1 (NK 1) receptor antagonist, indicated in adults, in combination with other antiemetic agents, for the prevention of acute and delayed nausea and vomiting associated with initial and repeat courses of highly emetogenic cancer chemotherapy (HEC) including high-dose cisplatin delayed nausea and vomiting associated with initial and repeat courses of moderately emetogenic cancer chemotherapy (MEC) Limitations of Use EMEND has not been studied for treatment of established nausea and vomiting.	Contraindications: Known hypersensitivity to any component of this drug, Concurrent use with pimozide. Side effects: Most common adverse reactions (≥2%) are: fatigue, diarrhea, neutropenia, asthenia, anemia, peripheral neuropathy, leukopenia, dyspepsia, urinary tract infection, pain in extremity.	Fosaprepitant 115mg/Vial Injection	USFDA	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
56.	Navana Pharmaceuticals Ltd.	Hyoscine Hydrobromide 300mcg Tablet	For the prevention of travel sickness.	Contraindications: Prostatic enlargement, paralytic ileus,	150 mcg	BNF-70 Page-344	*GB gvÎv c‡qvRb †bB wea|q Av‡e`b bv gÄyi Kiv ‡h‡Z*	*GB gvÎv c‡qvRb †bB wea|q Av‡e`b bvgÄyi Kiv nj |*

		Hyoscine Hydrobromide INN 300 mcg Antiemetic		pyloric stenosis, glaucoma and myasthenia gravis. In addition, Kwells should not be given to patients with a known sensitivity to hyoscine hydrobromide or any other component of the product. Side Effects: The listed adverse drug reactions are based on spontaneous reports, thus an organization according to CIOMS II categories of frequency is not pertinent. General: hyperthermia at high temperatures due to decreased sweating. Eye disorders: blurred vision, mydriasis. Gastrointestinal disorders: dry mouth. Immune system disorders: allergic reaction and anaphylactic reaction. Hypersensitivity reactions with respective laboratory and clinical manifestations, including asthma syndrome, mild to moderate reactions affecting skin, respiratory tract, gastrointestinal tract, and cardiovascular system, and symptoms such as rash, urticaria, oedema, pruritus, cardio-respiratory distress, have been reported. Nervous system disorders: drowsiness, dizziness, sedation and somnolence are commonly reported. Central nervous system stimulation including restlessness, hallucinations and confusion, have been less frequently reported following the administration of hyoscine. There have been rare reports of an increase in seizure frequency in epileptic patients			cv‡i |	
57.	Incepta Pharmaceuticals Ltd.	Rolapitant 90mg Tablet Rolapitant HCl INN 100mg eq. to 90mg Rolapitant Antiemetic Agents	Rolapitant is a substance P/neurokinin 1 (NK1) receptor antagonist indicated in combination with other antiemetic agents in adults for the prevention of delayed nausea and vomiting associated with initial and repeat courses of emetogenic cancer chemotherapy, including, but not limited to, highly	Contraindication: Concurrent use with thioridazine, a CYP2D6 substrate. Side effect: Most common adverse reactions (≥ 5%) are: • Cisplatin Based Highly Emetogenic Chemotherapy: neutropenia and hiccups Moderately Emetogenic Chemotherapy and Combinations of Anthracycline and	New	USFDA	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |

			emetogenic chemotherapy	Cyclophosphamide: decreased appetite, neutropenia and dizziness.				
58.	Orion Pharma Ltd.	Brivaracetam 10 mg/ml Oral Solution Brivaracetam INN 10 mg/ml Anti-epileptic	For the treatment of partial-onset seizures in patients 16 years of age and older with epilepsy	Contraindications: Hypersensitivity to Brivaracetam or any of the inactive ingredients in Brivaracetam (bronchospasm and angioedema have occurred). Side effects: Like other antiepileptic drugs, Brivaracetam may cause suicidal thoughts or actions in a very small number of people, about 1 in 500 people taking it. Brivaracetam can also cause problems with balance and coordination. Mental (psychiatric) symptoms. Brivaracetam can cause mood and behavior changes such as aggression, agitation, anger, anxiety, apathy, mood swings, depression, hostility, and irritability. Irritability and anxiety are common with Brivaracetam, and can be severe. The most common side effects of Brivaracetam include: sleepiness, dizziness, feeling tired nausea and vomiting.	New	USFDA	Abjygv`b Kiv †h‡Z cv‡i	Abjygv`b Kiv nj
59.	Orion Pharma Ltd.	Brivaracetam 100 mg Tablet Brivaracetam INN 100 mg Anti-epileptic	-do-	-do-	New	USFDA	Abjygv`b Kiv †h‡Z cv‡i	Abjygv`b Kiv nj
60.	Orion Pharma Ltd.	Brivaracetam 75 mg Tablet Brivaracetam INN 75 mg Anti-epileptic	-do-	-do-	New	USFDA	Abjygv`b Kiv †h‡Z cv‡i	Abjygv`b Kiv nj

61.	Orion Pharma Ltd. Globe Pharmaceuticals Ltd., Noakhali	Brivaracetam 50 mg Tablet Brivaracetam INN 50 mg Anti-epileptic	-do-	-do-	New	USFDA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
62.	Orion Pharma Ltd.	Brivaracetam 25 mg Tablet Brivaracetam INN 25mg Anti-epileptic	-do-	-do-	New	USFDA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
63.	Beacon Pharmaceutical Ltd., Orion Pharma Ltd.	Brivaracetam 50mg/5ml Vial IV Injection Brivaracetam INN 50mg/5ml Antiepileptic	For the treatment of partial-onset seizures in patients 16 years of age and older with epilepsy	Contraindications: Hypersensitivity to Brivaracetam or any of the inactive ingredients in Brivaracetam (bronchospasm and angioedema have occurred). Side effects: Like other antiepileptic drugs, Brivaracetam may cause suicidal thoughts or actions in a very small number of people, about 1 in 500 people taking it. Brivaracetam can also cause problems with balance and coordination. Mental (psychiatric) symptoms. Brivaracetam can cause mood and behavior changes such as aggression, agitation, anger, anxiety, apathy, mood swings, depression, hostility, and irritability. Irritability and anxiety are common with Brivaracetam, and can be severe. The most common side effects of Brivaracetam include: sleepiness, dizziness, feeling tired nausea and vomiting.	New	USFDA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
64.	Concord Pharmaceuticals Ltd.	Dapagliflozin 10 mg Tablet Dapagliflozin INN 10mg Anti-diabetic	Dapagliflozin is indicated as an adjunct to diet and exercise to improve glycemic control in adults with type 2 diabetes mellitus.	Contraindicaion: History of a serious hypersensitivity reaction to Dapagliflozin. Severe renal impairment, end-stage renal disease (ESRD), or patients on dialysis Side effects: Gastrointestinal Perforations and Fistulae, Surgery and Wound Healing Complications, Hemorrhage, Proteinuria	Dapagliflozin 5mg Tablet	USFDA BNF-70 Page No-610	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
65.	Concord Pharmaceuticals Ltd.	Voglibose 0.2 mg Tablet Voglibose INN 0.2 mg Anti-diabetic	Voglibose is indicated for the treatment of diabetes. It is specifically used for lowering post-prandial blood glucose levels thereby reducing the risk of	Contraindications: Inflammatory bowel disease; GI obstruction or patients predisposed to it; conditions which may deteriorate as a result of increased gas formation e.g., hernia; severe ketosis;	New		Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj

			macrovascular complications	diabetic coma or pre-coma; severe infection; hypersensitivity; pregnancy; lactation. Not to be used as monotherapy in IDDM. Side effects: Flatulence; abdominal distension; diarrhoea; pain; skin reactions; hypoglycemia; increased LFT. Potentially Fatal: Hepatotoxicity.				
66.	Square Pharmaceuticals Ltd., Pabna Unit, Salgaria, Pabna	Butoconazole Nitrate 2gm/100gm Vaginal Cream Butoconazole Nitrate USP 2gm/100gm Vaginal Cream **Antifungal**	Butoconazole 2% is indicated for the local treatment of vulvovaginal candidiasis (infections caused by Candida).	Contraindication: Butoconazole is contraindicated in patients with a history of hypersensitivity to any of the components of the product. ADR/Side effects:Of the 314 patients treated with Butoconazole for 1 day in controlled clinical trials, 18 patients (5.7%) reported complaints such as vulvar/vaginal burning, itching, soreness and swelling, pelvic or abdominal pain or cramping, or a combination of two or more of these symptoms. In 3 patients (1%) these complaints were considered treatment-related. Five of the 18 patients reporting adverse events discontinued the study because of them.	New	USFDA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
67.	Eskayef Bangladesh Limited.	Efinaconazole 10gm/100ml Topical solution Efinaconazole INN 10gm/100ml **Antifungal**	It is an azole antifungal indicated for the topical treatment of onychomycosis of the toenails due to Trichophyton rubrum and Trichophyton mentagrophytes.	Contraindications: None Side effects: The most common side effects (incidence >1%) were ingrown toenails, application site dermatitis, application site vesicles, and application site pain	New	USFDA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
68.	Beacon Pharmaceuticals Ltd.	Isavuconazole 100mg Capsule Isavuconazonium Sulfate INN 186.261mg eq. to Isavuconazole 100mg **Anifungal**	It is an azole antifungal indicated for use in the treatment of •Invasive aspergillosis •Invasive mucormycosis	Contraindication: •Hypersensitivity to Isavuconazole •Coadministration with strong CYP3A4 inhibitors, such as ketoconazole or highdose ritonavir •Coadministration with strong CYP3A4 inducers, such as rifampin, carbamazepine, St. John's wort, or long acting barbiturates •Use in patients with familial short QT syndrome Side-effect: Most frequent adverse	New	USFDA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj

				reactions: nausea, vomiting, diarrhea, headache, elevated liver chemistry tests , hypokalemia, constipation, dyspnea, cough, peripheral edema, and back pain				
69.	Beacon Pharmaceuticals Ltd.	Isavuconazole 200mg/Vial Injection Isavuconazonium Sulfate INN 372.522mg eq. to Isavuconazole 200mg/Vial **Anifungal**	It is an azole antifungal indicated for use in the treatment of •Invasive aspergillosis •Invasive mucormycosis	Contraindication: •Hypersensitivity to Isavuconazole •Coadministration with strong CYP3A4 inhibitors, such as ketoconazole or highdose ritonavir •Coadministration with strong CYP3A4 inducers, such as rifampin, carbamazepine, St. John's wort, or long acting barbiturates •Use in patients with familial short QT syndrome Side-effect: Most frequent adverse reactions: nausea, vomiting, diarrhea, headache, elevated liver chemistry tests , hypokalemia, constipation, dyspnea, cough, peripheral edema, and back pain	New	USFDA	*Abjygv`b Kiv †h‡Z cv‡i |*	*Abjygv`b Kiv nj |*
70.	Navana Pharmaceuticals Ltd.	Itraconazole 200mg Tablet Itraconazole USP 200mg Antifungal	It is an azole antifungal, are indicated for the treatment of onychomycosis of the toenail caused by Trichophyton rubrum or T. mentagrophytes.	Contraindications: Do not administer for the treatment of onychomycosis in patients with evidence of ventricular dysfunction such as congestive heart failure (CHF) or a history of CHF. Do not be administer for the treatment of onychomycosis to pregnant patients or to women contemplating pregnancy. Coadministration of cisapride, dofetilide, ergot alkaloids such as dihydroergotamine, ergotamine, ergometrine (ergonovine), and methylergometrine (methylergonovine), felodipine, levacetylmethadol (levomethadyl), lovastatin, methadone, oral midazolam, nisoldipine, pimozide, quinidine, simvastatin, and triazolam with Itraconazole 200mg Tablet is contraindicated. Anaphylaxis and hypersensitivity have been reported with use of itraconazole.	100mg Capsule	USFDA	*Abjygv`b Kiv †h‡Z cv‡i |*	*Abjygv`b Kiv nj |*

				Itraconazole 200mg Tablet is contraindicated in patients who have shown hypersensitivity to itraconazole products. Side Effects: Most common adverse reactions observed in the treatment phase of the onychomycosis clinical trial (>1%) are upper respiratory tract infections, increased hepatic enzymes, hypoacusis, headache, abdominal pain, diarrhea, nausea, fatigue, arrhythmia, cough, sore throat and back pain. Itraconazole has been associated with rare cases of serious hepatotoxicity, including liver failure and death				
71.	Eskayef Bangladesh Limited	Tavaborole 4.350gm/100ml Topical Solution Tavaborole INN 4.350 gm/ 100ml Antifungal	It is an oxaborole antifungal indicated for the topical treatment of onychomycosis of the toenails due to Trichophyton rubrum or Trichophyton mentagrophytes.	Contraindication: None Side effect: Common adverse reactions occurring in ≥1% in subjects treated with KERYDIN included application site exfoliation, ingrown toenail, application site erythema, and application site dermatitis.	New	USFDA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
72.	Jayson Pharmaceuticals Ltd.	Tavaborole 5% Topical Solution Tavaborole INN 5% Topical Antifungal	It is an oxaborole antifungal indicated for the topical treatment of onychomycosis of the toenails due to Trichophyton rubrum or Trichophyton mentagrophytes.	Contraindication: None Side effect: Common adverse reactions occurring in ≥1% in subjects treated with KERYDIN included application site exfoliation, ingrown toenail, application site erythema, and application site dermatitis.	New	USFDA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
73.	Eskayef Bangladesh Limited Globe Pharmaceuticals Ltd. Square Pharmaceuticals Ltd., Pabna Unit, Salgaria, Pabna	Bilastin 20mg Tablet Bilastin INN 20mg Antihistamine	It is indicated for the Symptomatic treatment of allergic rhino-conjunctivitis (seasonal and perennial) and urticaria	Contraindications: Hypersensitivity to the active substance bilastine or to any of the excipients. Side effects: Headache, Malaise; Less commonly abdominal pain, diarrhoea, increased appetite, weight gain, thirst, gastritis, prolongation of the QT interval, dyspnoea, anxiety, insomnia, vertigo, dizziness, pyrexia, oral herpes,, tinnitus.	New	BNF 70 Page: 244	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
74.	Acme Laboratories Ltd. Opsonin Pharma Limited, Bagura Road, Barisal	Cilnidipine 10 mg Tablet Cilnidipine INN 10 mg Anti-hypertensive	Cilnidipine is a calcium channel blocker indicated for patients at risk of vascular damage caused by hypertension. It helps to control hypertension (high blood pressure) by reduce the functioning of the body`s angiotensin receptors.	Contraindication: Cilnidipine is contraindicated in patients with severe aortic stenosis, cardiogenic shock, history of unstable angina or MI, heart failure and hypotension. It is also contraindicated in patients with known hypersensitivity to Cilnidipine or any other components of the product. Side effects: Possible side effects are	New		Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj

				nausea, headaches, trouble breathing, swelling, low blood pressure, edema (water retention).				
75.	Opsonin Pharma Limited, Bagura Road, Barisal	Cilnidipine 5 mg Tablet Cilnidipine INN 5 mg Anti-hypertensive	Cilnidipine is a calcium channel blocker indicated for patients at risk of vascular damage caused by hypertension. It helps to control hypertension (high blood pressure) by reduce the functioning of the body`s angiotensin receptors.	Contraindication: Cilnidipine is contraindicated in patients with severe aortic stenosis, cardiogenic shock, history of unstable angina or MI, heart failure and hypotension. It is also contraindicated in patients with known hypersensitivity to Cilnidipine or any other components of the product. Side effects: Possible side effects are nausea, headaches, trouble breathing, swelling, low blood pressure, edema (water retention).	New		Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
76.	Incepta Pharmaceuticals Ltd. Globe Pharmaceuticals Ltd., Noakhali Orion Pharma Ltd.	Tofacitinib 11 mg Extended Release Tablet Tofacitinib Citrate INN 17.60mg eq. to 11.0mg Tofacitinib Antirheumatic Drugs	iT is an inhibitor of Janus kinases (JAKs) indicated for the treatment of adult patients with moderately to severely active rheumatoid arthritis who have had an inadequate response or intolerance to methotrexate. It may be used as monotherapy or in combination with methotrexate or other nonbiologic disease-modifying antirheumatic drugs (DMARDs). •Limitations of Use: Use of Tofacitinib/Tofacitinib XR in combination with biologic DMARDs or potent immunosuppressants such as azathioprine and cyclosporine is not recommended.	Contraindication: None Side effect: The most commonly reported adverse reactions during the first 3months in controlled clinical trials (occurring in greater than or equal to 2% of patients treated with Tofacitinib monotherapy or in combination with DMARDs) were upper respiratory tract infections, headache, diarrhea and nasopharyngitis.	New	USFDA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
77.	Healthcare Pharmaceutical ltd., Rajendrapur, Gazipur Aristopharma Ltd.	Dexrabeprazole 10mg Tablet Dexrabeprazole Sodium INN 10.60mg eq. to Dexrabeprazole 10mg Anti Ulcerant	This medication is prescribed for gastro-esophageal reflux disease, gastric and duodenal ulcers.	Contraindications: It is contraindicated in patients with known hypersensitivity to dexrabeprazole, or to any excipients used in formulation. Side effects: Dexrabeprazole can cause side effects such as nausea, vomiting, diarrhoea, stomach pain and flatulence.	New		c‡qvRbxq †idv‡iY ‡bB wewfbœ Av‡e`b bv gÄyi Kiv ‡h‡Z cv‡i	c‡qvRbxq †idv‡iY bv_vKq Av‡e`b bvgÄyi Kiv হল
78.	Healthcare Pharmaceutical ltd., Rajendrapur, Gazipur	Dexrabeprazole 5 mg Tablet Dexrabeprazole Sodium INN	Gastro-esophageal Reflux Disease (GERD), erosive esophagitis, gastric and duodenal ulcers. It is more	Contraindications: It is contraindicated in patients with known hypersensitivity to dexrabeprazole, or to any excipients used	New		c‡qvRbxq †idv‡iY ‡bB wewfbœ Av‡e`b bv gÄyi Kiv ‡h‡Z cv‡i	c‡qvRbxq †idv‡iY bv_vKq Av‡e`b bvgÄyi Kiv হল

		5.30mg eq. to Dexrabeprazole 5.0mg Anti Ulcerant	effective and tolerable than rabeprazole.	in formulation. Side effects: Dexrabeprazole can cause side effects such as nausea, vomiting, diarrhoea, stomach pain and flatulence.				
79.	Healthcare Pharmaceutical ltd., Rajendrapur, Gazipur	Dexrabeprazole 10mg/IV Injection Dexrabeprazole Sodium INN 10.60mg eq. to Dexrabeprazole 10mg/vial Anti Ulcerant	Gastro-esophageal Reflux Disease (GERD), erosive esophagitis, gastric and duodenal ulcers. It is more effective and tolerable than rabeprazole.	Contraindications: It is contraindicated in patients with known hypersensitivity to dexrabeprazole, or to any excipients used in formulation. Side effects: Dexrabeprazole can cause side effects such as nausea, vomiting, diarrhoea, stomach pain and flatulence.	New		*c‡qvRbxq †idv‡iÝ ‡bB weavq Av‡e`b bv gÄyi Kiv ‡h‡Z cv‡i ।*	*c‡qvRbxq †idv‡iÝ bv _vKq Av‡e`b bvgÄyi Kiv* হল।
80.	Healthcare Pharmaceuticals Limited, Rajendrapur, Gazipur	Esomeprazole Magnesium (Trihydrate) USP 11.135 mg (Equivalent to Esomeprazole 10 mg) Sachat Esomeprazole Magnesium (Trihydrate) USP 11.135 mg (Equivalent to Esomeprazole 10 mg) Sachat Anti Ulcerant	Esomeprazole is a proton pump inhibitor indicated for the following: • Treatment of gastroesophageal reflux disease (GERD). • Risk reduction of NSAID-associated gastric ulcer. • H. pylorieradication to reduce the risk of duodenal ulcer recurrence. • Pathological hypersecretory conditions, including Zollinger-Ellison syndrome.	Contra-indication: Patients with known hypersensitivity to proton pump inhibitors (PPIs) (angioedema and anaphylaxis have occurred) Side-effects: Most common adverse reactions • Adults (≥18 years) (incidence>1%) are headache, diarrhea, nausea, flatulence, abdominal pain, constipation, and dry mouth. • Pediatric (1 to 17 years) (incidence>2%) are headache, diarrhea, abdominal pain, nausea, and somnolence Pediatric (1 month to less than 1 year) (incidence 1%) areabdominal pain, regurgitation, tachypnea, and increased ALT	Esomeprazole 20 mg , 40 mg Tablet,Capsule and 20 mg Sachat	USFDA	*c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv ‡h‡Z cv‡i ।*	*c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv* হল।
81.	Incepta Pharmaceuticals Ltd.	Esomeprazole 10mg/Sachet Delayed Release Granules for oral suspension Esomeprazole Magnesium Trihydrate BP 11.125mg eq. to Esomeprazole 10 mg delayed release Granules Anti Ulcerant	-do-	-do-	Esomeprazole 20mg DR Granules For Suspension	USFDA	*c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv ‡h‡Z cv‡i ।*	*c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv* হল।

82.	Eskayef Bangladesh Limited. Square Pharmaceuticals Ltd., Pabna Unit, Salgaria, Pabna ACI Ltd., Narayanganj	Rabeprazole Sodium 20mg/Vial Sterile Lyophilized Powder for Injection Rabeprazole Sodium INN 20mg/Vial **Antiulcerant**	Sequential-therapy (step-up) from oral rabeprazole, who is temporarily unable to take oral medication for any reason. Active duodenal ulcer and gastric ulcer with bleeding or severe erosions. Prevention of acid aspiration. Pathological hypersecretory conditions, including Zollinger-Ellison syndrome.	**Contraindications:** Hypersensitivity to rabeprazole or to any component of the formulation. **Side effects:** Headache, abdominal pain, diarrhea, dry mouth, dizziness, peripheral edema, hepatic enzyme increase, hepatitis, hepatic encephalopathy, myalgia, and arthralgia.	Rabeprazole 10mg & 20mg Tablet		c‡qvRbxq †idv‡iY ‡bB wea vq Av‡e`b bv gÄy Kiv ‡h‡Z cv‡i	c‡qvRbxq †idv‡iY bv_vKq Av‡e`b bvgÄy Kiv হল।
83.	Opsonin Pharma Limited, Bagura Road, Barisal	Rabeprazole sodium 20mg/10 ml Vial Injection Rabeprazole Sodium INN 21.22 mg eq. to Rabeprazole 20mg/10ml **Antiulcerant**	-do-	-do-	20 mg Tablet, 20 mg Capsule, 10 mg Tablet, 10 mg Capsule		c‡qvRbxq †idv‡iY ‡bB wea vq Av‡e`b bv gÄy Kiv ‡h‡Z cv‡i	c‡qvRbxq †idv‡iY bv_vKq Av‡e`b bvgÄy Kiv হল।
84.	ACI Ltd., Narayanganj	Rabeprazole Sodium 40 mg Tablet Rabeprazole Sodium INN 40mg **Antiulcerant**	-do-	-do-	10mg & 20 mg Tablet		c‡qvRbxq †idv‡iY ‡bB wea vq Av‡e`b bv gÄy Kiv ‡h‡Z cv‡i	c‡qvRbxq †idv‡iY bv_vKq Av‡e`b bvgÄy Kiv হল।
85.	Square Pharmaceuticals Ltd., Dhaka Unit, Kaliakoir, Gazipur	Olopatadine Hydrochloride 7.76mg eq. to 7.00mg/ml Olopatadine Opthalmic Solution Olopatadine Hydrochloride USP 7.76mg eq. to 7.00mg Olopatadine **Anti-Allergic**	Olopatadine is indicated for the treatment of ocular itching associated with allergic conjunctivitis.	Contraindications: None Side effects: Because clinical trials are conducted under widely varying conditions, adverse reaction rates observed in the clinical trials of a drug cannot be directly compared to rates in the clinical trials of another drug and may not reflect the rates observed in practice. These events were blurred vision, dry eye, superficial punctate keratitis, dysgeusia and abnormal sensation in eye.	1% and 0.2% Eye Drops 0.6% Nasal Spray	USFDA	Abygv`b Kiv ‡h‡Z cv‡i	Abygv`b Kiv nj।

86.	Healthcare Pharmaceutical ltd., Rajendrapur, Gazipur	Ceftibuten 200 mg Capsule Ceftibuten Dihydrate INN 217.54mg eq. to Ceftibuten 200mg **Antibiotic**	Ceftibuten is indicated for the treatment of individuals with mild-to-moderate infections caused by susceptible strains of the designated microorganisms in the specific conditions listed below Acute Bacterial Exacerbations of Chronic Bronchitis due to *Haemophilus influenzae* (including β-lactamase-producing strains), *Moraxella catarrhalis* (including β-lactamaseproducing strains), or *Streptococcus pneumoniae* (penicillin-susceptible strains only). NOTE: In acute bacterial exacerbations of chronic bronchitis clinical trials where *Moraxella catarrhalis* was isolated from infected sputum at baseline, ceftibuten clinical efficacy was 22% less than control. Acute Bacterial Otitis Media due to *Haemophilus influenzae* (including β-lactamase-producing strains), *Moraxella catarrhalis* (including β-lactamase-producing strains), or *Streptococcus pyogenes*. NOTE: Although ceftibuten used empirically was equivalent to comparators in the treatment of clinically and/or microbiologically documented acute otitis media, the efficacy against *Streptococcus pneumoniae* was 23% less than control. Therefore, ceftibuten should be given empirically only when adequate antimicrobial coverage against *Streptococcus pneumoniae* has been previously administered. Pharyngitis and Tonsillitis due to *Streptococcus pyogenes*. NOTE: Only penicillin by the intramuscular route of administration has been shown to be effective in	Contraindications: Ceftibuten is contraindicated in patients with known allergy to the Cephalosporin group of antibiotics. Side effects: Aphasia, jaundice, melena, psychosis, stridor, toxic epidermal necrolysis,serum ickness-like reactions and Stevens-Johnson syndrome.	400mg Capsule	USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

			the prophylaxis of rheumatic fever. Ceftibuten is generally effective in the eradication of *Streptococcus pyogenes* from the oropharynx; however, data establishing the efficacy of the CEDAX product for the prophylaxis of subsequent rheumatic fever are not available.					
87.	Healthcare Pharmaceutical ltd., Rajendrapur, Gazipur	Ceftibuten 180mg/5ml Powder for Suspention Ceftibuten Dihydrate INN 3.9157gm eq. to Ceftibuten 3.60gm/100ml Antibiotic	Do	Contraindications: Ceftibuten is contraindicated in patients with known allergy to the Cephalosporin group of antibiotics. Side effects: Aphasia, jaundice, melena, psychosis, stridor, toxic epidermal necrolysis,serum ickness-like reactions and Stevens-Johnson syndrome.	Ceftibuten 1.80gm/100ml Powder For Suspension	USFDA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
88.	Healthcare Pharmaceutical ltd., Rajendrapur, Gazipur	Dalbavancin HCl INN 1020mg eq. to Dalbavancin 1000mg/Vial Dalbavancin HCl INN 1020mg eq. to Dalbavancin 1000mg/Vial Antibiotic	Indicated for acute bacterial skin and skin structure infections (ABSSSI) caused by designated susceptible strains of Gram-positive microorganisms. To reduce the development of drug-resistant bacteria and maintain the effectiveness of dalbavancin and other antibacterial drugs, dalbavancin should be used only to treat infections that are proven or strongly suspected to be caused by susceptible bacteria.	Contra-indication: Hypersensitivity to dalbavancin Side-effects: The most common adverse reactions in patients treated with dalbavancin were nausea (5.5%), headache (4.7%), and diarrhea (4.4%).	500 mg/Vial Injection		c‡qvRbxq †idv‡iY ‡bB weavq Av‡e`b bv gÄyi Kiv ‡h‡Z cv‡i	c‡qvRbxq †idv‡iY bv_vKq Av‡e`b bvgÄyi Kiv হল
89.	ACI Ltd., Narayanganj	Ertapenem 1gm /Vial Injection Ertapenem USP 1 gm/Vial Antibiotic	Ertapenem is a penem antibacterial indicated in adult patients and pediatric patients (3 months of age and older) for the treatment of the following moderate to severe infections caused by susceptible bacteria: Complicated intra-abdominal infections , Complicated skin and skin structure infections, including diabetic foot infections without osteomyelitis, Community-acquired pneumonia, Complicated urinary tract infections including pyelonephritis, Acute pelvic	Contraindications: Known hypersensitivity to product components or anaphylactic reactions to β-lactams. Due to the use of lidocaine HCl as a diluent, Ertapenem administered intramuscularly is contraindicated in patients with a known hypersensitivity to local anesthetics of the amide type. Side Effects: Adults: The most common side effects in patients treated with Ertapenem, including those who were switched to therapy with an oralantimicrobial, were diarrhea,	New	USFDA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj

			infections including postpartum endomyometritis, septic abortion and post surgical gynecologic infections. Ertapenem is indicated in adults for the prophylaxis of surgical site infection following elective colorectal surgery.	nausea, headache and infused vein complication. In the prophylaxis indication the overall adverse experience profile was generally comparable to that observed for ertapenem in other clinical tria Pediatrics: Side effects in this population were comparable to adults. The most common side effects in pediatric patients treated with Ertapenem, including those who were switched to therapy with an oral antimicrobial, were diarrhea, vomiting and infusion site pain.				
90.	Healthcare Pharmaceutical ltd., Rajendrapur, Gazipur	Faropenem 150mg FC Tablet Faropenem Sodium INN 161.55mg eq. to Faropenem 150mg Antibiotic	LRTIs: Acute bronchitis, Pneumonia. ENT Infections: Otitis externa, Tympanitis, Sinusitis. Genito-urinary infections: Pyelonephritis, Cystitis, Prostatitis. SSTIs: Pustular acne, Folliculitis, Impetigo. Gynecological infections: Adnexitis, Bartholin gland inflammation.	Contraindications: Known hypersensitivity to any of the components of this product or to other drugs in the same class. Patients who have demonstrated anaphylactic reaction to beta-lactams.	New		c‡qvRbxq †idv‡iÝ ‡bB weavq Av‡e`b bv gÄyi Kiv ‡h‡Z cv‡i|	c‡qvRbxq †idv‡iÝ bv _vKq Av‡e`b bvgÄyi Kiv হল|
91.	Healthcare Pharmaceutical ltd., Rajendrapur, Gazipur	Faropenem 200mg FC Tablet Faropenem Sodium INN 215.40mg eq. to Faropenem 200mg Antibiotic	-do-	-do-	New		c‡qvRbxq †idv‡iÝ ‡bB weavq Av‡e`b bv gÄyi Kiv ‡h‡Z cv‡i|	c‡qvRbxq †idv‡iÝ bv _vKq Av‡e`b bvgÄyi Kiv হল|
92.	Jayson Pharmaceuticals Ltd.	Finafloxacin 0.3% Otic Suspension Finafloxacin 0.3% Antibiotic	it is a quinolone antimicrobial indicated for the treatment of acute otitis externa (AOE) caused by susceptible strains of Pseudomonas aeruginosa and Staphylococcus aureus	Contra-indication: None Side effect: The most common adverse reactions occurring in 1% of patients with Finafloxacin 0.3% were ear pruritus and nausea		USFDA	Abyg`b Kiv †h‡Z cv‡i|	Abyg`b Kiv nj|
93.	ACI Ltd., Narayanganj	Fosfomycin 3 gm/Sachet Powder for Oral Suspension Fosfomycin Tromethamine USP 5.641gm eq. to 3.0gm Fosfomycin/Sachet Antibiotic	Fosfomycin Tromethamine is indicated only for the treatment of uncomplicated urinary tract infections (acute cystitis) in women due to susceptible strains of *Escherichia coli and Enterococcus faecalis*. It is not indicated for the treatment of pyelonephritis or perinephric abscess. If persistence or reappearance of bacteriuria	Contraindications: Fosfomycin tromethamine is contraindicated in patients with known hypersensitivity to the drug. Side Effects: The common side-effects are diarrhea nausea, headache, dizziness, vaginitis and dyspepsia. WARNINGS Clostridium difficile associated diarrhea (CDAD) has been reported with use of	New	USFDA	Abyg`b Kiv †h‡Z cv‡i|	Abyg`b Kiv nj|

			occurs after treatment other therapeutic agents should be selected.	nearly all antibacterial agents and may range in severity from mild diarrhea to fatal colitis. Treatment with antibacterial agents alters the normal flora of the colon leading to overgrowth of C. difficile.				
94.	Aristopharma Ltd.	Moxifloxacin 0.50g/100g Sterile Eye Ointment Moxifloxacin Hydrochloride (Sterile & Micronised) BP 0.5454 g eq.to Moxifloxacin 0.5g/100g Antibiotic	It is indicated for the treatment of bacterial conjunctivitis.	Contraindication: Moxifloxacin ophthalmic ointment is contraindicated in patients with a history of hypersensitivity to moxifloxacin, to other quinolones, or to any of the components in this medication. Side-effect: The most frequently reported ocular undesirable effects were conjunctivitis, decreased visual acuity, dry eye, keratitis, ocular discomfort, ocular hyperemia, ocular pain, ocular pruritus, sub-conjunctival hemorrhage, and tearing.	Moxifloxacin 0.5% Eye Drops &Eye Ointment 0.5% (For Export Only)		c‡qvRbxq †idv‡iÝ ‡bB weavq Av‡e`b bv gÄyi Kiv ‡h‡Z c‡i |	c‡qvRbxq †idv‡iÝ bv _vKq Av‡e`b bvgÄyi Kiv হল|
95.	Acme Laboratories Ltd.	Garenoxacin 400mg Tablet Garenoxacin Mesylate INN 507.06 mg eq. to 400 mg Garenoxacin Antibiotic (Quinolone)	This medication is a quinolone antibiotic, indicated for the treatment of chronic bronchitis, sinusitis, pneumonia, intra-abdominal infections and others. It blocks the action of enzymes that are necessary for the bacteria to make DNA.	Contraindication: Contraindicated in patients with known hypersensitivity to Garenoxacin or any other ingredients of the product. Side effects: The most commonly reported side effects are headache, diarrhea, abdominal pain, loose stools.	New		c‡qvRbxq †idv‡iÝ ‡bB weavq Av‡e`b bv gÄyi Kiv ‡h‡Z c‡i |	c‡qvRbxq †idv‡iÝ bv _vKq Av‡e`b bvgÄyi Kiv হল|
96.	Acme Laboratories Ltd.	Garenoxacin 600mg Tablet Garenoxacin Mesylate INN 760.59 mg Eq. to 600 mg Garenoxacin Antibiotic (Quinolone)	This medication is a quinolone antibiotic, indicated for the treatment of chronic bronchitis, sinusitis, pneumonia, intra-abdominal infections and others. It blocks the action of enzymes that are necessary for the bacteria to make DNA.	Contraindication: Contraindicated in patients with known hypersensitivity to Garenoxacin or any other ingredients of the product. Side effects: The most commonly reported side effects are headache, diarrhea, abdominal pain, loose stools.	New		c‡qvRbxq †idv‡iÝ ‡bB weavq Av‡e`b bv gÄyi Kiv ‡h‡Z c‡i |	c‡qvRbxq †idv‡iÝ bv _vKq Av‡e`b bvgÄyi Kiv হল|
97.	Incepta Pharmaceuticals Ltd.	Cefixime 100mg Dispersible Tablet Cefixime Trihydrate USP 111.97mg eq. to 100mg Cefixime Antibiotics-Cephalosporins	Cefixime is indicated in the treatment of the following infections caused by susceptible organisms. 1. Urinary tract infections 2.Upper and Lower respiratory tract infections 3. Acute otitis media. 4. Gonococcal urethritis 5. Typhoid	Contraindication: Hypersensitivity to cephalosporin antibiotics. Side effect: Cefixime is generally well tolerated. Majority of adverse reactions in clinical trials were mild and self–limiting in nature. The most frequent side effects seen with Cefixime are diarrhoea and stool changes; diarrhoea has been more commonly associated with higher doses. Some cases of moderate to severe diarrhea have been reported; this has occasionally warranted cessation of therapy.	200mg, 400mg Tablet		c‡qvRbxq †idv‡iÝ ‡bB weavq Av‡e`b bv gÄyi Kiv ‡h‡Z c‡i |	c‡qvRbxq †idv‡iÝ bv _vKq Av‡e`b bvgÄyi Kiv হল|

				Other side effects include headache and dizziness. Mild transient changes in liver and renal function tests have been observed. Allergies in the form of rash, pruritus, urticaria, drug fever and arthralgia have been observed.				
98.	Incepta Pharmaceuticals Ltd.	Folinic Acid (Leucovorin Calcium) 5mg Tablet Calcium Folinate BP 5.405mg eq. to Folinic Acid 5mg Antidote (to folic acid antagonists); antineoplastic adjunct	a) To diminish the toxicity and counteract the effect of impaired METHOTREXATE elimination. b) To treat the megaloblastic anemias due to folate deficiency, as in sprue, nutritional deficiency, megaloblastic anemias of pregnancy and infancy.	Contraindication: Calcium folinate therapy is contraindicated for the following: • Known hypersensitivity to calcium folinate, or to any components of the product formulation. • Pernicious anemia or other megaloblastic anemias where Vitamin B-12 is deficient. A hematologic remission may occur while neurologic manifestations continue to progress. Side effect: Allergic sensitization, including anaphylactoid reactions and urticaria, has been reported following the administration of both oral and parenteral Calcium Folinate.	New	USFDA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
99.	Incepta Pharmaceuticals Ltd. Ziska Pharmaceuticals Ltd. Globe Pharmaceuticals Ltd., Noakhali Beximco Pharmaceuticals Ltd., Tongi ,Gazipur Eskayef Bangladesh Limited.	Lesinurad 200 mg Tablet Lesinurad INN 200 mg **Antigout Agent**	It is indicated in combination with a xanthine oxidase inhibitor for the treatment of hyperuricemia associated with gout in patients who have not achieved target serum uric acid levels with a xanthine oxidase inhibitor alone. Limitations of Use: Lesinurad is not recommended for the treatment of asymptomatic hyperuricemia. Lesinurad should not be used as monotherapy.	Contraindication: • Severe renal impairment (eCLcr less than 30 mL/min), end stage renal disease, kidney transplant recipients, or patients on dialysis • Tumor lysis syndrome or Lesch-Nyhan syndrome Side effect: Most common adverse reactions in 12-month controlled clinical trials (occurring in greater than or equal to 2% of patients treated with Lesinurad in combination with a xanthine oxidase inhibitor and more frequently than on a xanthine oxidase inhibitor alone) were headache, influenza, blood creatinine increased, and gastroesophageal reflux disease.	New	USFDA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
100.	Incepta Pharmaceuticals Ltd.	Alirocumab 150mg injection Alirocumab In House 150mg in a single-dose pre-filled syringe	<u>*Primary Hyperlipidemia*</u> ALIROCUMAB is a PCSK9 (Proprotein Convertase Subtilisin Kexin Type 9) inhibitor antibody indicated as adjunct to diet and	Contraindication: History of a serious hypersensitivity reaction to Alirocumab. Side effect: The most commonly occurring adverse reactions (≥5% of patients treated with Alirocumab and occurring more	New	USFDA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj

		Antihypercholesterolemic drug	maximally tolerated statin therapy for the treatment of adults with heterozygous familial hypercholesterolemia or clinical atherosclerotic cardiovascular disease, who require additional lowering of LDL-cholesterol (LDL-C). *Limitations Of Use* The effect of Alirocumab on cardiovascular morbidity and mortality has not been determined.	frequently than with placebo) are nasopharyngitis, injection site reactions, and influenza				
101.	Incepta Pharmaceuticals Ltd.	Alirocumab 75mg injection Alirocumab In House 75mg in single-dose pre-filled syringe Antihypercholesterolemic drug	*Primary Hyperlipidemia* ALIROCUMAB is a PCSK9 (Proprotein Convertase Subtilisin Kexin Type 9) inhibitor antibody indicated as adjunct to diet and maximally tolerated statin therapy for the treatment of adults with heterozygous familial hypercholesterolemia or clinical atherosclerotic cardiovascular disease, who require additional lowering of LDL-cholesterol (LDL-C). *Limitations Of Use* The effect of Alirocumab on cardiovascular morbidity and mortality has not been determined.	Contraindication: History of a serious hypersensitivity reaction to Alirocumab. Side effect: The most commonly occurring adverse reactions (≥5% of patients treated with Alirocumab and occurring more frequently than with placebo) are nasopharyngitis, injection site reactions, and influenza	New	USFDA	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
102.	Acme Laboratories Ltd.	Cilnidipine 5mg Tablet Cilnidipine INN 5 mg Anti-hypertensive	Cilnidipine is a calcium channel blocker indicated for patients at risk of vascular damage caused by hypertension. It helps to control hypertension (high blood pressure) by reduce the functioning of the body`s angiotensin receptors.	Contraindications: Hypersensitivity to any component of this product, Side effects: Most common adverse reactions Nausea, Headaches, Trouble breathing, Swelling, Low blood pressure, Edema (Water retention).	New		*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
103.	Opsonin Pharma Limited, Bagura Road, Barisal	Cefixime 100mg/ml Pediatric Drops Cefixime USP 100 mg/ml Anti-infective	Uncomplicated Urinary Tract Infections Otitis Media Pharyngitis and Tonsillitis Acute Exacerbations of Chronic Bronchitis Uncomplicated Gonorrhea (cervical/urethral) Typhoid	Contraindications: Cefixime is contraindicated in patients with known allergy to cefixime or other cephalosporins. Side effects: Most common adverse reactions are gastro intestinal such as diarrhea (16%), nausea (7%), loose stools (6%), abdominal pain (3%), dyspepsia (3%), and vomiting	200 & 400 mg Tablet & Capsule, 100mg/5ml PFS, 200 mg/5ml PFS, 25mg/ml Pediatric drops		*c‡qvRb †bB wewfbœ Av‡e`b bv gÄi Kiv †h‡Z cv‡i |*	*c‡qvRb †bB wewfbœ Av‡e`b bv gÄyi Kiv হল |*

104.	Opsonin Pharma Limited, Bagura Road, Barisal	Aceclofenac 150 mg/3ml Injection Aceclofenac BP 150mg/3ml Anti-inflammatory	Aceclofenac is indicated for the relief of mild to moderate pain, Post Surgical Pain & Post Traumatic Pain.	Contraindications: Aceclofenac is contraindicated in patients previously sensitive to aceclofenac or aspirin or other NSAIDs. It should not be administered to patients with active or suspected peptic ulcer or gastrointestinal bleeding and moderate to severe renal impairment. Side effects: Generally aceclofenac is well tolerated. The majority of side effects are reversible and mild which include gastrointestinal disorders (dyspepsia, abdominal pain, nausea and diarrhoea) and occasional occurrence of headache, dizziness or tiredness. Dermatological complaints including rash or itching, pruritus, abnormal hepatic enzyme levels and raised serum creatinine have occasionally been reported.	100 mg Tablet & 200 mg SR Tablet		c‡qvRb †bB wea‡q Av‡e`b bv gÄi Kiv †h‡Z cv‡i	c‡qvRb ‡bB wea‡q Av‡e`b bv gÄyiKiv হল
105.	Incepta Pharmaceuticals LTd (Dhamrai Unit)	Astaxanthin 4 mg Soft Gelatin Capsule Astaxanthin Oleoresin 5% INN 80mg eq. to 4 mg Astaxanthin Antioxidant	• Strong antioxidant • Improves cardiovascular health (Atherosclerosis, reduce cholesterol). • Improves immune function. • Improves condition of skin • Protects skin from damage caused by sun (Reduce wrinkles, pimples and other signs of aging) • Improves recovery from central nervous system injuries • Protects from Parkinson 's disease, Dementia and Alzheimer's • Protects eyes from cataracts and macular degeneration. • Reduces inflammation (Arthritis) • Reduces risk of infertility Also Astaxnthin effectively reduce oxidative damage to DNA, decrease the risk for many types of cancer and stabilize blood sugar.	Contraindication: Contraindicated for those with known allergies to Astaxanthin or any other component of the product. Side effect: No clear data foun	New		Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
106.	Beacon Pharmaceuticals	Trihexephenidyl Hydrochloride	It is indicated as an adjunct in the	Contraindication: Should be avoiding in	0.10gm/100ml Syrup	USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

	Ltd.	0.040gm/100ml Syrup Trihexephenidyl Hydrochloride USP 0.040gm/100ml Antiparkinsonian	treatment of all forms of parkinsonism (postencephalitic, arteriosclerotic, and idiopathic). It is often useful as adjuvant therapy when treating these forms of parkinsonism with levodopa. Additionally, it is indicated for the control of extrapyramidal disorders caused by central nervous system drugs such as the dibenzoxazepines, phenothiazines, thioxanthenes, and butyrophenones.	gastro-intestinal obstruction and myasthenia gravis. Side-effect: Side effects include constipation, dry mouth, nausea, vomiting, tachycardia, dizziness, confusion, euphoria, hellucinations, impaired memory, anxiety, restlessness, urinary retention, blurred vision, and rash. Angle-closure glaucoma may occur very rarely.		(2mg/5ml Elixir)		
107.	ACI Ltd., Narayanganj	Paliperidone 1.5 mg Extended Release Tablet Paliperidone INN 1.5 mg Antipsychotic	Paliperidone is an atypical antipsychotic agent indicated for Treatment of schizophrenia.In Adults: Efficacy was established in three 6-week trials and one maintenance trial.Adolescents (ages 12-17): Efficacy was established in one 6-week trial. Treatment of schizoaffective disorder as monotherapy and as an adjunct to mood stabilizers and/or antidepressants.Efficacy was established in two 6-week trials in adult patients.	Contraindications: Known hypersensitivity to paliperidone, risperidone, or to any components in the formulation. Side effects: Commonly observed adverse reactions (incidence ≥ 5% and at least twice that for placebo) were Adults with schizophrenia: extrapyramidal symptoms, tachycardia, and akathisia. Adolescents with schizophrenia: somnolence, akathisia, tremor, dystonia,cogwheel rigidity, anxiety, weight increased, and tachycardia. Adults with schizoaffective disorder: extrapyramidal symptoms, somnolence, dyspepsia, constipation, weight increased, and nasopharyngitis.	3/6/9mg ER Tablet	USFDA	Abyjgv`b Kiv †h‡Z cv‡i|	Abyjgv`b Kiv nj|
108.	Aristopharma Ltd.	Levosulpiride 25 mg Tablet Levosulpiride INN 25 mg Antipsychotic and prokinetic agent	This medication is an antipsychotic and prokinetic agent, prescribed for dyspepsia, gastro-esophageal reflux disease, and irritable bowel syndrome.	Contraindication: Contraindicated in patients with gastrointestinal bleeding, and intestinal obstruction. Side-effect: Genitourinary - Absence of menstrual period, breast enlargement in male, spontaneous milk secretion, and changes in libido.	New		c‡qvRbxq †idv‡iY ‡bB wea|q Av‡e`b bv gÄyi Kiv †h‡Z cv‡i|	c‡qvRbxq †idv‡iY bv_vKq Av‡e`b bvgÄyi Kiv হল|
109.	Healthcare Pharmaceutical ltd., Rajendrapur, Gazipur UniMed & UniHealth Mfg. Ltd., Gazipur	Daclatasvir 30mg Tablet Daclatasvir Dihydrochloride INN 32.964mg eq. to Daclatasvir 30 mg Antiviral	Daclatasvir is a hepatitis C virus (HCV) NS5A inhibitor indicated for use with sofosbuvir, with or without ribavirin, for the treatment of chronic Hepatitis C Virus.	Contraindications: Coadministration with medicinal products that strongly induce cytochrome P450 3A4 (CYP3A4) and Pglycoprotein transporter (P-gp) and thus may lead to lower exposure and loss of efficacy of Daclatasvir. These active substances includebut are not limited to phenytoin, carbamazepine, oxcarbazepine, phenobarbital, rifampicin, rifabutin,	60mg Tablet	USFDA	Abyjgv`b Kiv †h‡Z cv‡i|	Abyjgv`b Kiv nj|

				rifapentine, systemic dexamethasone, and the herbal product St John's wort (Hypericum perforatum). Side-effects: The most frequently reported adverse reactions were fatigue, headache, pruritus, insomnia, influenzalike, illness, dry skin, nausea, decreased appetite, alopecia, rash, asthenia, irritability, myalgia, anaemia, pyrexia, cough, dyspnoea, neutropenia, diarrhoea and arthralgia				
110.	UniMed & UniHealth Mfg. Ltd., Gazipur	Ribavirin 400mg Tablet Ribavirin USP 400mg Antiviral	severe respiratory syncytial virus bronchiolitis in infants and children; in combination with peginterferon alfa or interferon alfa for chronic hepatitis C in patients without liver decompensation	Contra-indication: Severe cardiac disease, including unstable or uncontrolled cardiac disease in previous 6 months; haemoglobinopathies; severe debilitating medical conditions; autoimmune disease (including autoimmune hepatitis); uncontrolled severe psychiatric condition; history of severe psychiatric condition in children Side-effects: Specific side-effects for inhaled treatment Worsening respiration, bacterial pneumonia, and pneumothorax reported; rarely non-specific anaemia and haemolysis Specific side-effects for oral treatment Haemolytic anaemia (anaemia may be improved by epoetin); also (in combination with peginterferon alfa or interferon alfa) nausea, vomiting, dyspepsia, abdominal pain, flatulence, constipation, diarrhoea, colitis, chest pain, palpitation, tachycardia, peripheral oedema, changes in blood pressure, syncope, flushing, cough, dyspnoea, headache, dizziness, asthenia, impaired concentration and memory, sleep disturbances, abnormal dreams, anxiety, depression, suicidal ideation (more frequent in children), psychoses, dysphagia, weight loss, dysphonia, paraesthesia, hypoaesthesia, ataxia, hypertonia, influenza-like symptoms, thyroid disorders, hyperglycaemia, menstrual disturbances, breast pain,	200mg Tablet	BNF-70 Page-545-546	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj	

				prostatitis, sexual dysfunction, micturition disorders, leucopenia, thrombocytopenia, lymphadenopathy, dehydration, hypocalcaemia, myalgia, arthralgia, hyperuricaemia, visual disturbances, eye pain, dry eyes, hearing impairment, tinnitus, earache, dry mouth, taste disturbances, mouth ulcers, stomatitis, glossitis, tooth disorder, gingivitis, alopecia, pruritus, dry skin, rash (including very rarely Stevens-Johnson syndrome and toxic epidermal necrolysis), increased sweating, psoriasis, photosensitivity, and acne; less commonly pancreatitis, gastro-intestinal bleeding, and hypertriglyceridaemia; rarely peptic ulcer, arrhythmias, cardiomyopathy, myocardial infarction, pericarditis, stroke, interstitial pneumonitis, pulmonary embolism, seizures, renal failure, vasculitis, rheumatoid arthritis, systemic lupus erythematosus, sarcoidosis, optic neuropathy, and retinal haemorrhage; very rarely aplastic anaemia and peripheral ischaemia; in children also growth retardation (including decrease in height and weight), pallor, tachypnoea, hyperkinesia, virilism, and skin discoloration				
111.	Incepta Pharmaceuticals Ltd.	Tenofovir Alafenamide 10 mg Tablet Tenofovir Alafenamide Fumarate INN 11.2mg eq. to Tenofovir Alafenamide 10 mg Antiviral	Chronic Hepatitis B Human Immunodeficiency Virus (HIV)	Contraindication: No clear data found Side effect: In the Phase II study of adarunavir/cobicistat/emtricitabine/tenofovir alafenamide fixed-dose combination tablet discussed under the previous question (GS-US-299-0102), most side effects were mild to moderate in severity. Some side effects that were reported by participants receiving the darunavir/cobicistat/emtricitabine/tenofovir alafenamide tablet included diarrhea, upper respiratory tract infection, fatigue, nausea, and rash.	Tenofovir Disoproxil Fumarate 300 mg Tablet		c‡qvRbxq †idv‡iY bv_vKvq AvcZZt ¯‹MZ ivLv †h‡Z cv‡i|	c‡qvRbxq †idv‡iY bv_vKq Avt‡e`b bvgÄyK iv হল|

112.	Beximco Pharmaceuticals Ltd., Tongi ,Gazipur	Salbutamol 5mg/2.5ml Respirator or Nebuliser Solution Salbutamol Sulphate BP 6.00mg eq.to Salbutamol 5mg/2.5ml Bronchodilators	It is indicated for use in the routine management of chronic bronchospasm unresponsive to conventional therapy and the treatment of acute severe asthma.	Contraindications: Hypersensitivity to the active substance salbutamol or to the excipients. Side effects: Nausea, Sweating, Restlessness, Headache, Dizziness etc.	5mg/ml; & 2.5 mg/3 ml Respirator solution	MHRA	Abjygv`b Kiv †h‡Z cv‡i	Abjygv`b Kiv nj
113.	ACI Ltd., Narayanganj	Armodafinil 150 mg Tablet Armodafinil INN 150mg CNS Stimulant	Armodafinil is indicated to improve wakefulness in adult patients with excessive sleepiness associated with obstructive sleep apnea (OSA), narcolepsy, or shift work disorder (SWD.). Limitation of use: In OSA, Armodafinil is indicated to treat excessive sleepiness and not as treatment for the underlying obstruction.	Contraindications: Armodafinil is contraindicated in patients with known hypersensitivity to modafinil and armodafinil. Side effects: Most common adverse reactions are (5%): headache, nausea, dizziness, insomnia Warning and Precautions Serious Rash, including Stevens-Johnson Syndrome: discontinue It is at the first sign of rash, unless the rash is clearly not drug- related	New	USFDA	Abjygv`b Kiv †h‡Z cv‡i	Abjygv`b Kiv nj
114.	ACI Ltd., Narayanganj	Armodafinil 250 mg Tablet Armodafinil INN 250mg CNS Stimulant	Armodafinil is indicated to improve wakefulness in adult patients with excessive sleepiness associated with obstructive sleep apnea (OSA), narcolepsy, or shift work disorder (SWD.). Limitation of use: In OSA, Armodafinil is indicated to treat excessive sleepiness and not as treatment for the underlying obstruction.	Contraindications: Armodafinil is contraindicated in patients with known hypersensitivity to modafinil and armodafinil. Side effects: Most common adverse reactions are (5%): headache, nausea, dizziness, insomnia	New	USFDA	Abjygv`b Kiv †h‡Z cv‡i	Abjygv`b Kiv nj
115.	Incepta Pharmaceuticals Ltd.	Alosetron HCl 1mg Tablet Alosetron HCl INN 1mg CNS-5HT3 Antagonist	Alosetron Hydrochloride Tablets are a selective serotonin 5 HT_3 antagonst is indicated only for women with severe diarrhea-predominant irritable bowel syndrome (IBS) who have: chronic IBS symptoms (generally lasting 6 months or longer), had anatomic or biochemical abnormalities of the gastrointestinal tract excluded, and not responded adequately to conventional therapy. Severe IBS includes diarrhea and 1	Contraindication: Do not initiate on patients with constipation History of chronic or severe constipation or sequelae from constipat on intestinal obstruction stricture toxic megacolon gastrointestinal perforation and/or adhesions ischemic colitis impaired intestinal circulation thrombophlebitis or hypercoagulable state Crohn's disease or ulce at ve colitis diverticulitis severe hepatic impairment Inability to understand or comply with the Patient Acknowledgement	New	USFDA	Abjygv`b Kiv †h‡Z cv‡i	Abjygv`b Kiv nj

			or more of the following frequent and severe abdominal pain/discomfort frequent bowel urgency or fecal incontinence disability or restriction of daily activities due to IBS.	Concomitant use of fluvoxamine. Side effect: Most common adverse reactions (incidence >2% and >placebo) in clinical studies were constipation abdominal discomfort and pain nausea and gastrointestinal discomfort and pain.				
116.	Incepta Pharmaceuticals Ltd.	Alosetron HCl 0.5mg Tablet Alosetron HCl INN 0.5mg CNS-5HT3 Antagonist	-do-	-do-	New	USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
117.	Popular Pharmaceuticals Limited	Hydroxyprogesterone Caproate 250mg/ml Vial Injection Hydroxyprogesterone Caproate USP 250mg/ml Contraceptive	It is a progestin indicated to reduce the risk of preterm birth in women with a singleton pregnancy who have a history of singleton spontaneous preterm birth. Limitation of use: It is not intended for use in women with multiple gestations or other risk factors for preterm birth.	Contraindications: Current or history of thrombosis or thromboembolic disorders •Known or suspected breast cancer, other hormone-sensitive cancer, or history of these conditions (4) •Undiagnosed abnormal vaginal bleeding unrelated to pregnancy •Cholestatic jaundice of pregnancy •Liver tumors, benign or malignant, or active liver disease •Uncontrolled hypertension Side Effects: Most common adverse reactions reported in ≥ 2% of subjects and at a higher rate in the Makena group than in the control group are injection site reactions (pain [35%], swelling [17%], pruritus [6%], nodule [5%]), urticaria (12%), pruritus (8%), nausea (6%), and diarrhea (2%).	New	USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
118.	Incepta Pharmaceuticals Ltd.	Fluticasone Furoate 200mcg Capsule intended for oral inhalation Fluticasone Furoate INN 200mcg Corticosteroid	It is a corticosteroid indicated for the maintenance treatment of asthma as prophylactic therapy in patients aged 12 years and older. Important Limitation of Use: -NOT indicated for the relief of acute bronchospasm.	Contraindication: Primary treatment of status asthmaticus or acute episodes of asthma requiring intensive measures. Severe hypersensitivity to milk proteins or any ingredients of Fluticasone Furoate Side effect: Most common adverse reactions (reported in greater than or equal to 5% of subjects) are: upper respiratory tract infection, nasopharyngitis, headache, and bronchitis	100mcg, 250mcg, 500mcg DPI Capsule	USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
119.	Incepta Pharmaceuticals Ltd.	Phenylephrine hydrochloride 10mg/ml vial Injection	It is an alpha-1 adrenergic receptor agonist indicated for the treatment of clinically important hypotension	Contraindication: None Side effect: Most common adverse reactions during treatment: nausea,	New	USFDA	c‡qvRb †bB wea¨vq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i	c‡qvRb †bB wea¨vq Av‡e`b bv gÄyi Kiv হল

		Phenylephrine Hydrochloride USP 10mg/ml Vial Decongestant	resulting primarily from vasodilation in the setting of anesthesia.	vomiting, and headache.				
120.	Incepta Pharmaceuticals Ltd.	Phenylephrine hydrochloride 50mg/5ml vial Injection Phenylephrine Hydrochloride USP 50mg/5ml Decongestant	It is an alpha-1 adrenergic receptor agonist indicated for the treatment of clinically important hypotension resulting primarily from vasodilation in the setting of anesthesia.	Contraindication: None Side effect: Most common adverse reactions during treatment: nausea, vomiting, and headache.	New	USFDA	c‡qvRb †bB wea‡q Av‡e`b bv gÄyi Kiv †h‡Z cv‡i	c‡qvRb ‡bB wea‡q Av‡e`b bv gÄyi Kiv হল
121.	Incepta Pharmaceuticals Ltd.	Barium Sulfate 20.0mg/ml Suspension Barium Sulfate BP 20.0mg/ml Diagnostic Agent	Indicated for use in computed tomography (CT) of the abdomen to delineate the gastrointestinal (GI) tract in adult and pediatric patients.	Contraindication: ➢ Known or suspected perforation of the GI tract. ➢ Known obstruction of the GI tract. ➢ Conditions associated with high risk of GI perforation or aspiration. ➢ Known severe hypersensitivity to barium sulfate or any of the excipients of this products Side effect: Common adverse reactions include nausea, vomiting, diarrhea and abdominal cramping	New	USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
122.	Incepta Pharmaceuticals Ltd.	Collagenase 250 Unit/gm Ointment Collagenase INN 250 Unit/gm Enzyme	Collagenase Ointment is indicated for debriding chronic dermal ulcers and severely burned areas.	Contraindication: Collagenase Ointment is contraindicated in patients who have shown local or systemic hypersensitivity to collagenase. Side effect: No allergic sensitivity or toxic reactions have been noted in clinical use when used as directed. However, one case of systemic manifestations of hypersensitivity to collagenase in a patient treated for more than one year with a combination of collagenase and cortisone has been reported.	New	USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
123.	Acme Laboratories Ltd.	Lactase 9000 Units/15 ml Pediatric Drops Lactase USP 0.300gm eq. to 9000 Units Lactase/15ml	Lactase Drops contains lactase enzyme which is also present in the human body naturally. Lactase enzyme breaks down the complex sugar Lactose, which is present in	Contraindication: Lactase Drops is contraindicated in patients with known hypersensitivity to lactase or any other components of the product.	Lactase 300 mg Chewable Tablet		Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

		Enzyme	milk, milk products, baby formulas and even breast milk, to simple digestible sugars like glucose and galactose. Lactase Drops breaks down large quantity of lactose, in the feed containing the milk, before it is fed to the baby. This helps easy digestion of lactose and prevents infant colic. Lactase Drops can be used every time the milk containing feed is given, normally till the baby is 3 to 4 months old. By this period baby's digestive system can start producing their own lactase enzyme subsequently. Lactase Drops can be withdrawn gradually by first using half the quantity, followed by using in alternate feeds, then once a day, then totally stopping. Lactase Drops can be restarted if required when discomfort is observed.	Side effects: There are no major side effects reported. The very few side effects may include rash, difficulty in breathing, tightness in chest etc.				
124.	Incepta Pharmaceuticals Ltd.	Lactase 50000 Units/15ml Paed. Drop Lactase USP 1.667gm eq. to Lactase 50000 Units/15ml Enzyme	For the relief of symptoms associated with lactose intolerance in infants	Contraindication: Lactase Drops is contraindicated in patients with known hypersensitivity to lactase or any other components of the product. Side effects: There are no major side effects reported. The very few side effects may include rash, difficulty in breathing, tightness in chest etc.	300mg Tablet		Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
125.	Square Pharmaceuticals Ltd., Dhaka Unit, Kaliakoir, Gazipur	Sodium Polystyrene Sulfonate 15.00gm/Sachet powder for suspension Sodium Polystyrene Sulfonate USP 15.00gm/Sachet Powder for suspension **Exchange Resin**	Sodium polystyrene sulfonate is indicated in patients with Hyperkalemia associated with anuria or severe oliguria. It reduces serum levels of potassium and removes excess potassium from the body. Sodium polystyrene sulfonate is indicated in all states of hyperkalemia due to acute and chronic renal failure; examples	Contraindication: Sodium Polystyrin Sulfonate is contraindicated in patients with hypokalemia, obstructive3 bowel disease, neonates with reduced gut motiliry (postoperatively or drug induced) and oral administration in neonates. Side effects: Some degree of gastric irritation, anorexia, nausea, vomiting and constipation may occur. Large doses in	Calcium Polystyrene Sulfonate 300gm/Container Suspension	USFDA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj

			include use following abortion, complicated labor, and incompatible blood transfusion, crush injury, prostatectomy, severe burns, surgical shock, and in cases of severe glomerulonephritis and pyelonephritis.Sodium polystyrene sulfonate can also be useful in patients requiring dialysis. Serum potassium levels in acute renal failure often reach dangerous heights before a rise in blood urea indicates the need for hemodialysis. Calcium polystyrene sulfonate can be used to reduce these potassium levels and thereby postpone the need for the use of the artificial kidney machine until other causes make it necessary.	elderly individuals mauy cause fecal inpaction				
126.	Square Pharmaceuticals Ltd., Pabna Unit, Salgaria, Pabna	Ambroxol HCl 0.6gm/100ml Syrup Ambroxol HCl BP 0.6gm/100ml **Expectorant (Cough & Cold)**	Secretolytic therapy in acute and chronic bronchopulmonary diseases associated with abnormal mucus secretion and impaired mucus transport.	Contra-indications: It should not be used in patients known to be hypersensitive to ambroxol or other components of the formulation. Side effects: It is generally well tolerated. Mild upper gastro-intestinal side effects (primarily pyrosis, dyspepsia, and occasionally nausea, vomiting) have been reported, principally following parenteral administration. Allergic reactions have occured rarely, primarily skin rashes. There have been extremely rare case reports of severe acute anaphylactic-type reactions but their relationship to ambroxol is uncertain. Some of these patients have also shown allergic reactions to other substances.	0.3gm/100ml Syrup, 6mg/ml Pad. Drops, 75 SR Capsule		*c‡qvRbxq †idv‡iÝ ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i |*	*c‡qvRbxq †idv‡iÝ bv _vKq Av‡e`b bvgÄyi Kiv* হল|
127.	ACI Ltd., Narayanganj	Purified Water 99.05% Purified Water INN 99.05% **Eye Cleanser**	Purified water is used to wash the eye to help relieve- Irritation, Discomfort, Stinging, Itching, Removes loose foreign materials, air pollutants (smog or pollens), and chlorinated water. For daily or emergency eye cleansing.	Contraindications : Hypersensitivity to any component of the formulation. Side Effects : Irritation or hypersensitivity to preservative. Frequency not defined.	New	USFDA OTC Product	*Aby‡gv`b Kiv †h‡Z cv‡i |*	*Aby‡gv`b Kiv nj |*
128.	Ziska Pharmaceuticals Ltd.	Estropipate 0.75mg Tablet	It is indicated in the: 1. Treatment of moderate to severe	Contraindications: Undiagnosed abnormal genital bleeding. 2.	New	USFDA	*Aby‡gv`b Kiv †h‡Z cv‡i |*	*Aby‡gv`b Kiv nj |*

		Estropipate USP 0.75mg **Female Reproductive Agent**	vasomotor symptoms associated with the menopause. 2. Treatment of moderate to severe symptoms of vulval and vaginal atrophy associated with the menopause. When prescribing solely for the treatment of symptoms of vulvar and vaginal atrophy, topical vaginal products should be considered. 3. Treatment of hypoestrogenism due to hypogonadism, castration or primary ovarian failure. 4. Prevention of postmenopausal osteoporosis. When prescribing solely for the prevention of postmenopausal osteoporosis, therapy should only be considered for women at significant risk of osteoporosis and for whom non-estrogen medications are not considered to be appropriate.	Known, suspected, or history of cancer of the breast. 3. Known or suspected estrogen-dependent neoplasia. 4. Active deep vein thrombosis, pulmonary embolism or history of these conditions. 5. Active or recent (e.g., within the past year) arterial thromboembolic disease (e.g., stroke, myocardial infarction). 6. Liver dysfunction or disease. 7. It should not be used in patients with known hypersensitivity to its ingredients. 8. Known or suspected pregnancy. There is no indication for this tablet in pregnancy. There appears to be little or no increased risk of birth defects in children born to women who have used estrogens and progestins from oral contraceptives inadvertently during early pregnancy. Side-effects: Genitourinary system Changes in vaginal bleeding pattern and abnormal withdrawal bleeding or flow; breakthrough bleeding; spotting; dysmenorrhea; increase in size of uterine leiomyomata; vaginitis; including vaginal candidiasischange in amount of cervical secretion: changes in cervical ectropion; ovarian cancer; endometrial hyperplasia; endometrial cancer. 2. Breasts Ogen® estropipate tablets, USP Tenderness, enlargement, pain, nipple discharge, galactorrhea; fibrocystic breast changes; breast cancer. 3. Cardiovascular Deep and superficial venous thrombosis; pulmonary embolism; thrombophlebitis; myocardial infarction; stroke; increase in blood pressure. 4. Gastrointestinal Nausea, vomiting; abdominal cramps, bloating; cholestatic jaundice; increased incidence of gallbladder disease; pancreatitis; enlargement of hepatic hemangiomas. 5. Skin Chloasma or melasma that may persist when drug is discontinued; erythema multiforme; erythema nodosum; hemorrhagic eruption; loss of scalp hair; hirsutism; pruritus, rash. 6. Eyes Retinal				

				vascular thrombosis; steepening of corneal curvature; intolerance to contact lenses. 7. Central nervous system Headache, migraine, dizziness; mental depression; chorea; nervousness; mood disturbances; irritability; exacerbation of epilepsy; dementia. 8. Miscellaneous Increase or decrease in weight; reduced carbohydrate tolerance; aggravation of porphyria; edema; changes arthralgias; leg cramps; urticaria; angioedema; anaphylactoid/anaphylactic reactions; hypocalcemia; exacerbation of asthma; increased libido; triglycerides.				
129.	Ziska Pharmaceuticals Ltd.	Estropipate 1.5 mg Tablet Estropipate USP 1.5 mg Female Reproductive Agent	-do-	-do-	New	USFDA	*Abÿgv`b Kiv †h‡Z cv‡i |*	*Abÿgv`b Kiv nj |*
130.	Ziska Pharmaceuticals Ltd.	Estropipate 3.0 mg Tablet Estropipate USP 3.0mg Female Reproductive Agent	-do-	-do-	New	USFDA	*Abÿgv`b Kiv †h‡Z cv‡i |*	*Abÿgv`b Kiv nj |*
131.	Incepta Pharmaceuticals Ltd.	Levoleucovorin 50 mg /Vial Lyophilized powder for Injection Levoleucovorin Calcium Pentahydrate INN 64mg eq. to Levoleucovorin 50mg/Vial **Folate Analog**	Levoleucovorin is a folate analog indicated for: Rescue after high-dose methotrexate therapy in osteosarcoma. Diminishing the toxicity and counteracting the effects of impaired methotrexate elimination and of inadvertent overdosage of folic acid antagonists. Use in combination chemotherapy with 5-fluorouracil in the palliative treatment of patients with advanced metastatic colorectal cancer. Limitations of Use Levoleucovorin is not approved for pernicious anemia and megaloblastic anemias. Improper use may cause a hematologic remission while neurologic manifestations continue to progress.	**Contraindications:** Levoleucovorin is contraindicated for patients who have had previous allergic reactions attributed to folic acid or folinic acid. **Side effect:** Allergic reactions were reported in patients receiving Levoleucovorin. Vomiting (38%), stomatitis (38%) and nausea (19%) were reported in patients receiving Levoleucovorin as rescue after high-dose methotrexate therapy. The most common adverse reactions (>50%) in patients with advanced colorectal cancer receiving Levoleucovorin in combination with 5-FU were diarrhea, nausea and stomatitis	New	USFDA	*Abÿgv`b Kiv †h‡Z cv‡i |*	*Abÿgv`b Kiv nj |*
132.	Square Pharmaceuticals	Deflazacort 1mg Tablet	It is a glucocorticoid used as an anti-	Contraindications:	6 mg Tablet		*c‡qvRbxq †id‡iY ‡bB ïeavq*	*c‡qvRbxq †id‡iY bv_vKq*

	Ltd., Pabna Unit, Salgaria, Pabna	Deflazacort INN 1 mg Glucocorticoid	inflammatory and immunosuppressant. It can be used to treat a wide range of allergic and inflammatory conditions, including severe asthma and rheumatoid arthritis.	Contraindicated in patients who are receiving immunosuppressive vaccines, who have infections and hypersensitivity. Side effects: GI disturbances, musculoskeletal, endocrine, neuropsychiatric, ophthalmic, fluid and electrolyte disturbances; susceptible to infection, impaired healing, hypersensitivity, skin atrophy, striae, telangiectasia, acne, myocardial rupture following recent MI, thromboembolism.			Av‡e`b bv gÄy¨K iv †h‡Z cv‡i	Av‡e`b bvgÄy¨K iv হল|
133.	Beacon Pharmaceuticals Ltd. Julphar Bangladesh Ltd., Sreepur, Gazipur, Dhaka	Deflazacort 24mg Tablet Deflazacort INN 24mg **Glucocorticoid**	Rheumatoid arthritis, juvenile chronic arthritis, asthma and other airway diseases, pemphigus, uveitis, nephrotic syndrome,Immune suppression in transplantation, anaphylaxis, severe hypersensitivity reactions, dermatomyositis, mixed connective tissue disease, polyarteritis nodosa, bullous pemphigoid, ulcerative colitis, optic neuritis, autoimmune haemolytic anaemia, idiopathic thrombocytopenic purpura, acute and lymphatic leukaemia, malignant lymphoma.	**Contraindications:** Contraindicated in patients who are receiving immunosuppressive vaccines, who have infections and hypersensitivity. **Side effects:** GI disturbances, musculoskeletal, endocrine, neuropsychiatric, ophthalmic, fluid and electrolyte disturbances; susceptible to infection, impaired healing, hypersensitivity, skin atrophy, striae, telangiectasia, acne, myocardial rupture following recent MI, thromboembolism.	6mg Tablet		Abyjgv`b K iv †h‡Z cv‡i	Abyjgv`b K iv nj
134.	Square Pharmaceuticals Ltd., Pabna Unit, Salgaria, Pabna	Deflazacort 30mg Tablet Deflazacort INN 30 mg **Glucocorticoid**	-do-	-do-	6 mg Tablet		c‡qvRbxq †i dv‡iY ‡bB weavq Av‡e`b bv gÄy¨K iv †h‡Z cv‡i	c‡qvRbxq †i dv‡iY bv _vKq Av‡e`b bvgÄy¨K iv হল|
135.	Beacon Pharmaceutical Ltd., Julphar Bangladesh Ltd., Sreepur, Gazipur, Dhaka	Deflazacort 6mg/5ml Suspension Deflazacort INN 6mg/5ml **Glucocorticoid**	-do-	-do-	6mg Tablet		c‡qvRbxq †i dv‡iY ‡bB weavq Av‡e`b bv gÄy¨K iv †h‡Z cv‡i	c‡qvRbxq †i dv‡iY bv _vKq Av‡e`b bvgÄy¨K iv হল|
136.	Incepta Pharmaceuticals Ltd.	Prednisolone 5mg Delayed Release Tablet Prednisolone BP 5mg	Endocrine Disorders: Primary or secondary adrenocortical insufficiency, Congenital adrenal hyperplasia, Hypercalcemia associated with cancer, Nonsuppurative thyroiditis. Rheumatic Disorders : As adjunctive	**Contraindication:** Systemic fungal infections and known hypersensitivity to components. **Side effect:** Fluid and Electrolyte Disturbance: Sodium retention, Fluid retention, Congestive heart failure in susceptible patients, Potassium loss,	5/10/15/20mgTablet	USFDA	Abyjgv`b K iv †h‡Z cv‡i	Abyjgv`b K iv nj

			therapy for short-term administration in Psoriatic arthritis, Rheumatoid arthritis, Including juvenile rheumatoid arthritis, Ankylosing spondylitis, Acute and subacute bursitis, Acute nonspecific tenosynovitis, Acute gouty arthritis, Post-traumatic osteoarthritis, Synovitis of osteoarthritis, Epicondylitis. Collagen Diseases: During an exacerbation or as maintenance therapy in selected cases of Systemic lupus erythematosus, Systemic-dermatomyositis (polymyositis), Acute rheumatic carditis. Dermatologic Diseases: Pemphigus, Bullous dermatitis herpetiformis, Severe erythema multiforme (Stevens-Johnson syndrome), Exfoliative dermatitis, Mycosis fungoides, Severe psoriasis, Severe seborrheic dermatitis.	Hypokalemic alkalosis, Hypertension. Musculoskeletal: Muscle weakness, Steroid myopathy, Loss of muscle mass, Osteoporosis, Tendon rupture, particularly of the Achilles tendon, Vertebral compression fractures, Aseptic necrosis of femoral and humeral heads, Pathologic fracture of long bones. Gastrointestinal: Peptic ulcer with possible perforation and hemorrhage, Pancreatitis, Abdominal distention, Ulcerative esophagitis, Increases in alanine transaminase (ALT, SGPT), aspartate transaminase (AST, SGOT) and alkaline phosphatase have been observed following corticosteroid treatment. These changes are usually small, not associated with any clinical syndrome and are reversible upon discontinuation. Dermatologic: Impaired wound healing, Thin fragile skin, Petechiae and ecchymoses, Facial erythema,				
137.	Incepta Pharmaceuticals Ltd.	Eltrombopag 12.5 mg Tablet Eltrombopag Olamine INN 15.95mg eq. to Eltrombopag 12.5mg **Hematopoietic**	Eltrombopag is a thrombopoietin receptor agonist indicated for the treatment of: •thrombocytopenia in adult and pediatric patients 6 years and older with chronic immune (idiopathic) thrombocytopenia (ITP) who have had an insufficient response to corticosteroids, immunoglobulins, or splenectomy. Thrombocytopenia in patients with chronic hepatitis C to allow the initiation and maintenance of interferon-based therapy. Patients with severe aplastic anemia who have had an insufficient response to immunosuppressive therapy. Limitations of Use: • Eltrombopag should be used only in patients with ITP whose degree of thrombocytopenia and clinical	Contraindication: None Side effect: In adult patients with ITP, the most common adverse reactions (greater than or equal to 5% and greater than placebo) were: nausea, diarrhea, upper respiratory tract infection, vomiting, increased ALT, myalgia, and urinary tract infection. In pediatric patients age 6 years and older with ITP, the most common adverse reactions (greater than or equal to 10% and greater than placebo) were upper respiratory tract infection, nasopharyngitis, and rhinitis. •In patients with chronic hepatitis C-associated thrombocytopenia, the most common adverse reactions (greater than or equal to 10% and greater than placebo) were: anemia, pyrexia, fatigue, headache, nausea, diarrhea, decreased appetite, influenza-like illness, asthenia, insomnia, cough, pruritus, chills, myalgia, alopecia,	New	USFDA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj

			condition increase the risk for leeding. • Eltrombopag should be used only in patients with chronic hepatitis C whose degree of thrombocytopenia prevents the initiation of interferon-based therapy or limits the ability to maintain interferon-based therapy. •Safety and efficacy have not been established in combination with direct-acting antiviral agents used without interferon for treatment of chronic hepatitis C infection.	and peripheral edema. In patients with severe aplastic anemia, the most common adverse reactions (greater than or equal to 20%) were: nausea, fatigue, cough, diarrhea, and headache.				
138.	Beacon Pharmaceuticals Ltd.	Erythropoietin 2000 IU/0.3ml Injection Erythropoietin Beta BP 2000 IU/0.3ml Hematopoietic	Treatment of anemia associated with chronic renal failure (renal anemia) in patients on dialysis. Treatment of symptomatic renal anemia in Patients not yet undergoing dialysis. Treatment of anemia in adult patients with solid tumors receiving chemotherapy. Treatment of anemia in adult patients with adult patients with multiple myeloma, low grade non- Hodgkin's lymphoma or chronic lymphocytic leukemia, who have a relative erythropoietin deficiency and are receiving anti-tumor therapy. Deficiency is defined as an inappropriately low serum erythropoietin level in relation to the degree of anemia. Increasing the yield of autologous blood from patients in a pre-donation program. Its use in this indication must be balanced against the reported increased risk of thromboembolic events. Treatment should only be given to patients with moderate anemia (Hb 10-13 g/dL [6.21-8.07 mmol/L], no iron deficiency) if blood conserving procedures are not available or insufficient when the scheduled major elective surgery requires a large	Contraindication: pure red cell aplasia following erythropoietin therapy;uncontrolled hypertension; patients unable to receive thromboprophylaxis; avoid injections containing benzyl alcohol in neonates Side-effect: diarrhoea, nausea, vomiting; dosedependent increase in blood pressure or aggravation of hypertension; in isolated patients with normal or low blood pressure, hypertensive crisis withencephalopathy-like symptoms and generalised tonic-clonic seizures requiring immediate medical attention; headache; dose-dependent increase in platelet count (but thrombocytosis rare) regressing during treatment	2000 IU/.5 ml; 3000 IU/0.75 ml, 4000 IU/.4ml; 5000 IU/0.5ml; 10,000 IU/ml Injection	BNF-69 Page-668	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

			volume of blood (4 or more units of blood for females or 5 or more units for males).					
139.	Beacon Pharmaceuticals Ltd.	Erythropoietin 3000 IU/0.3ml Injection Erythropoietin Beta BP 3000IU/0.3ml Hematopoietic	Do	-do-		BNF-69 Page-668	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
140.	Beacon Pharmaceuticals Ltd.	Erythropoietin 5000 IU/0.3ml Injection Erythropoietin Beta BP 5000 IU/0.3ml Hematopoietic	Do	-do-		BNF-69 Page-668	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
141.	Beximco Pharmaceuticals Ltd., Tongi ,Gazipur	Dienogest 2.00mg Tablet Dienogest INN 2.00mg **Hormone**	It is used for the treatment of Endometriosis.	Contraindications:Thromboembolic disorder, Diabetes mellitus with vascular involvement, Hormone-dependent malignancies, Undiagnosed vaginal bleeding etc Side effects: Headache or migraine, breast discomfort, hair loss, nausea, acne etc.	New		Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
142.	Incepta Pharmaceuticals Ltd.	Lenalidomide 15mg Capsule Lenalidomide INN 15mg **Immune Modulator**	Lenalidomide is a thalidomide analogue indicated for the treatment of patients with: •Multiple myeloma (MM), in combination with dexamethasone, in patients who have received at least one prior therapy. •Transfusion-dependent anemia due to low- or intermediate-1-risk myelodysplastic syndromes (MDS) associated with a deletion 5q abnormality with or without add itional cytogenetic abnormalities. •Mantle cell lymphoma (MCL) whose disease has relapsed or progressed after two prior therapies, one of which included bortezomib Limitations of Use:	Contraindication: • Pregnancy • Demonstrated hypersensitivity to lenalidomide Side effect: MM: Most common adverse reactions (≥20%) include fatigue, neutropenia, constipation, diarrhea, muscle cramp, anemia, pyrexia, peripheral edema, nausea, back pain, upper respiratory tract infection, dyspnea, dizziness, thrombocytopenia, tremor and rash. MDS: Most common adverse reactions (>15%) include thrombocytopenia, neutropenia, diarrhea, pruritus, rash, fatigue, constipation, nausea, nasopharyngitis, arthralgia, pyrexia, back pain, peripheral edema, cough, dizziness,	New	USFDA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj

			Lenalidomide is not indicated and is not recommended for the treatment of patients with chronic lymphocytic leukemia (CLL) outside of controlled clinical trials	headache, muscle cramp, dyspnea, pharyngitis, and epistaxis. MCL: Most common adverse reactions (≥15%) include neutropenia, thrombocytopenia, fatigue, diarrhea,anemia, nausea, cough, pyrexia, rash, dyspnea, pruritus, constipation, peripheral edema and leukopenia				
143.	Incepta Pharmaceuticals Ltd.	Lenalidomide 2.5mg capsule Lenalidomide INN 2.5mg Immune Modulator	-do-	-do-	New	USFDA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
144.	Incepta Pharmaceuticals Ltd.	Lenalidomide 20mg Capsule Lenalidomide INN 20mg Immune Modulator	-do-	-do-	New	USFDA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
145.	Incepta Pharmaceuticals Ltd.	Lenalidomide 5mg Capsule Lenalidomide INN 5mg Immune Modulator	-do-	-do-	New	USFDA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
146.	Aristopharma Ltd.	Cyclosporine 0.50g/100ml Ophthalmic Emulsion (0.5%) Cyclosporine USP 0.50g/100ml Immunomodulator	It is indicated to increase tear production in patients whose tear production is presumed to be suppressed due to ocular inflammation associated with keratoconjunctivitis sicca. Increased tear production was not seen in patients currently taking topical anti-inflammatory drugs or using punctal plugs.	Contra-indication : It is contraindicated in patients with known or suspected hypersensitivity to any of the ingredients in the formulation Side-effect: Clinical Trials Experience Because clinical trials are conducted under widely varying conditions, adverse reaction rates observed in the clinical trials of a drug cannot be directly compared to rates in the clinical trials of another drug and may not reflect the rates observed in practice	Cyclosporine 50mg/100ml (0.05%) Ophthalmic Emulsion		Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj

147.	Incepta Pharmaceuticals Ltd.	Thalidomide 150 mg Capsule Thalidomide INN 150 mg Immunomodulator	Thalidomide in combination with dexamethasone is indicated for the treatment of patients with newly diagnosed multiple myeloma (MM). Thalidomide is indicated for the acute treatment of the cutaneous manifestations of moderate to severe erythema nodosum leprosum (ENL). Thalidomide is not indicated as monotherapy for such ENL treatment in the presence of moderate to severe neuritis. Thalidomide is also indicated as maintenance therapy for prevention and suppression of the cutaneous manifestations of ENL recurrence.	Contraindication: Pregnancy: Demonstrated hypersensitivity to the drug or its components Side effect: MM: The most common adverse reactions (≥ 20%) are fatigue, hypocalcemia, edema, constipation, neuropathy-sensory,dyspnea, muscle weakness, leukopenia, neutropenia, rash/desquamation, confusion, anorexia, nausea, anxiety/agitation, asthenia, tremor, fever, weight loss, thrombosis/embolism, neuropathy-motor, weight gain, dizziness, and dry skin. ENL: The most common adverse reactions (≥ 10%) are somnolence, rash, and headache	New	USFDA	Abjygv`b Kiv †h‡Z cv‡i	Abjygv`b Kiv nj
148.	ACI Ltd., Narayanganj	Lenograstim 264.00 mcg/vial (eq. to 33.6 MIU) sterile bulk ready to fill powder Lenograstim INN 264.00mcg eq. to 33.6MIU/Vial Immunostimulant/ Hematopoietic growth factor	Reduction in the duration of neutropenia and associated complications following bone-marrow transplantation for non-myeloid malignancy (specialist use only) Reeducation in duration of neutropenia and associated complications following peripheral stem cells transplantation for non-myeloid malignancy (specialist use only)	Contraindication: Lenograstim should not be administered to patients with known hypersensitivity to the product or its constituents. Lenograstim should not be used to increase the dose intensity of cytotoxic chemotherapy beyond established dosage regimens and time courses since the drug could reduce myelotoxicity but not overall toxicity of cytotoxic drugs. Lenograstim should not be administered concurrently with cytotoxic chemotherapy. Lenograstim should not be administered to patients suffering from myeloid malignancy. Side-effects: Mucositis, Splenic rupture, Toxic epidermal necrolysis	New	BNF 70, Page: 842	Abjygv`b Kiv †h‡Z cv‡i	Abjygv`b Kiv nj
149.	Ziska Pharmaceuticals Ltd.	Ferric Carboxymaltose 750mg/15 ml vial Injection Ferric Carboxymaltose INN 750 mg/15 ml Mineral	It is an iron replacement product indicated for the treatment of iron deficiency anemia in adult patients: who have intolerance to oral iron or have had unsatisfactory response to oral iron; who have non-dialysis dependent chronic kidney disease.	Contraindications: Hypersensitivity to Injectafer or any of its inactive components. Side effects: The most common adverse reactions (>2%) are nausea, hypertension, flushing, hypophosphatemia, and dizziness	New	USFDA	Abjygv`b Kiv †h‡Z cv‡i	Abjygv`b Kiv nj
150.	Beacon Pharmaceutical	Potassium Iodate BP 253 µgm	➢ Provides Iodine during	**Contraindication**: Avoid if patients have	New		c‡qvRbxq †id‡iY ‡bB ïeavq	c‡qvRbxq †id‡iY bv_vKq

	Ltd.,	(Iodine BP 150µgm) Tablet Potassium Iodate BP 253µgm eq. to Iodine 150µgm Mineral	pregnancy and Breastfeeding to prevent neural tube defects ➢ Ensure normal foetal development and infant health ➢ Iodine deficiency disorder	thyroid condition associating with hyperthyroidism Side effect: Diarrhea, mild fever, mild headache, nausea, stomach pain, vomiting			Av‡e`b bv gÄÿ Kiv †h‡Z cv‡i	Av‡e`b bvgÄÿ Kiv হল|
151.	Ziska Pharmaceuticals Ltd.	Zinc oxide 8.0gm/100gm Lip Balm Zinc oxide BP 8.0gm/100gm Mineral	It helps to prevent sun burn on lip and decrease the risk of skin cancer and early skin aging that is caused by sun.	Contraindications: None Side effects: Spending time in the sun increases your risk of skin cancer and early skin aging	New		c‡qvRbxq †idv‡iÝ ‡bB weavq Av‡e`b bv gÄÿ Kiv †h‡Z cv‡i	c‡qvRbxq †idv‡iÝ bv_vKq Av‡e`b bvgÄÿ Kiv হল|
152.	Incepta Pharmaceuticals Ltd.	Zinc-L- Carnosine 37.5mg Capsule Zinc-L- Carnosine INN 37.5mg Mineral	It enhances the stomach's mucosal defenses, providing significant improvements in gastric ulcer patients.	Contraindication: Zinc-carnosine is contraindicated in those patients with a known hypersensitivity to it or any of its ingredients. Side effect: Zinc intake in excess has been reported to impair immune function. Some people report that zinc lozenges lead to stomach ache, nausea, mouth irritation, and a bad taste. One source reports that gastrointestinal upset, metallic taste in the mouth, blood in the urine, and lethargy can occur from chronic oral zinc supplementation over 150 mg per day, but those claims are unsubstantiated. In topical form, zinc has no known side effects when used as recommended.	New		c‡qvRbxq †idv‡iÝ ‡bB weavq Av‡e`b bv gÄÿ Kiv †h‡Z cv‡i	c‡qvRbxq †idv‡iÝ bv_vKq Av‡e`b bvgÄÿ Kiv হল|
153.	Globe Pharmaceuticals Ltd., Noakhali	Erdosteine 300 mg Capsule Erdosteine INN 300 mg Mucolytic	Symptomatic treatment of acute exacerbations of chronic bronchitis.	Cautions: History of peptic ulceration (may disrupt the gastric mucosal barrier). Side effects: Very rare- Abdominal pain, diarrhea, headache, nausea, rash, taste disturbance, urticaria, vomiting	New	BNF 70 Page: 256	Abÿgv`b Kiv †h‡Z cv‡i	Abÿgv`b Kiv nj|
154.	Eskayef Bangladesh Limited Beacon Pharmaceuticals Ltd.	Flibanserin 100mg FC Tablet Flibanserin 100mg **Multifunctional Serotonin**	It is indicated for the treatment of premenopausal women with acquired, generalized hypoactive sexual desire disorder (HSDD) as characterized by low sexual desire	**Contraindications:** Alcohol, Moderate or strong cytochrome P450 3A4 (CYP3A4) inhibitors, Hepatic impairment **Side effects:** Most common adverse	New	USFDA	c‡qvRbxq †idv‡iÝ ‡bB weavq Av‡e`b bv gÄÿ Kiv †h‡Z cv‡i	c‡qvRbxq †idv‡iÝ bv_vKq Av‡e`b bvgÄÿ Kiv হল|

	Globe Pharmaceuticals Ltd., Noakhali\ Acme Laboratories Ltd. Orion Pharma Ltd. Ziska Pharmaceuticals Ltd. Julphar Bangladesh Ltd., Sreepur, Gazipur, Dhaka Square Pharmaceuticals Ltd., Pabna Delta Pharma Ltd.	**Agonist Antagonist**	that causes marked distress or interpersonal difficulty and is NOT due to: • A co-existing medical or psychiatric condition, • Problems within the relationship • The effects of a medication or other drug substance. <u>Limitations of Use</u>: Flibanserin is not indicated for the treatment of HSDD in postmenopausal women or in men. Flibanserin is not indicated to enhance sexual performance.	reactions (incidence ≥2%) are dizziness, somnolence, nausea, fatigue, insomnia, and dry mouth.				
155.	ACI Ltd., Narayanganj	Fesoterodine Fumarate 4.0mg Extended Release (ER) Tablet Fesoterodine Fumarate INN 4 mg Muscarinic Antagonist	It is indicated for the treatment of overactive bladder with symptoms of urge urinary incontinence, urgency, and frequency.	Contraindications: Fesoterodine is contraindicated in patients with urinary retention, gastric retention, or uncontrolled narrow-angle glaucoma. Fesoterodine is also contraindicated in patients with known hypersensitivity to the drug or its ingredients or to tolterodine tartrate tablets. Side Effects: The common side-effects are dry mouth and constipation.	New	USFDA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
156.	Incepta Pharmaceuticals Ltd.	OnabotulinumtoxinA 50Units/Vial Injection OnabotulinumtoxinA (Ready to fill sterile solution) INN 1.0ml containing OnabotulinumtoxinA 50 Units/Vial Neuromuscular Blocking Agent	OnabotulinumtoxinA Cosmetic is an acetylcholine release inhibitor and a neuromuscular blocking agent indicated for: •Temporary improvement in the appearance of moderate to severe glabellar lines associated with corrugator and/or procerus muscle activity in adult patients •Temporary improvement in theappearance of moderate to severe lateral canthal lines associated with orbicularis oculi activity in adult patients	Contraindication: ➢ Hypersensitivity to any botulinum toxin preparation or to any of the components in the formulation ➢ Infection at the injection site Side effect: The most common adverse reactions are: •Glabellar Lines: eyelid ptosis (3%) •Lateral Canthal Lines: eyelid edema (1%)	New	USFDA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
157.	Incepta Pharmaceuticals Ltd.	OnabotulinumtoxinA 100Units/Vial Injection	-do-	-do-	New	USFDA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj

		OnabotulinumtoxinA (Ready to fill sterile solution) INN 2.0ml containing OnabotulinumtoxinA 100 Units/Vial Neuromuscular Blocking Agent						
158.	Beximco Pharmaceuticals Ltd., Tongi ,Gazipur	Diclofenac Sodium 4% Cutaneous Spray Diclofenac Sodium BP 40.00mg/gm **NSAID**	Diclofenac sodium is used for the local symptomatic relief of mild to moderate pain & inflammation following acute blunt trauma of small and medium-sized joints and periarticular structures	**Contraindications:** Hypersensitivity to diclofenac, acetylsalicylic acid or other NSAIDs drugs **Side-effects:** Pruritus has been reported at a frequency of 0.9% in a clinical trial.	25mg, 50mg Tablet 12.5mg, 25mg, 50mg Suppository & 75mg/3ml Injection 1% Gel	MHRA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
159.	Globe Pharmaceuticals Ltd., Noakhali	Ibuprofen 5% Gel Ibuprofen BP 5.0gm/100gm Nonsteroidal anti-inflammatory drug (NSAID)	Pain and inflammation in analgesic and anti-inflammatory for backache, pain of non-serious arthritic conditions, muscular pain, sprains, strains, sports injuries and neuralgia.	Contraindications: Contraindicated in case of hypersensitivity to any of the constituents. Hypersensitivity to aspirin, or other non-steroidal anti-inflammatory drugs, asthma, rhinitis or urticaria. Not to be used on broken or damaged skin. Side effects: Adverse reactions are very rare, extremely uncommon when ibuprofen is administered topically. If they occur, treatment should be discontinued:- Hypersensitivity: Hypersensitivity reactions may occur. Gastro-intestinal: Side effects such as abdominal pain and dyspepsia have been reported. Renal: Renal impairment can occur in patients with a history of kidney problems.	200mg, 400mg Tablet 300mg Capsule 100mg/5ml Suspension	MHRA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
160.	Beximco Pharmaceuticals Ltd., Tongi ,Gazipur	Ketorolac Tromethamine 10mg Mouth Dissolving Tablet Ketorolac Tromethamine USP 10.0mg **NSAID**	Ketorolac Tromethamine indicated for the management of moderately severe acute pain that requires analgesia at the opioid level, usually in a postoperative setting.	Contraindications: Peptic ulcer, history of GI bleeding, aspirin/anti-inflammatory allergy, dehydration, nasal polyps etc. . Side effects: GI upset, ulcers, bleeding, thirst, thrombocytopenia, abnormal Liver Function Tests (LFTs), anorexia etc.	10mg Tablet		Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
161.	Beacon Pharmaceuticals Ltd.	Naloxegol 25mg Film coated Tablet Naloxegol Oxalate INN 28.453mg eq. to Naloxegol 25mg **Opioid antagonist**	It is an opioid antagonist indicated for the treatment of opioid-induced constipation (OIC) in adult patients with chronic non-cancer pain	Contraindication: •Patients with known or suspected gastrointestinal obstruction and at increased risk of recurrent obstruction Concomitant use with strong CYP3A4 inhibitors (e.g., clarithromycin, ketoconazole) Side-effect: The most common adverse	New	USFDA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj

				reactions in clinical trials (≥3%) are: abdominal pain, diarrhea, nausea, flatulence, vomiting, and headache				
162.	Mundipharma (Bangladesh) Private Limited., Gazipur.	Oxycodone Hydrochloride 40mg Controlled Release Tablet Oxycodone Hydrochloride Controlled Release USP 40mg Opoid Analgesic	Indicated for pain severe enough to require daily, around-the-clock, long-term opioid treatment	Contraindication: • Patients who are hypersensitive to the active substance (oxycodone) or other opioid analgesics or to any ingredient in the formulation,acute appendicitis or pancreatitis, acute asthma or other obstructive airway, acute respiratory depression, elevated carbon dioxide levels in the blood, and cor pulmonale. Side effects: The most frequently observed adverse are asthenia, constipation, dizziness, dry mouth, headache, nausea, pruritus, somnolence, sweating and vomiting.	New	USFDA & BNF-70; Page: 370(MR)	Abţgv`b Kiv †h‡Z cv‡i|	Abţgv`b Kiv nj|
163.	Globe Pharmaceuticals Ltd., Noakhali	Atosiban 37.5mg/ 5 ml Concentrate for Solution for Infusion Atosiban acetate INN 39.77 mg eq. to Atosiban 37.5 mg Oxytocin receptor antagonist	Atosiban is indicated to delay imminent pre-term birth in pregnant adult women with: − regular uterine contractions of at least 30 seconds duration at a rate of ≥4 per 30 minutes − a cervical dilation of 1 to 3 cm (0-3 for nulliparas) and effacement of ≥ 50% − a gestational age from 24 until 33 completed weeks − a normal foetal heart rate	Contraindications: Abrupto placenta, antepartum hemorrahage (requering immediate delivery), eclampsia, intrauterine fetal death, itrauterine infection, intrauterine growth restriction with abnormal fetal heart rate, placenta praveia, premature ruptrue of membranes after 30 weeks' gestation, severe preeclampsia Side effects: Common or very common- Dizziness, headache, hot flashes, hyperglycemia, hypotension, injection site reaction, nausea, tachycardia, vomiting Uncommon- Fever, insomnia, pruritus, rash.	New	MHRA BNF 70 Page:708	Abţgv`b Kiv †h‡Z cv‡i|	Abţgv`b Kiv nj|
164.	Globe Pharmaceuticals Ltd., Noakhali	Atosiban 6.75mg/ 0.9 ml Solution for injection Atosiban acetate INN 7.158 mg eq. to Atosiban 6.75 mg Oxytocin receptor antagonist	-do-	-do-	New	MHRA BNF 70 Page:708	Abţgv`b Kiv †h‡Z cv‡i|	Abţgv`b Kiv nj|
165.	Healthcare Pharmaceutical ltd., Rajendrapur, Gazipur	Avanafil 200 mg Tablet Avanafil INN 200 mg	Erectile Dysfunction	Contraindications: Organic nitrate or nitric oxide donors.	50mg & 100mg Tablet		c‡qvRbxq †idv‡iÝ ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z	c‡qvRbxq †idv‡iÝ bv _vKq Av‡e`b bvgÄyi Kiv হল|

		PDE5 Inhibitor		Side effects: headache, dizziness, flushing, stuffy nose, runny nose, sinus pain, sore throat, diarrhea, constipation, stomach upset, and back pain			cv‡i	
166.	Globe Pharmaceuticals Ltd., Noakhali	Lanthanum 500 mg Chewable Tablet Lanthanum Carbonate Hydrate INN 856.47 mg eq. to elemental Lanthanum 500 mg Phosphate binder	Lanthanum is a phosphate binder indicated to reduce serum phosphate level in patients with end stage renal disease (ESRD).	Contraindications: Lanthanum is contraindicated in bowel obstruction, including ileus and fecal impaction. Side effects: The most common adverse reactions for Lanthanum are gastrointestinal events, such as nausea, vomiting and abdominal pain, which are minimized by taking with food and generally abated with time with continued dosing.	New	USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
167.	Navana Pharmaceuticals Ltd. Nipro JMI Pharma Ltd. Eskayef Bangladesh Limited.	Lorcaserin Hydrochloride 10mg Tablet Lorcaserin Hydrochloride Hemihydrate INN 10.40mg eq. to 10mg Lorcaserin Hydrochloride Serotonin 2C Receptor Agonist	It is a serotonin 2C receptor agonist indicated as an adjunct to a reduced-calorie diet and increased physic al activity for chronic weight management in adults with an initial body mass index (BMI) of: 30 kg/m^2 or greater (obese) or 27 kg/m^2 or greater (overweight) in the presence of at least one weight-related comorbid condition, (e.g., hypertension, dyslipidemia, type 2 diabetes) Limitations of Use: The safety and efficacy of coadministration with other products for weight loss have not been established The effect of BELVIQ on cardiovascular morbidity and mortality has not been established.	Contraindication: Pregnancy Category X Side-effect: Most common adverse reactions (greater than 5%) in non-diabetic patients are headache, dizziness, fatigue, nausea, dry mouth, and constipation, and in diabetic patients are hypoglycemia, headache, back pain, cough, and fatigue.	New	USFDA	c‡qvRb ‡bB wea|q Av‡e`b bv gÄyi Kiv †h‡Z cv‡i	c‡qvRb ‡bB wea|q Av‡e`b bv gÄyi Kiv হল
168.	Beacon Pharmaceutical Ltd.,	Carisoprodol 250mg Tablet Carisoprodol USP 250mg	Carisoprodol is indicated for the relief of discomfort associated with acute, painful musculoskeletal	Contraindication: • Acute intermittent porphyria	New	USFDA	c‡qvRb ‡bB wea|q Av‡e`b bv gÄyi Kiv †h‡Z cv‡i	c‡qvRb ‡bB wea|q Av‡e`b bv gÄyi Kiv হল

	Incepta Pharmaceuticals Ltd.	Skeletal Muscle Relaxant	conditions. Important Limitations: • Should only be used for acute treatment periods up to two or three weeks. • Not recommended in pediatric patients less than 16 years of age.	• Hypersensitivity reaction to a carbamate such as meprobamate Side effect: Most common adverse reactions (incidence > 2%) are drowsiness, dizziness, and headache.				
169.	Incepta Pharmaceuticals Ltd.	Carisoprodol 350mg Tablet Carisoprodol USP 350mg Skeletal Muscle Relaxant	Carisoprodol is indicated for the relief of discomfort associated with acute, painful musculoskeletal conditions. Important Limitations: • Should only be used for acute treatment periods up to two or three weeks. • Not recommended in pediatric patients less than 16 years of age.	Contraindication: • Acute intermittent porphyria • Hypersensitivity reaction to a carbamate such as meprobamate Side effect: Most common adverse reactions (incidence > 2%) are drowsiness, dizziness, and headache.	New	USFDA	c‡qvRb ‡bB wea‡q Av‡e`b bv gÄyi Kiv †h‡Z cv‡i	c‡qvRb ‡bB wea‡q Av‡e`b bv gÄyi Kiv হল
170.	Incepta Pharmaceuticals Ltd.	Cisatracurium 10mg/5ml vial Injection Cisatracurium Besylate BP 13.38mg eq. to Cisatracurium 10mg Skeletal muscle relaxant (nondepolarizing)	Cisatracurium Besylateis an intermediate-onset/intermediate-duration neuromuscular blocking agent indicated for in patients and outpatients as an adjunct to general anesthesia, to facilitate tracheal intubation, and to provide skeletal muscle relaxation during surgery or mechanical ventilation in the ICU.	Contraindication: Cisatracurium Besylateis contraindicated in patients with known hypersensitivity to the product and its components. The 10 mL multiple-dose vials of Cisatracurium Besylateis contraindicated for use in premature infants because the formulation contains benzyl alcohol. Side effect: The following adverse experiences were judged by investigators during the clinical trials to have a possible causal relationship to administration of Cisatracurium Besylate: Incidence Greater than 1% None. Incidence Less than 1% Cardiovascular bradycardia(0.4%) hypotension(0.2%), flushing (0.2%). Respiratory bronchospasm (0.2%). Dermatological rash (0.1%)	New	USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
171.	Incepta Pharmaceuticals Ltd.	Cisatracurium 50mg/5ml Vial Injection Cisatracurium Besylate BP 66.90mg eq. to Cisatracurium 50mg	Do	-do-	New	USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

		Skeletal muscle relaxant (nondepolarizing)						
172.	Acme Laboratories Ltd.	Sucralose 12 mg/Sachet Sucralose BP 12mg/Sachet Sweetener	Sucralose is the low-calorie sweetener made from sugar. It has a clean sweet taste like sugar. It is about 400-1000 times sweeter than that of sugar. It is used in place of sugar in processed food and beverages and in tabletop sweetener. Sucralose tablet can be used by all segments of the population including children, pregnant and lactating women, and people with diabetes.	Contraindication: No contraindication has been reported. Side effects: Sucralose is usually well tolerated. However, bloating, abdominal pain, gas, sometimes painful, nausea, heartburn, diarrhea, headaches, migraines, heart palpitations, shortness of breath, depression, anxiety and panic attacks, spaced-out or drugged sensation, joint pain, especially in the knees and dizziness may occur.	8 mg Tablet & 6.5gm/Sachet	USFDA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
173.	Techno Drugs Ltd.	Hyaluronidase 1500 IU/Vial Injection Hyaluronidase BP 1500 IU/Vial Tissue Permeability Modifier	It is indicated as an adjuvant ➢ in subcutaneous fluid administration for achieving hydration ➢ to increase absorption and dispersion of other injected drugs ➢ in subcutaneous urography for improving resorption of radiopaque agents.	Contraindications: Hypersensitivity Side Effects: Spread of Localized Infection, Ocular Damage, Enzyme Inactivation with Intravenous Administration.	150 IU/vial Injection		c‡qvRbxq †idv‡iY bv_vKq Av‡e`b bvgÄyi Kiv †h‡Z cv‡i	c‡qvRbxq †idv‡iY bv_vKq Av‡e`b bvgÄyi Kiv হল
174.	Beximco Pharmaceuticals Ltd., Tongi ,Gazipur	Mirabegron 50.00mg Extended Release Tablet Mirabegron INN 50.00mg **Urinary Tract Antispasmodics**	Mirabegron is indicated for the treatment of overactive bladder (OAB) with symptoms of urge urinary incontinence, urgency, and urinary frequency.	Contraindications: MIRABEGRON is contraindicated in patients who have known hypersensitivity reactions to mirabegron or any component of the tablet. Side effects: The most common side effect is hypertension, nasopharyngitis, urinary tract infection and headache.	25mg Extended Release Tablet	USFDA BNF-70, Page-672 (MR Tablet)	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
175.	Incepta Pharmaceuticals Ltd.	Aflibercept 40 mg/ml Injection Aflibercept INN 40 mg/ml Vascular endothelial growth factor (VEGF) inhibitor	Aflibercept is indicated for the treatment of patients with: ●Neovascular (Wet) Age-Related Macular Degeneration (AMD) ●Macular Edema Following Retinal Vein Occlusion (RVO) ● Diabetic Macular Edema (DME)	Contraindication: •Ocular or periocular infection •Active intraocular inflammation •Hypersensitivity Side effect: The most common adverse reactions (≥5%) reported in patients receiving Aflibercept were conjunctival	New	USFDA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj

			•Diabetic Retinopathy (DR) in Patients with DME	hemorrhage, eye pain, cataract, vitreous floaters, intraocular pressure increased, and vitreous detachment.				
176.	ACI Ltd., Narayanganj	Isoproterenol Hydrochloride 0.020 gm/100 ml vial injection Isoproterenol Hydrochloride USP 0.020gm/100ml Vial Vasopressor, bronchodilator, antiasthmatic	Isoproterenol hydrochloride injection is indicated: For mild or transient episodes of heart block that do not require electric shock or pacemaker therapy. For serious episodes of heart block and Adams-Stokes attacks (except when caused by ventricular tachycardia or fibrillation). For use in cardiac arrest until electric shock or pacemaker therapy, the treatments of choice, is available. For bronchospasm occurring during anesthesia. As an adjunct to fluid and electrolyte replacement therapy and the use of other drugs and procedures in the treatment of hypovolemic and septic shock, low cardiac output (hypoperfusion) states, congestive heart failure, and cardiogenic shock.	Contraindication: Use of isoproterenol hydrochloride injection is contraindicated in patients with tachyarrhythmias; tachycardia or heart block caused by digitalis intoxication; ventricular arrhythmias which require inotropic therapy; and angina pectoris. Side Effects: The most common side effects of isoproterenol are nervousness, headache, dizziness, nausea, visual blurring, tachycardia, palpitations, angina, Adams-Stokes attacks, pulmonary edema, hypertension, hypotension, ventricular arrhythmias, tachyarrhythmias, dyspnea, flushing of the skin, sweating, mild tremors, weakness and pallor.	New	USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
177.	MedRx Life Science Ltd, BSCIC, Nandanpur, B. Baria	Biotin 1mg Film Coated Tablet Biotin BP 1mg Vitamin	• Used for the maintenance of normal hair and skin. • For nutritional supplementation, effective aid in treating dietary shortage or imbalance. • Biotin has also been used to treat seborrhea (skin rash) in babies. • It contributes to forming energy, normal functioning of the nervous system, macronutrient metabolism and psychological function. • Contributes to the	Contraindications: None known. Side Effects: Biotin is a safe and nontoxic vitamin. It has not been associated with any serious side effects, even in large doses. The FDA reports that biotin is safe and well tolerated when taken by mouth in recommended doses.	New		c‡qvRbxq †idv‡iY ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i	c‡qvRbxq †idv‡iY bv _vKq Av‡e`b bvgÄyi Kiv হল

			maintenance of normal mucus membranes.					
178.	MedRx Life Science Ltd, BSCIC, Nandanpur, B. Baria	Biotin 2mg Film Coated Tablet Biotin BP 2mg Vitamin	• Used for the maintenance of normal hair and skin. • For nutritional supplementation, effective aid in treating dietary shortage or imbalance. • Biotin has also been used to treat seborrhea (skin rash) in babies. • It contributes to forming energy, normal functioning of the nervous system, macronutrient metabolism and psychological function. • Contributes to the maintenance of normal mucus membranes.	Contraindications: None known. Side Effects: Biotin is a safe and nontoxic vitamin. It has not been associated with any serious side effects, even in large doses. The FDA reports that biotin is safe and well tolerated when taken by mouth in recommended doses.	New		c‡qvRbxq †idv‡iY ‡bB ïeavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i	c‡qvRbxq †idv‡iY bv_vKq Av‡e`b bvgÄyi Kiv হল।
179.	Beximco Pharmaceuticals Ltd., Tongi ,Gazipur	Cholecalciferol 20,000 IU Capsule Cholecalciferol (As vitamin D_3 1,00,000 IU/g) USP 200mg eq. to Vitamin D_3 20, 000 IU **Vitamin**	Vitamin D is essential for normal bone growth and development and to maintain bone density. Vitamin D acts as a hormone and increases reabsorption of Calcium and Phosphorus by the kidneys and increased bone turnover. Prevention and Treatment of vitamin D deficiency states	Contraindications: Cholecalciferol is contraindicated in all diseases associated with hypercalcaemia. Side effects: Common side effect are including; hypercalcaemia syndrome or Calcium intoxication (depending on the severity and duration of hypercalcaemia), anorexia, headache, nausea, vomiting, abdominal pain or stomach ache and constipation with the administration of Cholecaciferol.	400IU, 1000IU Tablet 25 mcg/5ml Syrup, 5mg/ml Injection	MHRA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
180.	Beximco Pharmaceuticals Ltd., Tongi ,Gazipur	Cholecalciferol 40,000 IU Capsule Cholecalciferol (As vitamin D_3 1,00,000 IU/g) USP 400.000mg eq.to Vitamin D_3 40, 000 IU	-do-	-do-	400IU, 1000IU Tablet 25 mcg/5ml Syrup, 5mg/ml Injection	MHRA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

		Vitamin						
181.	Beximco Pharmaceuticals Ltd., Tongi ,Gazipur	Cholecalciferol 2,000 IU Tablet Cholecalciferol (As vitamin D_3 1,00,000 IU/g) USP 20.00mg eq.to Cholecalciferol 2, 000 IU **Vitamin**	-do-	-do-	400IU, 1000IU Tablet 25 mcg/5ml Syrup, 5mg/ml Injection	MHRA	Abyjgv`b Kiv †h‡Z cv‡i|	Abyjgv`b Kiv nj|
182.	Healthcare Pharmaceutical ltd., Rajendrapur, Gazipur	Cholecalciferol 40.0mg/100ml Oral Drops Cholecalciferol (Vitamin D_3 1.0M IU/gm) USP 40.0mg/100ml **Vitamin**	-do-	-do-	New		c‡qvRbxq †idv‡iÝ ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i|	c‡qvRbxq †idv‡iÝ bv_vKq Av‡e`b bvgÄyi Kiv হল|
183.	Drug International Ltd.	Cholecalciferol (Vitamin D_3) 800IU Soft Gelatin Capsule Cholecalciferol (Vitamin D_3) BP 800 IU **Vitamin**	-do-	-do-	1000 IU Tablet	MHRA	Abyjgv`b Kiv †h‡Z cv‡i|	Abyjgv`b Kiv nj|
184.	Globe Pharmaceuticals Ltd., Noakhali	Vitamin D_3 10800 IU/ ml Pediatric Drops Vitamin D3 (Cholecalciferol) USP 10800 IU/ ml **Vitamin**	• -do-	-do-	Cholecalciferol (Vit. D3) 1000 IU, Tablet Cholecalciferol (Vit. D3) 5 mg/ml, Injection Cholecalciferol (Vit. D3) 25 mcg/5 ml, Syrup		c‡qvRbxq †idv‡iÝ ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i|	c‡qvRbxq †idv‡iÝ bv_vKq Av‡e`b bvgÄyi Kiv হল|
185.	Globe Pharmaceuticals Ltd., Noakhali	Vitamin D3 14400IU/ml Oral Drops Vitamin D3 (Cholecalciferol) USP 14400 IU/ml Vitamin	-do-	-do-			c‡qvRbxq †idv‡iÝ ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i|	c‡qvRbxq †idv‡iÝ bv_vKq Av‡e`b bvgÄyi Kiv হল|

186.	Globe Pharmaceuticals Ltd., Noakhali	Vitamin D3 21600 IU/ml Oral Drops Vitamin D3 (Cholecalciferol) USP 21600 IU/ml Vitamin	-do-	-do-	New		c‡qvRbxq †idv‡iÝ ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i |	c‡qvRbxq †idv‡iÝ bv _vKq Av‡e`b bvgÄyi Kiv হল |
187.	MedRx Life Science Ltd, BSCIC, Nandanpur, B. Baria	Vitamin E 10000 IU/100gm Cream Vitamin E (Alpha-tocopherol) BP 10,000 IU/100gm Vitamin	• Vitamin E is an antioxidant which softens the skin and locks in moisture to help minimize the appearance of wrinkles. • This cream is a valuable aid for dry, rough skin and provides soothing relief after sun or wind exposure. • It's naturally absorbing, moisturizing and non-greasy	Contraindications: Vitamin E is likely safe for most healthy people when applied to the skin. Side Effects: Most people do not experience any side effects when taking the cream.	200 mg Cap. 200 mg Tab., 800 mg Cap., 400 mg Cap.		c‡qvRbxq †idv‡iÝ ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i |	c‡qvRbxq †idv‡iÝ bv _vKq Av‡e`b bvgÄyi Kiv হল |
188.	UniMed & UniHealth Mfg. Ltd., Gazipur	Mirabegron 50mg Prolonged-Release Tablet Mirabegron INN 50mg β3-Adrenoceptor Agonist	Urinary frequency, urgency, and urge incontinence	Contra-indications : Severe hypertension Side-effects: Ttachycardia, urinary-tract infection; less commonly dyspepsia, gastritis, palpitation, atrial , fibrillation, hypertension, vulvovaginal infection and pruritus, joint swelling, rash, pruritus	25mg Extended Release Tablet	USFDA BNF-70, Page-672 (MR Tablet)	Abygw`b Kiv †h‡Z cv‡i |	Abygw`b Kiv nj |
189.	Incepta Pharmaceuticals Ltd.	L-Ornithine L-Aspartate 3gm/Sachet Granules L-Ornithine L-Aspartate INN 3 gm/Sachet	For the treatment of concomitant diseases and sequelae of acute and chronic liver diseases (e.g. liver cirrhosis) with the symptoms of latent and manifest hepatic encephalopathy.	Contraindication: Hypersensitivity to L-ornithine-L-aspartate or any other excipients of these products. Severe renal insufficiency (a serum creatinine level in excess of 3 mg/100 ml can be regarded as the guide value). Side effect: Uncommon nausea, vomiting, stomach ache, flatulence, diarrhoea; very rare pain in the limbs. E110 can trigger allergic reactions.	New		Abygw`b Kiv †h‡Z cv‡i |	Abygw`b Kiv nj |
	Incepta Pharmaceuticals Ltd.	L-Ornithine L-Aspartate 5gm/10ml Ampoule Injection L-Ornithine L-Aspartate INN 5gm/10ml	-do-	-do-	New		Abygw`b Kiv †h‡Z cv‡i |	Abygw`b Kiv nj |

190.	Beacon Pharmaceutical Ltd.	Obeticholic Acid 10mg Tablet Obeticholic Acid INN 10mg	Liver diseases, including primary biliary cirrhosis and non-alcoholic fatty liver disease (steatohepatitis)	Contraindication: Contraindicated for those with known allergies to Obeticholic Acid or any other component of the product. Side effect: The most common side effect of obeticholic acid is itching. Pruritus was the most common adverse event, occurring in 50% of patients receiving placebo and a similar percentage of patients at 10 mg; however, > 80% of patients receiving the two higher doses complained of pruritus, leading to discontinuation of the drug in up to 24% of cases.	New		c‡qvRbxq †idv‡iÝ ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i|	c‡qvRbxq †idv‡iÝ bv _vKq Av‡e`b bvgÄyi Kiv হল|
191.	Incepta Pharmaceuticals Ltd.	Obeticholic Acid 25mg Tablet Obeticholic Acid INN 25mg	-do-	-do-	New		c‡qvRbxq †idv‡iÝ ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i|	c‡qvRbxq †idv‡iÝ bv _vKq Av‡e`b bvgÄyi Kiv হল|
192.	Incepta Pharmaceuticals Ltd.	Obeticholic Acid 50mg Tablet Obeticholic Acid INN 50mg	-do-	-do-	New		c‡qvRbxq †idv‡iÝ ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i|	c‡qvRbxq †idv‡iÝ bv _vKq Av‡e`b bvgÄyi Kiv হল|
193.	Concord Pharmaceuticals Ltd.	Citric Acid Monohydrate 0.625% Linctus Citric acid Monohydrate BP 0.625 gm/100ml Alkalinizing Agents	The management of mild non-specific cough.	Contraindicaion: Glucose malabsorption syndrome. Glucose intolerance due to sucrose or isomaltase deficiency. Hypersensitivity to any of the constituents. Side effects: It is possible that citric acid ingested in large quantities or frequently may cause gastric irritation, or erosion of dental enamel	New	MHRA (Numark Paediatric Simple Linctus 31.25 mg/5 ml Oral solution)	Ab‡gv`b Kiv †h‡Z cv‡i|	Ab‡gv`b Kiv nj|
194.	Concord Pharmaceuticals Ltd.	Citric acid Monohydrate 2.5 % Linctus Citric acid Monohydrate BP 2.50mg/100ml Alkalinizing Agents	The management of mild non-specific cough.	Contraindicaion: Glucose malabsorption syndrome. Glucose intolerance due to sucrose or isomaltase deficiency. Hypersensitivity to any of the constituents. Side effects: It is possible that citric acid ingested in large quantities or frequently may cause gastric irritation, or erosion of dental enamel	New	MHRA (Numark Simple Linctus BP 125mg/5 ml Oral solution)	Ab‡gv`b Kiv †h‡Z cv‡i|	Ab‡gv`b Kiv nj|
195.	ACI Ltd., Narayanganj	Acetaminophen 300 mg + Hydrocodone Bitartrate 10mg Tablet Acetaminophen USP 300mg + Hydrocodone Bitartrate USP 10 mg	Do	Contraindication:This product should not be administered to patients who have previously exhibited hypersensitivity tohydrocodone or acetaminophen. patients known to be hypersensitive to other opioids may exhibit cross sensitivity to hydrocodone	New	USFDA	c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i|	c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv হল|

		Analgesic		Side-effects: The most frequently reported adverse reactions are lightheadedness, dizziness, sedation, nausea andvomiting. These effects seem to be more prominent in ambulatory than in nonambulatory patients, andsome of these adverse reactions may be alleviated if the patient lies down. Other adverse reactions include: Central nervous system: drowsiness, mental clouding, lethargy, impairment of mental and physical performance, anxiety, fear, dysphoria, psychic dependence, mood changes. Gastrointestinal system: prolonged administration of hydrocodone bitartrate and acetaminophen tabletsmay produce constipation. Genitourinary system: ureteral spasm, spasm of vesical sphincters and urinary retention have beenreported with opiates. Respiratory depression: Hydrocodone Bitartrate may produce dose-related respiratory depression byacting directly on the brain stem respiratory centers. Special senses: cases of hearing impairment or permanent loss have been reported predominantly inpatients with chronic overdose. Dermatological: skin rash, pruritus. WARNINGS Hepatotoxicity: Acetaminophen has been associated with cases of acute liver failure, at times resulting in liver transplant and death. Most of the cases of liver injury are associated with the use of acetaminophen at doses that exceed 4000 milligrams per day, and often involve more than one acetaminophen containing product. The excessive intake of acetaminophen may be intentional to cause self-harm or unintentional as patients attempt to obtain more pain relief or unknowingly take other				

				acetaminophen-containing products. The risk of acute liver failure is higher in individuals with underlying liver disease and in individuals who ingest alcohol while taking acetaminophen. Instruct patients to look for acetaminophen or APAP on package labels and not to use more than one product that contains acetaminophen. Instruct patients to seek medical attention immediately upon ingestion of more than 4000 milligrams of acetaminophen per day, even if they feel well. Serious skin reactions: Rarely, acetaminophen may cause serious skin reactions such as acute generalized exanthematous pustulosis (AGEP), Stevens - Johnson syndrome (SJS), and toxic epidermal necrolysis (TEN), which can be fatal. Patients should be informed about the signs of serious skin reactions, and use of the drug should be discontinued at the first appearance of skin rash or any other sign of hypersensitivity. Hypersensitivity/anaphylaxis: There have been post-marketing reports of hypersensitivity and anaphylaxis associated with use of acetaminophen. Clinical signs included swelling of the face, mouth, and throat, respiratory distress, urticaria, rash, pruritus, and vomiting. There were infrequent reports of life-threatening anaphylaxis requiring emergency medical attention. Instruct patients to discontinue this combination immediately and seek medical care if they experience these symptoms. Do not prescribe this combination for patients with acetaminophen allergy. Respiratory Depress ion: At high doses or in sensitive patients, hydrocodone may produce doserelated respiratory depression by acting directly on				

				the brain stem respiratory center. Hydrocodone also affects the center that				
196.	ACI Ltd., Narayanganj	Acetaminophen 300 mg + Hydrocodone Bitartrate 5mg Tablet Acetaminophen USP 300mg + Hydrocodone Bitartrate USP 5 mg Analgesic	-DO-	Do-	New	USFDA	*c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i|*	*c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv* हल|
197.	ACI Ltd., Narayanganj	Acetaminophen 300mg + Hydrocodone Bitartrate 7.5mg Tablet Acetaminophen USP 300mg + Hydrocodone Bitartrate USP 7.5 mg Analgesic	-do-	-Do-	New	USFDA	*c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i|*	*c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv* हल|
198.	ACI Ltd., Narayanganj	Acetaminophen 325 mg + Hydrocodone Bitartrate 5 mg Tablet Acetaminophen USP 325mg + Hydrocodone Bitartrate USP 5 mg Analgesic	Do	-do-	New	USFDA	*c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i|*	*c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv* हल|
199.	ACI Ltd., Narayanganj	Acetaminophen 325 mg + Hydrocodone Bitartrate 10mg Tablet Acetaminophen USP 325mg + Hydrocodone Bitartrate USP 10 mg Analgesic	Do	-do-	New	USFDA	*c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i|*	*c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv* हल|
200.	ACI Ltd., Narayanganj	Acetaminophen 325 mg + Hydrocodone bitartrate 2.5mg Tablet Acetaminophen USP 325mg + Hydrocodone bitartrate USP 2.5	-do-	-do-	New	USFDA	*c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i|*	*c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv* हल|

		mg Analgesic						
201.	ACI Ltd., Narayanganj	Acetaminophen 325 mg + Hydrocodone Bitartrate 7.5mg Tablet Acetaminophen USP 325mg + Hydrocodone Bitartrate USP 7.5 mg Analgesic	-do-	-do-	New	USFDA	c‡qvRb ‡bB wea‡q Av‡e`b bv gÄÿ Kiv †h‡Z cv‡i	c‡qvRb ‡bB wea‡q Av‡e`b bv gÄÿ Kiv হল
202.	Incepta Pharmaceuticals Ltd.	Glucosamine HCl 750mg + Chrondroitin Sulfate 400mg + Methyl Sulfonyl Methane (MSM) 375mg + Vitamin D3 1000IU Tablet Glucosamine HCl USP 750mg + Chrondroitin Sulfate BP 400mg + Methyl Sulfonyl Methane (MSM) USP 375mg + Vitamin D3 BP 1000IU Analgesic	Indicated for the treatment of osteoarthritis of knee, hip, spine, hand, and other locations. It is also beneficial in rheumatoid arthritis, sport injuries, migraine, different skin problems (e.g., psoriasis), vascular complications (e. g., atherosclerosis), kidney stones, and inflammatory bowel disease (e.g., ulcerative colitis, leaky gut syndrome).	Contraindication: There are no known contraindications for Glucosamine and Chondroitin. But proven hypersensitivity (e. g. allergic to shellfish or sulfur) to Glucosamine and Chondroitin is a contraindication. Side effect: Both Glucosamine and Chondroitin Sulfates are virtually nontoxic. Side effects are rare and are limited to stomach upset, nausea or diarrhea. These usually disappear when the tablet is taken with meals.	New		c‡qvRb ‡bB wea‡q Av‡e`b bv gÄÿ Kiv †h‡Z cv‡i	c‡qvRb ‡bB wea‡q Av‡e`b bv gÄÿ Kiv হল
203.	Incepta Pharmaceuticals Ltd.	Glucosamine Sulfate USP 1000mg + Chrondroitin Sulfate BP 800mg + MSM USP 500mg + Vitamin C BP 200mg/15ml Syrup Glucosamine Sulfate USP 1000mg + Chrondroitin Sulfate BP 800mg + MSM USP 500mg + Vitamin C BP 200mg/15ml Analgesic	-do-	-do-	New		c‡qvRb ‡bB wea‡q Av‡e`b bv gÄÿ Kiv †h‡Z cv‡i	c‡qvRb ‡bB wea‡q Av‡e`b bv gÄÿ Kiv হল
204.	Ziska Pharmaceuticals Ltd.	Oxycodone HCl 10mg + Acetaminophen 325 mg Tablet Oxycodone HCl USP 10mg + Acetaminophen USP 325 mg Analgesic	It is indicated for the relief of moderate to moderately severe pain.	Contraindications: This tablet should not be administered to patients with known hypersensitivity to oxycodone, acetaminophen, or any other component of this product. Oxycodone is contraindicated in any situation where opioids are contraindicated including patients with significant respiratory depression (in unmonitored settings or the absence of resuscitative equipment) and	Acetaminophen 500mg Tablet	USFDA	c‡qvRb ‡bB wea‡q Av‡e`b bv gÄÿ Kiv †h‡Z cv‡i	c‡qvRb ‡bB wea‡q Av‡e`b bv gÄÿ Kiv হল

				patients with acute or severe bronchial asthma or hypercarbia. Oxycodone is contraindicated in the setting of suspected or known paralytic ileus. Side effects: depression, apnea, respiratory arrest, circulatory depression, hypotension, and shock The most frequently observed non-serious adverse reactions include lightheadedness, dizziness, drowsiness or sedation, nausea, and vomiting. These effects seem to be more prominent in ambulatory than in nonambulatory patients, and some of these adverse reactions may be alleviated if the patient lies down. Other adverse reactions include euphoria, dysphoria, constipation, and pruritus. Hypersensitivity reactions may include: Skin eruptions, urticarial, erythematous skin reactions. Hematologic reactions may include: Thrombocytopenia, neutropenia, pancytopenia, hemolytic anemia. Rare cases of agranulocytosis has likewise been associated with acetaminophen use. In high doses, the most serious adverse effect is a dose-dependent, potentially fatal hepatic necrosis. Renal tubular necrosis and hypoglycemic coma also may occur				
205.	Ziska Pharmaceuticals Ltd.	Oxycodone HCl 5.0mg + Acetaminophen 325 mg Tablet Oxycodone HCl USP 5.0mg + Acetaminophen USP 325 mg Analgesic	-do-	-do-	Acetaminophen 500mg Tablet	USFDA	c‡qvRb ‡bB wea‡q Av‡e`b bv gÄÿ Kiv †h‡Z cv‡i	c‡qvRb ‡bB wea‡q Av‡e`b bv gÄÿ Kiv হল।
206.	Ziska Pharmaceuticals Ltd.	Oxycodone HCl5.0mg + Ibuprofen 400mg Tablet Oxycodone HCl USP 5.0mg + Ibuprofen USP 400mg	Carefully consider the potential benefits and risks of this tablet and other treatment options before deciding to use. Use the lowest effective dose for the shortest	Contraindications: This Tablet should not be administered to patients who have previously exhibited hypersensitivity to oxycodone HCl, ibuprofen, or any of it's components. It	Ibuprofen 200mg/ 400 mg Tablet	USFDA	c‡qvRb ‡bB wea‡q Av‡e`b bv gÄÿ Kiv †h‡Z cv‡i	c‡qvRb ‡bB wea‡q Av‡e`b bv gÄÿ Kiv হল।

		Analgesic	duration consistent with individual patient treatment goals. This tablet is indicated for the short term (no more than 7 days) management of acute, moderate to severe pain.	should not be administered in any situation where opioids are contraindicated. This includes patients with significant respiratory depression (in unmonitored settings or the absence of resuscitative equipment) and patients with acute or severe bronchial asthma or hypercarbia. Patients known to be hypersensitive to other opioids may exhibit crosssensitivity to oxycodone. Combunox is contraindicated in any patient who has or is suspected of having paralytic ileus. It should not be given to patients who have experienced asthma, urticaria, or allergic-type reactions after taking aspirin or other NSAIDs. Severe anaphylactoid reactions to NSAIDs, some of which were fatal, have been reported in such patients. It is contraindicated for the treatment of peri-operative pain in the setting of coronary artery bypass graft (CABG) surgery. Side effects: Cardiovascular Thrombotic Events, Congestive Heart Failure and Edema Hypertension, Gastrointestinal Effects - Risk of Ulceration, Bleeding, and Perforation, Misuse Abuse and Diversion of Opioids, Respiratory Depression, Hypotensive Effect, Head Injury and Increased Intracranial Pressure, Acute Abdominal Conditions, Anaphylactoid Reactions, Advanced Renal Disease, Skin Reactions.				
207.	Ziska Pharmaceuticals Ltd.	Oxycodone HCl 7.5mg + Acetaminophen 325 mg Tablet Oxycodone HCl USP 7.5mg + Acetaminophen USP 325 mg	It is indicated for the relief of moderate to moderately severe pain.	Contraindications: This tablet should not be administered to patients with known hypersensitivity to oxycodone, acetaminophen, or any other component of this product. Oxycodone is contraindicated in any situation where	Acetaminophen 500mg Tablet	USFDA	c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i|	c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv হল|

		Analgesic		opioids are contraindicated including patients with significant respiratory depression (in unmonitored settings or the absence of resuscitative equipment) and patients with acute or severe bronchial asthma or hypercarbia. Oxycodone is contraindicated in the setting of suspected or known paralytic ileus. Side effects: depression, apnea, respiratory arrest, circulatory depression, hypotension, and shock The most frequently observed non-serious adverse reactions include lightheadedness, dizziness, drowsiness or sedation, nausea, and vomiting. These effects seem to be more prominent in ambulatory than in nonambulatory patients, and some of these adverse reactions may be alleviated if the patient lies down. Other adverse reactions include euphoria, dysphoria, constipation, and pruritus. Hypersensitivity reactions may include: Skin eruptions, urticarial, erythematous skin reactions. Hematologic reactions may include: Thrombocytopenia, neutropenia, pancytopenia, hemolytic anemia. Rare cases of agranulocytosis has likewise been associated with acetaminophen use. In high doses, the most serious adverse effect is a dose-dependent, potentially fatal hepatic necrosis. Renal tubular necrosis and hypoglycemic coma also may occur				
208.	Square Formulations Ltd., Gorai, Tangail	Paracetamol 500mg + Diphenhydramine HCl 25mg Tablet Paracetamol BP 500mg + Diphenhydramine HCl BP 25mg **Analgesic + Antihistamine**	For the short term treatment of bedtime pain, for example rheumatic and muscle pain, backache, neuralgia, toothache, migraine, headache and period pain which is causing difficulty in getting to sleep.	Contra-indication: Hypersensitivity to paracetamol, diphenhydramine hydrochloride or other constituents. Porphyria. Antihistamines are contraindicated in premature infants or neonates who have increased susceptibility to antimuscarinic effects.	Diphenhydramine 50mg Tablet	MHRA	*c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z c‡i***।**	*c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv* হল।

				Side-effects: Like all medicines, it can have side effects, but not everybody gets them. Older people are more prone to these side effects. When using this product you may experience: • Drowsiness, dizziness, tiredness, blurred vision, or difficulty concentrating, Dry mouth. Stop taking this medicine and tell your doctor immediately if you experience: • Allergic reactions which may be severe such as skin rash and itching sometimes with swelling of the mouth or face or shortness of breath • Chest tightness or thickening of phlegm, • Difficulty in passing urine, headaches, • Skin rash or peeling or mouth ulcers, • Upset Stomach • Breathing problems. These are more likely if you have experienced them before when taking other painkillers (such as ibuprofen and aspirin), • Seizures or difficulty of muscle coordination • Changes in heart rhythm • Unexplained bruising or bleeding.				
209.	Delta Pharma Ltd.	Sacubitril 49mg + Valsartan 51mg Film Coated Tablet Sacubitril INN 49 mg + Valsartan USP 51 mg Angiotensin II receptor blocker	This is a combination of sacubitril, a neprilysin inhibitor, and valsartan, an angiotensin II receptor blocker, indicated to reduce the risk of cardiovascular death and hospitalization for heart failure in patients with chronic heart failure (NYHA Class II-IV) and reduced ejection fraction.	Contra-indications: Hypersensitivity to any component. History of angioedema related to previous ACE inhibitor or ARB therapy. Concomitant use with ACE inhibitors Concomitant use with aliskiren in patients with diabetes. Side-effect: Adverse reactions occurring ≥5% are hypotension, hyperkalemia, cough, dizziness, and renal failure. WARNINGS AND PRECAUTIONS Observe for signs and symptoms of angioedema and hypotension. Monitor renal function and potassium in susceptible patients.	Valsartan 80mg/16mg Tablet	USFDA	Abÿgv`b Kiv †h‡Z cv‡i	Abÿgv`b Kiv nj
210.	Delta Pharma Ltd.	Sacubitril 97mg + Valsartan 103mg Film Coated Tablet Sacubitril INN 97mg + Valsartan	This is a combination of sacubitril, a neprilysin inhibitor, and valsartan, an angiotensin II receptor blocker, indicated to reduce the risk of	Contra-indications: Hypersensitivity to any component. History of angioedema related to previous ACE inhibitor or ARB therapy. Concomitant use with ACE inhibitors.	Valsartan 80mg/16mg Tablet	USFDA	Abÿgv`b Kiv †h‡Z cv‡i	Abÿgv`b Kiv nj

		USP 103mg Angiotensin II receptor blocker	cardiovascular death and hospitalization for heart failure in patients with chronic heart failure (NYHA Class II-IV) and reduced ejection fraction.	Concomitant use with aliskiren in patients with diabetes. Side-effect: Adverse reactions occurring ≥5% are hypotension, hyperkalemia, cough, dizziness, and renal failure. WARNINGS AND PRECAUTIONS Observe for signs and symptoms of angioedema and hypotension. Monitor renal function and potassium in susceptible patients.				
211.	Square Pharmaceuticals Ltd., Pabna Unit, Salgaria, Pabna	Calcium Carbonate (Heavy) 1000.00mg eq. to 400mg Elemental Calcium + Simethicone 100 DC Ph. Gr. 100mg eq. to 60mg Simethcone Tablet Calcium Carbonate (Heavy) BP 1000.00mg eq. to 400mg Elemental Calcium + Simethicone 100 DC Ph. Gr. 100mg eq. to 60mg Simethcone USP Tablet **Antacid**	For the relieve of acid digestion, heartburn, sour stomach, upset of stomach associated with these symptoms, bloating and pressure commonly referred to as gas.	Contraindications: Constipation, diarrhoea. ADR/Side effects: Constipation, diarrhoea.	New		c‡qvRb ‡bB wea‡q Av‡e`b bv gÄÿ Kiv †h‡Z cv‡i	c‡qvRb ‡bB wea‡q Av‡e`b bv gÄÿ Kiv হল।
212.	Square Pharmaceuticals Ltd., Pabna Unit, Salgaria, Pabna	Sodium Alginate 250mg + Sodium Bicarbonate 106.500mg + Calcium Carbonate (Heavy) 187.500mg Tablet Sodium Alginate USP 250mg + Sodium Bicarbonate BP 106.500mg + Calcium Carbonate (Heavy) BP 187.500mg Tablet Gastrointestinal agent (Alginate + Antacid)	It is indicated for the treatment of gastro-oesophageal reflux i.e. acid regurgitation, heartburn, indigestion and for symptoms of excess stomach acid (hyperacidity) It acts in a dual mechanism mood, quickly neutralizes excess stomach acid and also forms a protective layer over stomach content	Contraindications: Hypersensitivity to the active substances or to any of the excipients, including the esters of hydroxybenzoates (parabens). Side effects: Very rarely (<1/10,000) patients sensitive to the ingredients may develop allergic manifestations such as urticaria or bronchospasm, anaphylactic or anaphylactoid reactions	New	MHRA	c‡qvRb ‡bB wea‡q Av‡e`b bv gÄÿ Kiv †h‡Z cv‡i	c‡qvRb ‡bB wea‡q Av‡e`b bv gÄÿ Kiv হল।
213.	Square Pharmaceuticals Ltd., Pabna Unit, Salgaria, Pabna Beacon Pharmaceutical Ltd.,	Sodium Alginate 5gm + Sodium Bicarbonate 2.130gm + Calcium Carbonate (light) 3.250gm/100ml Suspension Sodium Alginate USP 5gm + Sodium Bicarbonate BP 2.130gm + Calcium Carbonate (light) BP 3.250gm/100ml Suspension	It is indicated for the treatment of gastro-oesophageal reflux i.e. acid regurgitation, heartburn, indigestion and for symptoms of excess stomach acid (hyperacidity) It acts in a dual mechanism mood, quickly neutralizes excess stomach acid and also forms a protective	Contraindications: Hypersensitivity to the active substances or to any of the excipients, including the esters of hydroxybenzoates (parabens). Side effects: Very rarely (<1/10,000) patients sensitive to the ingredients may develop allergic manifestations such as urticaria or bronchospasm, anaphylactic or anaphylactoid reactions	New	MHRA	c‡qvRb ‡bB wea‡q Av‡e`b bv gÄÿ Kiv †h‡Z cv‡i	c‡qvRb ‡bB wea‡q Av‡e`b bv gÄÿ Kiv হল।

		Gastrointestinal agent (Alginate + Antacid)	layer over stomach content					
214.	GlaxoSmithKline Bangladesh Limted Fouzderhat Industrial Area PO: North Kattali Chittagong-4217	Soduim Bicarbonate 2.295gm + Citric Acid Anhydrous 2.165gm + Sodium Carbonte Anhydrous 0.495gm/5gm Powder in Sachet Soduim Bicardonate USP 2.295gm + Citric Acid Anhydrous USP 2.165gm + Sodium Carbonte Anhydrous USP 0.495gm/5.0 gm Antacid	It is effective for the symptomatic relief of: • Heartburn • Acid indigestion • Sour stomach	Contraindication: Persons on a sodium restricted diet e.g. those suffering from hypertension or congestive heart failure should not use this product unless directed by a doctor. Patients with impaired hepatic and renal Eno is contraindicated in patients with a prior hypersensitivity reaction to any ingredient of the preparation. Side Effects: Stomach/gut irritations which could cause wind or bloating.	New		c‡qvRb †bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i		c‡qvRb †bB weavq Av‡e`b bv gÄyi Kiv হল|
215.	Aristopharma Ltd.	Chlordiazepoxide HCl 5mg + Clidinium Bromide 2.5 mg Capsule Chlordiazepoxide HCl USP 5mg + Clidinium Bromide USP 2.5mg Antianxity/Anticholinergic	Adjunctive management of irritable bowel syndrome, peptic ulcer and other gastrointestinal disorders associated with hypersecretion, hypermotility and spasm and accompanied by anxiety or tension states	Contraindication: Hypersensitivity to chlordiazepoxide and /or Clidinium Bromide. Glaucoma, prostatic hypertrophy and benign bladder neck obstruction. Side-effects: As for chlordiazepoxide. In addition, clidinium Bromide may cause dryness of the mouth, blurred vision, urinary hesitancy; constipation particularly when combined with other spasmolytis and / or low residue diet.	New		c‡qvRb †bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i		c‡qvRb †bB weavq Av‡e`b bv gÄyi Kiv হল|
216.	Eskayef Bangladesh Limited	Etoricoxib 60mg + Thiocolchicoside 4mg FC Tablet Etoricoxib INN 60mg + Thiocolchicoside INN 4mg Antiarthritic + Muscle relaxant	It is used medically to treat orthopedic, traumatological and rheumatologic disorders as well as to treat muscular spasms. It is also used to control the pain and swelling suffered by individuals with four medical conditions: • Rheumatoid arthritis • Gout • Osteoarthritis • Ankylosing spondylitis	Contraindications: Inflammatory bowel disease, Severe congestive heart failure, Active peptic ulceration, Cerebrovascular disease, Lactation Children and adolescent < 16 yr. Side effects: Side effects of Etoricoxib and Thiocolchicoside are most likely to be minor. Like Constipation, diarrhea, dizziness. If you suffer from serious side effects, then concern your doctor as soon as possible.	Etoricoxib 60mg Tablet Etoricoxib 90mg Tablet Etoricoxib 120mg Tablet		c‡qvRb †bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i		c‡qvRb †bB weavq Av‡e`b bv gÄyi Kiv হল|
217.	Eskayef Bangladesh Limited.	Etoricoxib 60mg + Thiocolchicoside 8mg FC Tablet Etoricoxib INN 60mg +	-do-	-do-	Etoricoxib 60mg Tablet Etoricoxib 90mg Tablet Etoricoxib 120mg		c‡qvRb †bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i		c‡qvRb †bB weavq Av‡e`b bv gÄyi Kiv হল|

		Thiocolchicoside INN 8mg Antiarthritic + Muscle relaxant			Tablet			
218.	UniMed & UniHealth Mfg. Ltd., Gazipur	Fluticasone Propionate 125mcg + Formoterol Fumarate dehydrate 5mcg Actuation Metered Dose Inhalation Fluticasone Propionate BP 125mcg + Formoterol Fumarate dehydrate BP 5mcg per Actuation Antiasthmatic	For the treatment of prophylaxis of asthma	Contra-indication: Beta2 agonists should be used with caution in hyperthyroidism, cardiovascular disease, arrhythmias, susceptibility to QT-interval prolongation, and hypertension. Beta2 agonists should be used with caution in diabetes—monitor blood glucose (risk of ketoacidosis, especially when beta2 agonist given intravenously). Side-effects Side-effects of the beta2 agonists include fine tremor (particularly in the hands), nervous tension, headache, muscle cramps, and palpitation.	New	BNF-70 Page-231	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
219.	UniMed & UniHealth Mfg. Ltd., Gazipur	Fluticasone Propionate 250mcg + Formoterol Fumarate dehydrate 10mcg Actuation Metered Dose Inhalation Fluticasone Propionate BP 250mcg + Formoterol Fumarate dehydrate BP 10mcg per Actuation Antiasthmatic	-do-	-do-	New	BNF-70 Page-231	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
220.	UniMed & UniHealth Mfg. Ltd., Gazipur	Fluticasone Propionate 50mcg + Formoterol Fumarate dehydrate 5mcg Actuation Metered Dose Inhalation Fluticasone Propionate BP 50mcg + Formoterol Fumarate dehydrate BP 5mcg per Actuation Antiasthmatic	-do-	-do-	New	BNF-70 Page-231	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
221.	Ziska Pharmaceuticals Ltd. Aristopharma Ltd. Beximco Pharmaceuticals	Doxylamine Succinate 10mg+ Pyridoxine HCl 10mg Delayed Release Tablet Doxylamine Succinate USP	It is a fixed dose combination drug product of doxylamine succinate, an antihistamine, and pyridoxine hydrochloride, a Vitamin B6 analog, indicated for the treatment of	Contraindications: Known hypersensitivity to doxylamine succinate, other ethanolamine derivative antihistamines, pyridoxine hydrochloride or any inactive ingredient in the formulation Monoamine	New	USFDA (DR Tablet)	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

	Ltd., Tongi ,Gazipur	10mg + Pyridoxine HCl BP 10 mg **Anticholinergic + Vitamin**	nausea and vomiting of pregnancy in women who do not respond to conservative management.	oxidase (MAO) inhibitors. Side effects : The most common adverse reaction with this tablet (≥5 percent and exceeding the rate in placebo) is somnolence.				
222.	Ziska Pharmaceuticals Ltd.	Canagliflozin Hemihydrate 150mg + Metformin HCl 1000mg Tablet Canagliflozin Hemihydrate INN 153.0mg eq. to Canagliflozin 150mg + Metformin HCl BP 1000mg Antidiabetic	It is a sodium-glucose co-transporter 2 (SGLT2) inhibitor and biguanide combination product indicated as an adjunct to diet and exercise to improve glycemic control in adults with type 2 diabetes mellitus who are not adequately controlled on a regimen containing metformin or canagliflozin or in patients already being treated with both canagliflozin and metformin Limitation of use: Not for treatment of type 1 diabetes or diabetic ketoacidosis.	Contraindications: Renal impairment, ESRD, or on dialysis Metabolic acidosis, including diabetic ketoacidosis, History of serious hypersensitivity reaction to canagliflozin or metformin. Side effects Most common adverse reactions associated with canagliflozin (5% or greater incidence): female genital mycotic infections, urinary tract infection, and increased urination, Most common adverse reactions associated with metformin (5% or greater incidence) are diarrhea, nausea, vomiting, flatulence, asthenia, indigestion, abdominal discomfort, and headache.	New	USFDA	*c‡qvRb ‡bB wea‡q Av‡e`b bv gÄÿ Kiv †h‡Z cv‡i |*	*c‡qvRb ‡bB wea‡q Av‡e`b bv gÄÿ Kiv হল |*
223.	Ziska Pharmaceuticals Ltd.	Canagliflozin 150mg + Metformin HCl 500 mg Tablet Canagliflozin Hemihydrate INN 153mg eq. to Canagliflozin 150mg + Metformin HCl BP 500mg Antidiabetic	-do-	-do-	New	USFDA	*c‡qvRb ‡bB wea‡q Av‡e`b bv gÄÿ Kiv †h‡Z cv‡i |*	*c‡qvRb ‡bB wea‡q Av‡e`b bv gÄÿ Kiv হল |*
224.	Ziska Pharmaceuticals Ltd.	Canagliflozin 50mg + Metformin HCl 500 mg tablet Canagliflozin Hemihydrate INN 51.0mg eq. to Canagliflozin 50mg + Metformin HCl BP 500mg Antidiabetic	-do-	-do-	New	USFDA	*c‡qvRb ‡bB wea‡q Av‡e`b bv gÄÿ Kiv †h‡Z cv‡i |*	*c‡qvRb ‡bB wea‡q Av‡e`b bv gÄÿ Kiv হল |*
225.	Beximco Pharmaceuticals Ltd., Tongi ,Gazipur	Empagliflozin 5mg + Metformin HCl 1000mg Tablet Empagliflozin INN 5mg+ Metformin HCl BP 1000mg	It is indicated as an adjunct to diet and exercise to improve glycemic control in adults with type 2 diabetes mellitus who are not adequately controlled on a regimen containing empagliflozin or metformin.	Contraindications: Renal impairment which may also result from conditions such as cardiovascular collapse, acute myocardial infarction, and septicemia; end stage renal disease (ESRD) or patients on dialysis,	New	USFDA	*c‡qvRb ‡bB wea‡q Av‡e`b bv gÄÿ Kiv †h‡Z cv‡i |*	*c‡qvRb ‡bB wea‡q Av‡e`b bv gÄÿ Kiv হল |*

		Antidiabetic		Side effects: Common side effect are including; Lactic Acidosis, Hypotension, Impairment in Renal Function, Hypoglycemia with Concomitant Use with Insulin etc.				
226.	Beximco Pharmaceuticals Ltd., Tongi ,Gazipur	Empagliflozin 5mg + Metformin HCl 500mg Tablet Empagliflozin INN 5mg + Metformin HCl BP 500mg **Antidiabetic**	-do-	-do-	New	USFDA	*c‡qvRb ‡bB wea‡q Av‡e`b bv gÄyÿ Kiv †h‡Z cv‡i |*	*c‡qvRb ‡bB wea‡q Av‡e`b bv gÄyÿ Kiv* হল |*
227.	ACI Ltd., Narayanganj	Gliclazide 60 mg + Metformin HCl 500mg Extended Release Tablet Gliclazide BP 60 mg + Metformin HCl BP 500 mg **Antidiabetic**	Gliclazide plus Metformin is indicated for the non-insulin dependent diabetes mellitus; diabetes with or without obesity in adults	Contraindications: It is contraindicated in patients with hypersensitivity to any ingredients of this product. It is also contraindicated in Insulin-dependent diabetes mellitus, renal or hepatic failure, alcoholism, NIDDM complicated by severe ketosis and acidosis, diabetic precoma and coma, patients undergoing surgery, after severe trauma or during infections, chronic obstructive pulmonary disease, coronary heart disease, cardiac failure, peripheral vascular disease, pregnancy, known hypersensitivity to any of the ingredients. Side Effects: The most common side effects are Nausea, diarrhoea, gastric pain, constipation, vomiting, metallic taste in mouth, rash, pruritus, urticaria, erythema, flushing, headache and dizziness.	New		*K‡¤‡bkb c‡qvRb ‡bB wea‡q Av‡e`b bv gÄyÿ Kiv †h‡Z cv‡i |*	*K‡¤‡bkb c‡qvRb ‡bB wea‡q Av‡e`b bvgÄyÿ Kiv nj |*
228.	UniMed & UniHealth Mfg. Ltd., Gazipur Popular Pharmaceuticals Limited Incepta Pharmaceuticals Ltd. Acme Laboratories Ltd. Drug International Ltd.	Linagliptin 2.5mg + Metformin Hydrochloride 500mg Tablet Linagliptin INN 2.5mg + Metformin Hydrochloride BP 500mg Antidiabetic	It is a dipeptidyl peptidase-4 (DPP-4) inhibitor and biguanide combination product indicated as an adjunct to diet and exercise to improve glycemic control in adults with type 2 diabetes mellitus when treatment with both linagliptin and metformin is appropriate.	Contra-indications: Renal impairment Metabolic acidosis, including diabetic ketoacidosis,, Hypersensitivity to linagliptin or metformin Side effects: Adverse reactions reported in 5% of patients treated with Metformin + Linagliptin and more commonly than in patients treated with placebo are nasopharyngitis and diarrhea Hypoglycemia was more commonly reported in patients treated with the combination of Metformin + Linagliptin, and	New	USFDA	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

	Nipro JMI Pharma Ltd. Eskayef Bangladesh Limited.			SU compared with those treated with the combination of SU and metformin Pancreatitis was reported more often in patients randomized to linagliptin (1 per 538 person years versus zero in 433 person years for comparator).				
229.	UniMed & UniHealth Mfg. Ltd., Gazipur Incepta Pharmaceuticals Ltd. Acme Laboratories Ltd. Eskayef Bangladesh Limited. Drug International Ltd. Nipro JMI Pharma Ltd. Popular Pharmaceuticals Limited	Linagliptin 2.5mg + Metformin Hydrochloride 850mg Tablet Linagliptin INN 2.5mg + Metformin Hydrochloride BP 850mg Antidiabetic	-do-	-do-	New	USFDA, BNF-70 Page-597	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
230.	Incepta Pharmaceuticals Ltd. Nipro JMI Pharma Ltd. Eskayef Bangladesh Limited.	Linagliptin 2.5 mg +Metformin Hydrochloride 1000 mg Tablet Linagliptin INN 2.5 mg + Metformin Hydrochloride BP 1000 mg Antidiabetic	-do-	-do-	New	USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
231.	Square Pharmaceuticals Ltd., Pabna Unit, Salgaria, Pabna	Simvastatin 10 mg + Sitagliptin 100 mg FC Tablet Simvastatin USP 10 mg + Sitagliptin Phosphate Monohydrate INN 128.50mg eq. to 100 mg Sitagliptin	Sitagliptin: Sitagliptin is indicated as an adjunct to diet and exercise to improve glycemic control in adults with type 2 diabetes mellitus. Simvastatin: Reduce elevated total cholesterol (total-C), low-density lipoprotein cholesterol (LDL-C),	Contraindications: History of a serious hypersensitivity reaction, such as anaphylaxis or angioedema, to any component of this medication. Concomitant administration of strong CYP3A4 inhibitors (e.g., itraconazole, ketoconazole, posaconazole, HIV protease inhibitors, erythromycin, clarithromycin, telithromycin	New	USFDA	Kwc¤‡bkb c‡qvRb ‡bB wea|q Av‡e`b bv gÄyi Kiv †h‡Z cv‡i	Kwc¤‡bkb c‡qvRb ‡bB wea|q Av‡e`b bvgÄyi Kiv nj

		(Lipid lowering + Antidiabetic)	apolipoprotein B (Apo B) and triglycerides (TG) and to increase high-density lipoprotein cholesterol (HDL-C) in patients with primary hyperlipidemia (Fredrickson type IIa, heterozygous familial and nonfamilial) or mixed dyslipidemia (Fredrickson type IIb). Moreover Simvastatin is used,To reduce the risk of total mortality by reducing CHD deaths. To reduce the risk of non-fatal myocardial infarction and stroke. To reduce the need for coronary and non-coronary revascularization procedures.	and nefazodone).Concomitant administration of gemfibrozil, cyclosporine or diazole. Active liver disease, which may include unexplained persistent elevations in hepatic transaminase levels. ADR/Side effects: Sitagliptin: In controlled clinical studies as monotherapy and combination therapy with metformin, pioglitazone, or rosiglitazone and metformin, the overall incidence of adverse reactions, hypoglycemia, and discontinuation of therapy due to clinical adverse reactions with sitagliptin were similar to placebo. Simvastatin: The most common adverse reactions that led to treatment discontinuation were: gastrointestinal disorders (0.5%), myalgia (0.1%), and arthralgia (0.1%). The most commonly reported adverse reactions (incidence ≥5%) in simvastatin controlled clinical trials were: upper respiratory infections (9.0%), headache (7.4%), abdominal pain (7.3%), constipation (6.6%), and nausea (5.4%).				
232.	Healthcare Pharmaceutical ltd., Rajendrapur, Gazipur	Netupitant 300mg + Palonosetron 0.5 mg Capsule Netupitant INN 300mg + Palonosetron HCl INN 0.56 mg eq. to Palonosetron 0.5mg Antiemetic	Indicated for the prevention of acute & delayed nausea & vomiting associated with initial & repeat courses of cancer chemotherapy	Contraindication: Patients with a history of drug hypersensitivity to any of the ingredients in the product. Side Effects: Most common adverse reactions are headache, asthenia, dyspepsia, fatigue, constipation and erythema	New	USFDA	*Kw¤‡bkb c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i|*	*Kw¤‡bkb c‡qvRb ‡bB weavq Av‡e`b bvgÄyi Kiv nj|*
233.	Eskayef Bangladesh Limited	Albendazole 400mg + Ivermectin 6mg Tablet Albendazole BP 400mg + Ivermectin BP 6mg **Anthelmintic**	It is traditionally used against worms. It is mainly used in humans in the treatment of onchocerciasis, but is also effective against other worm infestations (strongyloidiasis), some epidermal parasitic skin diseases (scabies). It is effective against Flatworms,	Contraindications: Contraindicated in persons with a history of hypersensitivity. Side effects: Abdominal pain,	New		*Kw¤‡bkb c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i|*	*Kw¤‡bkb c‡qvRb ‡bB weavq Av‡e`b bvgÄyi Kiv nj|*

			Flukes, Tapeworm, Cysticercosis, Enterobiasis, Trichuriasis Ascariasis and Hookworm.					
234.	Square Pharmaceuticals Ltd., Pabna Unit, Salgaria, Pabna	Cetirizine HCl 5mg + Pseudoephedrine HCl 120mg Extended Release Tablet Cetirizine HCl BP 5mg + Pseudoephedrine HCl USP 120mg **Antihistamine + Decongestant**	Temporarily relieves symptoms due to hay fever or other upper respiratory allergies: Runny nose, sneezing, itchy, watery eyes, itching of the nose or throat, nasal congestion, reduces swelling of nasal passages, temporarily relieves sinus congestion and pressure, temporarily restores freer breathing through the nose.	Contraindications: It should not be used in patients having allergic reaction to this product or any of its ingredients or to an antihistamine containing hydroxyzine. It is also contraindicated in patetients receiving monoamine oxidase (MAO) inhibitor therapy. Side effects: Weakness, tremors (uncontrolled shaking), or sleep problems (insomnia); severe restless feeling, hyperactivity; extreme feeling of fear or confusion; increased blood pressure (severe headache, blurred vision, trouble concentrating, chest pain, numbness, seizure); problems with vision; or urinating less than usual or not at all. Less serious side effects may include: dizziness, drowsiness; tired feeling; dry mouth; nausea, stomach pain, constipation; problems with concentration; or ringing in your ears.	Cetirizine 10 mg Tablet; Pseudoephedrine 60mg Tablet	USFDA	*Kw¤‡bkb c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i|*	*Kw¤‡bkb c‡qvRb ‡bB weavq Av‡e`b bvgÄyi Kiv nj |*
235.	Square Pharmaceuticals Ltd., Pabna Unit, Salgaria, Pabna	Fexofenadine HCl 180mg + Pseudoephedrine HCl 240mg Extended release Bilayer Tablet Fexofenadine HCl BP 180mg + Pseudoephedrine HCl BP 240mg **Antihistamine + Decongestant**	It is indicated for the relief of symptoms associated with seasonal allergic rhinitis in adults and children 12 years of age and older. Symptoms treat ed effectively include sneezing, rhinorrhea, itchy nose/palate/ and/or throat, itchy/watery/red eyes, and nasal congestion.	Contra-indications: It should generally be avoided in patients with renal insufficiency. Due to its pseudoephedrine component, this is contraindicated in patients with narrow-angle glaucoma or urinary retention. It is also contraindicated in patetients receiving monoamine oxidase (MAO) inhibitor therapy. Side effects: Side effects of this combination include stimulation of the nervous system leading to nervousness, restlessness, excitability, dizziness, headache, fear, anxiety, tremor, and even hallucinations and convulsions (seizures).	60mg + 120mg ER Tablet	USFDA	*Kw¤‡bkb c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i|*	*Kw¤‡bkb c‡qvRb ‡bB weavq Av‡e`b bvgÄyi Kiv nj |*
236.	Opsonin Pharma Limited, Bagura Road, Barisal	Azelastine HCl 137 mcg + Fluticasone Propionate 50mcg/ 0.137ml Nasal Spray Azelastine HCl BP 137mcg +	This intranasal spray is indicated for the symptomatic relief of rhinitis.	Contraindications: Patients with hypersensitive to Fluticasone propionate and Azelastine hydrochloride should avoid this medication. Side effects: The most common side	New	USFDA, BNF -70 Page-985	*Ab‡gv`b Kiv †h‡Z cv‡i|*	*Ab‡gv`b Kiv nj |*

		Fluticasone Propionate BP 50mcg/0.137 ml **Antihistamine + Steroid**		effects of this combination nasal preparation include -changes in taste, nosebleeds and headache.				
237.	Popular Pharmaceuticals Limited	Ebastine 10mg + Montelukast 10mg Tablet Ebastine BP 10mg + Montelukast Sodium BP 10.40mg eq. to Montelukast 10mg **Antihistamine+ Antiasthmatic**	Ebastine and Montelukast combination is indicated for the prophylaxis and chronic treatment of Asthma, Exercise-Induced Bronchoconstriction & Allergic Rhinitis	Contraindications: Hypersensitivity to any component of this product Side effects: The most frequently occurring adverse effects of Ebastine are: headache, dry mouth, drowsiness, nausea, and insomnia. The most frequently occurring adverse effects of Montelukast are: headache; stomach pain, heartburn, upset stomach, nausea, diarrhea; tooth pain; tired feeling; fever, stuffy nose, sore throat, cough, hoarseness; or mild rash.	Ebastine 10mg Tablet Montelukast 10mg Tablet		Kw¤‡bkb c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i|	Kw¤‡bkb c‡qvRb ‡bB weavq Av‡e`b bvgÄyi Kiv nj|
238.	Popular Pharmaceuticals Limited	Bisoprolol Fumarate 2.50mg + Amlodipine 5mg Film Coated Tablet Bisoprolol Fumarate USP 2.50mg + Amlodipine Besilate BP 6.93mg eq. to Amlodipine 5mg **Antihypertensive**	It is indicated for the treatment of hypertension, alone or with other antihypertensive agents. (Amlodipine+Bisoprolol) may also be used as initial therapy in patients who are likely to need multiple antihypertensive agents to achieve their blood pressure goals. It is also used to treat angina pectoris, stable chronic heart failure.	Contraindications: Combination of Amlodipine and Bisoprolol is contraindicated in patients who are hypersensitive to any component of this product or to any of its ingredients. Side effects: The common side effects include edema, upper respiratory tract infection, hypotension, dizziness, headache, nausea, vomiting, diarrhoea, constipation, hypersensitivity reactions (itching, flush, rash) etc.	New		Kw¤‡bkb c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i|	Kw¤‡bkb c‡qvRb ‡bB weavq Av‡e`b bvgÄyi Kiv nj|
239.	Popular Pharmaceuticals Limited	Bisoprolol Fumarate 5.0 mg + Amlodipine 5.0mg Film Coated Tablet Bisoprolol Fumarate USP 5.00mg + Amlodipine Besilate BP 6.93mg eq. to Amlodipine 5mg Antihypertensive	-do-	-do-	New		Kw¤‡bkb c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i|	Kw¤‡bkb c‡qvRb ‡bB weavq Av‡e`b bvgÄyi Kiv nj|
240.	ACI Ltd., Narayanganj	Indapamide 1.5 mg + Amlodipine 5 mg Extended Release Tablet Indapamide BP 1.5 mg + Amlodipine Besilate BP 6.935mg eqv. to 5mg Amlodipine	It is indicated as substitution therapy for treatment of essential hypertension in patients already controlled with indapamide and amlodipine given concurrently at the same dose level.	Contraindications: Hypersensitivity to the active substances to other sulphonamides to dihydropyridine derivatives or to any of the excipients, severe renal failure (creatinine clearance below 30 ml/min), hepatic encephalopa, thy or severe impairment of liver function, hypokalaemia,	New		Kw¤‡bkb c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i|	Kw¤‡bkb c‡qvRb ‡bB weavq Av‡e`b bvgÄyi Kiv nj|

		Anti Hypertensive		lactation, severe hypotension, shock (including cardiogenic shock), obstruction of the outflow tract of the left ventricle (e.g., high grade aortic stenosis), haemodynamically unstable heart failure after acute myocardial infarction. Side Effects: The most commonly reported adverse reactions with indapamide and amlodipine given separately are somnolence, dizziness, headache, palpitations, flushing, abdominal pain, nausea, ankle swelling, oedema and fatigue.				
241.	Sun Pharmaceutical (Bangladesh) Ltd.	Olmesartan Medoxomil 20mg + Amlodipine 5mg + Hydrochlorothiazide 12.5mg Tablet Olmesartan Medoxomil BP 20 mg + Amlodipine Besylate 6.940 mg eq. to Amlodipine 5 mg + Hydrochlorothiazide BP 12.5 mg Antihypertensive	Indicated for the treatment of hypertension, to lower blood pressure. Lowering blood pressure reduces the risk of fatal and nonfatal cardiovascular events, primarily strokes and myocardial infarctions. The combination is not indicated for initial therapy.	Contraindications: Anuria: Hypersensitivity to sulfonamide-derived drugs. Do not co-administer aliskiren with this combination in patients with diabetes . Side Effects: Bloating or swelling of the face, arms, hands, lower legs, or feet rapid weight gain tingling of the hands or feet unusual weight gain or loss	New	USFDA	*K|¤‡bkb c‡qvRb ‡bB |eavq Av‡e`b bv gÄy¨ K i v †h‡Z cv‡i |*	*K|¤‡bkb c‡qvRb ‡bB |eavq Av‡e`b bvgÄy¨ K i v n j |*
242.	Sun Pharmaceutical (Bangladesh) Ltd.	Olmesartan Medoxomil 40mg + Amlodipine 5mg + Hydrochlorothiazide 12.5mg Tablet Olmesartan Medoxomil BP 20 mg + Amlodipine Besylate 6.940 mg eqv. to Amlodipine 5 mg + Hydrochlorothiazide BP 12.5 mg Antihypertensive	-do-	-do-	New	USFDA	*K|¤‡bkb c‡qvRb ‡bB |eavq Av‡e`b bv gÄy¨ K i v †h‡Z cv‡i |*	*K|¤‡bkb c‡qvRb ‡bB |eavq Av‡e`b bvgÄy¨ K i v n j |*
243.	ACI Ltd., Narayanganj	Olmesartan medoxomil 40mg + Hydrochlorothiazide 25mg Film Coated Tablet Olmesartan Medoxomil USP 40mg + Hydrochlorothiazide BP 25mg	Treatment of hypertension. This fixed dose combination is not indicated for initial therapy	Contraindications: This combination is contraindicated in patients who are hypersensitive to any component of this product. Because of Hydrochlorothiazide component it is also contraindicated in patients with anuria and other sulfonamide derived drugs. Side Effects: The common side-effects are nausea, headache, dizziness,	Hydrochlorothiazide 12.5 mg + Olmesartan Medoxomil 40 mg Tablet	USFDA	*K|¤‡bkb c‡qvRb ‡bB |eavq Av‡e`b bv gÄy¨ K i v †h‡Z cv‡i |*	*K|¤‡bkb c‡qvRb ‡bB |eavq Av‡e`b bvgÄy¨ K i v n j |*

		Anti hypertensive		hyperuricemia, upper respiratory tract infection and urinary tract infection. Other adverse effects are chest pain, back pain, peripheral edema, abdominal pain, dyspepsia, gastroenteritis, and diarrhea. Warning When pregnancy is detected, discontinue as soon as possible. Drugs that act directly on the renin -angiotensin system can cause injury and death to the developing fetus.				
244.	ACI Ltd., Narayanganj	Perindopril Arginine 10 mg + Indapamide 2.5mg + Amlodipine 10mg Film Coated Tablet Perindopril Arginine INN 10mg + Indapamide BP 2.5mg + Amlodipine Besilate BP 13.87mg eq. to 10mg Amlodipine Anti Hypertensive	It is indicated as substitution therapy for treatment of essential hypertension, in patients already controlled with perindopril/indapamide fixed dose combination and amlodipine, taken at the same dose level.	Contraindications: Dialysis patients, Patients with untreated decompensated, heart failure , Severe renal impairment (creatinine clearance below 30 mL/min), Moderate renal impairment (creatinine clearance below 60 mL/min) for doses containing 10mg/2.5mg of perindopril/ indapamide combination (i.e., Triplixam 10mg/2.5mg/5mg and 10mg/2.5mg/10mg), Hypersensitivity to the active substances, to other sulphonamides, to dihydropyridine derivatives, any other ACE-inhibitor or to any of the excipients, History of angioedema (Quincke's oedema) associated with previous ACE inhibitor therapy, Hereditary/idiopathic angioedema, Second and third trimesters of pregnancy ,Lactation), Hepatic encephalopathy, Severe hepatic impairment , Hypokalaemia , Severe hypotension , Shock including cardiogenic shock , Obstruction of the outflow, tract of the left ventricle (e.g. high grade aortic stenosis) , Haemodynamically unstable heart failure after acute myocardial infarction , Concomitant use with aliskiren in patients with diabetes mellitus or renal impairment (GFR < 60mL/min/1.73m2). Side Effects : The most commonly reported adverse reactions with perindopril, indapamide and amlodipine given separately are: dizziness,	Perindopril 4mg + Indapamide 1.25mg Tablet Perindopril 2.0mg + Indapamide 625mcg Tablet		*K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bv gÄy¨ K i v †h‡Z cv‡i |*	*K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bvgÄy¨ K i v n j |*

				headache, paresthesia, vertigo, somnolence, visual disturbances, tinnitus, palpitations, flushing, hypotension				
245.	ACI Ltd., Narayanganj	Perindopril arginine 10mg + Indapamide 2.5mg + Amlodipine 5mg Film Coated Tablet Perindopril Arginine INN 10mg eq. to perindopril 6.790mg + Indapamide BP 2.5mg + Amlodipine Besilate BP 6.935mg eqv. to 5mg Amlodipine Anti Hypertensive	-do-	-do-	Perindopril 4mg + Indapamide 1.25mg Tablet Perindopril 2.0mg + Indapamide 625mcg Tablet		K‡¤‡bkb c‡qvRb ‡bB wea‡q Av‡e`b bv gÄyi Kiv †h‡Z cv‡i	K‡¤‡bkb c‡qvRb ‡bB wea‡q Av‡e`b bvgÄyi Kiv nj
246.	ACI Ltd., Narayanganj	Perindopril arginine 2.5mg + Indapamide 0.625 mg + Amlodipine 5 mg Film Coated Tablet Perindopril Arginine INN 2.5mg eq. to perindopril 1.6975 mg + Indapamide BP 0.625 mg + Amlodipine Besilate BP 6.935mg eq. to 5mg Amlodipine Anti Hypertensive	It is indicated as substitution therapy for treatment of essential hypertension, in patients already controlled with perindopril/indapamide fixed dose combination and amlodipine, taken at the same dose level.	Contraindications: Dialysis patients, Patients with untreated decompensated, heart failure , Severe renal impairment (creatinine clearance below 30 mL/min), Moderate renal impairment (creatinine clearance below 60 mL/min) for doses containing 10mg/2.5mg of perindopril/ indapamide combination (i.e., Triplixam 10mg/2.5mg/5mg and 10mg/2.5mg/10mg), Hypersensitivity to the active substances, to other sulphonamides, to dihydropyridine derivatives, any other ACE-inhibitor or to any of the excipients, History of angioedema (Quincke's oedema) associated with previous ACE inhibitor therapy, Hereditary/idiopathic angioedema, Second and third trimesters of pregnancy ,Lactation), Hepatic encephalopathy, Severe hepatic impairment , Hypokalaemia , Severe hypotension , Shock including cardiogenic shock , Obstruction of the outflow, tract of the left ventricle (e.g. high grade aortic stenosis) , Haemodynamically unstable	New		K‡¤‡bkb c‡qvRb ‡bB wea‡q Av‡e`b bv gÄyi Kiv †h‡Z cv‡i	K‡¤‡bkb c‡qvRb ‡bB wea‡q Av‡e`b bvgÄyi Kiv nj

				heart failure after acute myocardial infarction , Concomitant use with aliskiren in patients with diabetes mellitus or renal impairment (GFR < 60mL/min/1.73m2). Side Effects : The most commonly reported adverse reactions with perindopril, indapamide and amlodipine given separately are: dizziness, headache, paresthesia, vertigo, somnolence, visual disturbances, tinnitus, palpitations, flushing, hypotension				
247.	ACI Ltd., Narayanganj	Perindopril Arginine 5mg + Indapamide 1.25mg + Amlodipine 5 mg Film Coated Tablet Perindopril Arginine INN 5mg eq. to 3.395mg + Indapamide BP 1.25mg + Amlodipine Besilate BP 6.935mg eq. to 5.0mg Amlodipine Anti Hypertensive	It is indicated as substitution therapy for treatment of essential hypertension, in patients already controlled with perindopril/indapamide fixed dose combination and amlodipine, taken at the same dose level.	Contraindications: Dialysis patients, Patients with untreated decompensated, heart failure , Severe renal impairment (creatinine clearance below 30 mL/min), Moderate renal impairment (creatinine clearance below 60 mL/min) for doses containing 10mg/2.5mg of perindopril/ indapamide combination (i.e., Triplixam 10mg/2.5mg/5mg and 10mg/2.5mg/10mg), Hypersensitivity to the active substances, to other sulphonamides, to dihydropyridine derivatives, any other ACE-inhibitor or to any of the excipients, History of angioedema (Quincke's oedema) associated with previous ACE inhibitor therapy, Hereditary/idiopathic angioedema, Second and third trimesters of pregnancy ,Lactation), Hepatic encephalopathy, Severe hepatic impairment , Hypokalaemia , Severe hypotension , Shock including cardiogenic shock , Obstruction of the outflow, tract of the left ventricle (e.g. high grade aortic stenosis) , Haemodynamically unstable heart failure after acute myocardial	New		*K‡¤‡bkb c‡qvRb ‡bB wea‡q Av‡e`b bv gÄyi Kiv †h‡Z cv‡i |*	*K‡¤‡bkb c‡qvRb ‡bB wea‡q Av‡e`b bvgÄyi Kiv nj |*

				infarction , Concomitant use with aliskiren in patients with diabetes mellitus or renal impairment (GFR < 60mL/min/1.73m2). Side Effects : The most commonly reported adverse reactions with perindopril, indapamide and amlodipine given separately are: dizziness, headache, paresthesia, vertigo, somnolence, visual disturbances, tinnitus, palpitations, flushing, hypotension				
248.	ACI Ltd., Narayanganj	Perindopril Arginine 5 mg + Indapamide 1.25 mg + Amlodipine 10 mg Film Coated Tablet Perindopril Arginine INN 5mg eq. to perindopril 3.395 mg + Indapamide BP 1.25 mg + Amlodipine Besilate BP 13.87mg eq. to 10mg Amlodipine Anti Hypertensive	It is indicated as substitution therapy for treatment of essential hypertension, in patients already controlled with perindopril/indapamide fixed dose combination and amlodipine, taken at the same dose level.	Contraindications: Dialysis patients, Patients with untreated decompensated, heart failure , Severe renal impairment (creatinine clearance below 30 mL/min), Moderate renal impairment (creatinine clearance below 60 mL/min) for doses containing 10mg/2.5mg of perindopril/ indapamide combination (i.e., Triplixam 10mg/2.5mg/5mg and 10mg/2.5mg/10mg), Hypersensitivity to the active substances, to other sulphonamides, to dihydropyridine derivatives, any other ACE-inhibitor or to any of the excipients, History of angioedema (Quincke's oedema) associated with previous ACE inhibitor therapy, Hereditary/idiopathic angioedema, Second and third trimesters of pregnancy ,Lactation), Hepatic encephalopathy, Severe hepatic impairment , Hypokalaemia , Severe hypotension , Shock including cardiogenic shock , Obstruction of the outflow, tract of the left ventricle (e.g. high grade aortic stenosis) , Haemodynamically unstable heart failure after acute myocardial infarction , Concomitant use with aliskiren	Perindopril 4mg + Indapamide 1.25mg Tablet Perindopril 2.0mg + Indapamide 625mcg Tablet		*K‖¤‡bkb c‡q‖Rb ‡bB ‖ea‖q Av‡e`b b‖ gÄÿ K i‖ †h‡Z c‖‡i |*	*K‖¤‡bkb c‡q‖Rb ‡bB ‖ea‖q Av‡e`b b‖gÄÿ K i‖ n j |*

				in patients with diabetes mellitus or renal impairment (GFR < 60mL/min/1.73m2). Side Effects : The most commonly reported adverse reactions with perindopril, indapamide and amlodipine given separately are: dizziness, headache, paresthesia, vertigo, somnolence, visual disturbances, tinnitus, palpitations, flushing, hypotension				
249.	Pharmasia Limited.	Telmisartan 80mg + Amlodipine 5mg Tablet Telmisartan BP 80mg + Amlodipine BP 5mg Antihypertensive	It is an angiotensin II receptor blocker (ARB) and a dihydropyridine calcium channel blocker (DHP-CCB) combination product indicated for the treatment of hypertension alone or with other antihypertensive agents to lower blood pressure. Lowering blood pressure reduces the risk of fatal and nonfatal cardiovascular events, primarily strokes and myocardial infarctions. This tablets are indicated as initial therapy in patients likely to need multiple antihypertensive agents to achieve their blood pressure goals	Contraindication: Known hypersensitivity (e.g., anaphylaxis or angioedema) to telmisartan, amlodipine or any other component of this product. Do not co-administer aliskiren with TWYNSTA in patients with diabetes Side effects: The most common reasons for discontinuation of therapy with tablets were peripheral edema, dizziness, and hypotension, each leading to discontinuation of ≤0.5% of its treated patients. Adverse reactions that occurred at a ≥2% higher incidence on this tablets than placebo were peripheral edema (4.8% vs 0%), dizziness (3.0% vs 2.2%), and back pain (2.2% vs 0%).	Telmisartan 40mg + Amlodipine 5mg Tablet; Telmisartan 80mg Tablet	USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
250.	Acme Laboratories Ltd.	Sacubitril 24 mg + Valsartan 26 mg Tablet. Sacubitril INN 24 mg + Valsartan USP 26 mg Cardiovascular Agent (Angiotensin II Receptor Antagonist)	Indicated to reduce the risk of cardiovascular death and hospitalization for heart failure in patients with chronic heart failure (NYHA Class II-IV) and reduced ejection fraction. Sacubitril and Valsartan Tablet is usually administered in conjunction with other heart failure therapies, in place of an ACE inhibitor	Contraindication: Hypersensitivity to any component; History of angioedema related to previous ACE inhibitor or ARB therapy. Concomitant use with ACE inhibitors. Concomitant use with aliskiren in patients with diabetes. Side effect: Adverse reactions occurring ≥5% are hypotension, hyperkalemia, cough, dizziness, and renal failure.	New	USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

251.	Acme Laboratories Ltd.	Sacubitril 49 mg + Valsartan 51 mg Tablet. Sacubitril INN 49 mg + Valsartan USP 51 mg Cardiovascular Agent (Angiotensin II Receptor Antagonist)	-do-	-do-	New	USFDA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
252.	Acme Laboratories Ltd.	Sacubitril 97 mg + Valsartan 103 mg Tablet Sacubitril INN 97 mg + Valsartan USP 103 mg Cardiovascular Agent (Angiotensin II Receptor Antagonist)	-do-	-do-	New	USFDA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
253.	Delta Pharma Ltd. Incepta Pharmaceuticals Ltd.	Sacubitril 24 mg + Valsartan 26mg Film Coated Tablet Sacubitril INN 24 mg + Valsartan USP 26 mg Angiotensin II receptor blocker	Indicated to reduce the risk of cardiovascular death and hospitalization for heart failure in patients with chronic heart failure (NYHA Class II-IV) and reduced ejection fraction. Sacubitril and Valsartan Tablet is usually administered in conjunction with other heart failure therapies, in place of an ACE inhibitor	Contraindication: Hypersensitivity to any component; History of angioedema related to previous ACE inhibitor or ARB therapy. Concomitant use with ACE inhibitors. Concomitant use with aliskiren in patients with diabetes. Side effect: Adverse reactions occurring ≥5% are hypotension, hyperkalemia, cough, dizziness, and renal failure.	Valsartan 80mg/16mg Tablet	USFDA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
254.	Incepta Pharmaceuticals Ltd.	Sacubitril 49 mg + Valsartan 51 mg Tablet Sacubitril INN 49 mg + Valsartan USP 51 mg Antihypertensive (Angiotensin II receptor blocker)	-do-	-do-	New	USFDA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
255.	Incepta Pharmaceuticals Ltd.	Sacubitril 97 mg + Valsartan 103 mg Tablet Sacubitril INN 97 mg + Valsartan USP 103 mg Antihypertensive (Angiotensin II receptor blocker)	Do		New	USFDA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj

256.	Eskayef Bangladesh Limited.	Omeprazole 20mg + Domperidone 10mg Capsule Omeprazole USP 20mg + Domperidone BP 10mg Antiulcerant + Antiemetic	Dyspepsia, Gastro-oesophageal reflux disease.	Contraindications: • Pregnancy–Domperidone is contraindicated in conditions associated with rise in prolactin level. • Omeprazole is contraindicated in hypersensitive patients. Side effects: • Domperidone-Dry mouth, itching, headache, diarrhoea. • Omeprazole-Anaemia, UTI, skin rash, pruritus, vertigo.	Omeprazole 20mg Capsule Domperidone 10mg Tablet		K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bv gÄyi Ki‖ †h‡Z cv‡i	K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bvgÄyi Ki‖ nj	
257.	Eskayef Bangladesh Limited	Pantoprazole 40mg + Domperidone 10mg Capsule Pantoprazole Sodium sesquihydrate USP 45.232mg eq. to 40mg Pantoprazole + Domperidone BP 10mg Antiulcerant + Antiemetic	Antacids, Anti-reflux Agents & Antiulcerants / GIT Regulators, Antiflatulents & Anti-Inflammatories.	Contraindications: • Pantoprazole is contraindicated in heptic-impairment; hypersensitivity. • Domperidone is contraindicated in the history of hypersensitivity to dimenhydrinate or related compounds. Side effects: Nausea, drowsiness, flatulence and dry mouth.	Pantoprazole 40mg Tablet Domperidone 10mg Tablet		K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bv gÄyi Ki‖ †h‡Z cv‡i	K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bvgÄyi Ki‖ nj	
258.	Popular Pharmaceuticals Limited Aristopharma Ltd.	Esomeprazole 20 mg + Domperidone 30 mg Capsule Esomeprazole Magnesium Trihydrate USP 22.300 mg eq. to 20mg Esomeprazole + Domperidone BP 30mg Antiulcerant (PPI) + Antiemetic	It is indicated for the management of: Functional Dyspepsia (Non-ulcer Dyspepsia) Dyspeptic symptom complex associated with gastroesophageal reflux disease (GERD): epigastric sense of fullness, feeling of abdominal distension, upper abdominal pain, nausea, vomiting, belching, flatulence, and early satiety,	Contraindications: It is contraindicated in patients with known hypersensitivity to Esomeprazole or other substituted benzimidazoles or to Domperidone or other dopamine antagonists. It should not be used whenever stimulation of gastrointestinal motility might be dangerous such as in the presence of gastrointestinal haemorrhage, mechanical obstruction, or perforation. Side Effects: Esomeprazole Common adverse events reported with Esomeprazole in clinical trials include headache, nausea, vomiting, diarrhoea, abdominal pain, flatulence, constipation and dry mouth. Domperidone: The most frequent reactions	New		K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bv gÄyi Ki‖ †h‡Z cv‡i	K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bvgÄyi Ki‖ nj	

				to Domperidone are those related to elevated prolactin levels including breast tenderness, galactorrhoea, gynaecomaslia and amenorrhoea.				
259.	Popular Pharmaceuticals Limited	Esomeprazole 40 mg + Domperidone 30 mg Capsule Esomeprazole Magnesium Trihydrate USP 44.600 mg eq. to 40mg Esomeprazole + Domperidone BP 30 mg Antiulcerant (PPI) + Antiemetic	-do-	-do-	New		*K|¤‡bkb c‡qvRb ‡bB |eavq Av‡e`b bv gÄÿ Kiv †h‡Z cv‡i |*	*K|¤‡bkb c‡qvRb ‡bB |eavq Av‡e`b bvgÄÿ Kiv nj |*
260.	Popular Pharmaceuticals Limited	Cefixime 200mg + Clavulanic Acid 125mg Tablet Cefixime BP 200mg + Diluted Clavulanate Potassium BP 297.840mg eq. to Clavulanic Acid 125mg Antibiotic	Cefixime-Clavulanic Acid should be used only to treat or prevent infections that are proven or strongly suspected to be caused by susceptible bacteria. Cefixime-Clavulanic Acid is indicated for the treatment of – 1) Uncomplicated Urinary Tract Infections. 2) Otitis Media. 3) Pharyngitis and Tonsillitis. 4) Acute Bronchitis and Acute Exacerbations of Chronic Bronchitis. 5) Uncomplicated gonorrhea etc.	Contraindications: Cefixime is contraindicated in patients with known allergy to the Cephalosporin class of antibiotics. Clavulanic Acid does not inactivate all β-Lactamases. Most chromosomally mediated β-Lactamases, the enzyme produced by pseudomonas aeruginosa, are resistant to its action. Other organism have different mechanisms of acquired resistance to β-Lactam antibiotics, against which clavulanic acid is ineffective. Side effects: Cefixime-Clavulanic Acid are diarrhea and stool changes. Events like nausea/vomiting, transient elevation in liver transaminases, alkaline phosphatase and jaundice can also occur. Thrombocytosis, thrombocytopenia, leucopenis, hypereosinophilia, neutropenia and agranulocytosis may also occur. Other adverse events that may occur are abdominal pain, abdominal cramps, flatulence, indigestion, headache, vaginitis, vulvar itch, rash, hives, itch, dysuria, chills, chest pain, shortness of breath, mouth ulcers, swollen tongue, sleepiness, thirst, anorexia.	New		*K|¤‡bkb c‡qvRb ‡bB |eavq Av‡e`b bv gÄÿ Kiv †h‡Z cv‡i |*	*K|¤‡bkb c‡qvRb ‡bB |eavq Av‡e`b bvgÄÿ Kiv nj |*
261.	Healthcare Pharmaceutical ltd., Rajendrapur, Gazipur	Ceftriaxone1.0gm + Tazobactam 125 mg/vial IV & IM Injection	Indicated for the treatment of the following infections when caused by susceptible organisms:	Contraindication: Contraindicated in patients with known allergy to the cephalosporin or beta lactam	New		*K|¤‡bkb c‡qvRb ‡bB |eavq Av‡e`b bv gÄÿ Kiv †h‡Z*	*K|¤‡bkb c‡qvRb ‡bB |eavq Av‡e`b bvgÄÿ Kiv nj |*

		Ceftriaxone Sodium and Tazobactum Sodium sterile Powder Ph. Grade 1388.88mg containning Ceftriaxone USP 1.0gm + Tazobactam USP 125 mg/vial Antibiotic	1.Lower Respiratory Tract Infections, 2.Acute Bacterial Otitis Media, 3.Skin and Skin Structure Infections, 4.Urinary Tract Infections, 5.Uncomplicated Gonorrhea, 6.Pelvic Inflammatory Disease, 7.Bacterial Septicemia, 8.Bone and Joint Infections, 9.Intra-Abdominal Infections, 10.Surgical Prophylaxis	class of antibiotics. Side Effects: Most common adverse reactions are pain, induration, tenderness, pruritus, eosinophilia, thrombocytosis, leukopenia, diarrhea & headache or dizziness was reported occasionally.			*cv‡i |*	
262.	Healthcare Pharmaceutical ltd., Rajendrapur, Gazipur	Ceftriaxone2.0gm + Tazobactam 250mg/vial IV & IM Injection Ceftriaxone Sodium and Tazobactum Sodium sterile Powder Ph. Grade 2777.76mg containning Ceftriaxone USP 2.0gm + Tazobactam USP 250 mg/vial Antibiotic	-do-	-do-	New		*K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i |*	*K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bvgÄyi Kiv nj |*
263.	ACI Ltd., Narayanganj	Ticarcillin disodium 3000 mg + Clavulanate potassium 100mg/Vial IV Infusion Ticarcillin disodium USP 3000 mg + Clavulanate potassium USP 100mg/Vial Antibiotic	To reduce the development of drug-resistant bacteria and maintain the effectiveness of Ticarcillin and Clavulanate and other antibacterial drugs, Ticarcillin and Clavulanate should be used only to treat infections that are proven or strongly suspected to be caused by bacteria. Ticarcillin and Clavulanate is a combination of a ß-lactam antibacterial and a ß-lactamase inhibitor indicated for the treatment of the following infections due to designated, susceptible bacteria: Septicemia, Lower respiratory infections, Bone and joint infections, Skin and skin structure infections, Urinary tract infections, Gynecologic	Contraindications: History of a serious hypersensitivity reaction (anaphylaxis or Stevens-Johnson syndrome) to Ticarcillin and Clavulanate or to other ß-lactams (e.g., penicillins and cephalosporins). Side Effects: The most common side effects are rash, nausea, diarrhea, and phlebitis at injection site.	Ticarcillin 3000mg + Clavulanic Acid 200mg/Vial Injection	USFDA	*K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i |*	*K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bvgÄyi Kiv nj |*

			infections, Intra-abdominal infections.					
264.	Aristopharma Ltd.	Dexamethasone Phosphate 0.10gm + Moxifloxacin 0.5gm/100gm Sterile Ophthalmic Solution Dexamethazone Sodium Phosphate USP 0.1093 gm eq.to Dexamethasone Phosphate 0.10gm + Moxifloxacin Hydrochloride BP 0.5454 gm eq.to Moxifloxacin 0.5gm/100g Antibiotic & Steroid	It is indicated for the treatment of eye infections caused by susceptible microorganisms and preventing of inflammation and bacterial infection can occur after ocular surgery	Contraindication: Epithelial herpes simplex keratitis (Dendritic keratitis), vaccinia, varicella, and many other diseases viral cornea and conjunctiva. Eye infections by mycobacteria. Ocular fungal diseases. Hypersensitivity to the components of the formula or other Quinolone derivatives. Glaucoma and /or diseases with thinning of the cornea and sclera. Side-effect: Adverse reactions that may occur with use of corticosteroids are: glaucoma with injury optic nerve, defects in visual acuity and visual field, cataract formation, infections Secondary ocular after suppression of host response and perforation of the globe. Ocular adverse events reported more frequently with the use of the ophthalmic solution 0.5% Moxifloxacin were conjunctivitis, decrease in visual perception, dry eye, inflammation of cornea, ocular discomfort, eye redness, eye pain, eye itching, bleeding subconjunctival and tearing. These events occurred in approximately 1 to 6% of patients. In surgical procedures, some of these events are consequences of own ocular surgery.	Moxifloxacin 0.5% Eye Drops, Dexamethasone Phosphate 0.1% Eye Drops		K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bv gÄy K iv †h‡Z cv‡i	
265.	Acme Laboratories Ltd.	Cefixime 1.0 gm + Clavulanic Acid 0.625 gm/50 ml Powder for Suspension (PFS) Cefixime Trihydrate BP 1.120 gm (Eqv. to Cefixime 1.0gm) + Diluted Potassium Clavulanate [As Potassium Clavulanate and Silicon Dioxide (Syloid) 1:1] BP 1.490 gm (Eqv. to Clavulanic Acid 0.625 gm)/50 ml Antibiotic (Cephalosporin + Beta Lactam Inhibitor)	Cefixime and Clavulanic Acid should be used only to treat or prevent infections that are proven or strongly suspected to be caused by susceptible bacteria. Cefixime and Clavulanic Acid is indicated for the treatment of: Uncomplicated Urinary Tract Infections caused by *Escherichia coli* and *Proteus mirabilis*. Otitis Media caused by *Haemophilus influenzae* (beta-lactamase positive and negative strains), *Moraxella (Branhamella) catarrhalis*, (most of which are beta-	Contraindication: It is contraindicated in patients with known allergy to Cefixime and Clavulanic acid or to the cephalosporin group of antibiotics. Side effects: The most frequent side effects seen with Cefixime and Clavulanic Acid are diarrhoea and stool changes. Events like nausea/vomiting, transient elevation in liver transaminases, alkaline phosphatase and jaundice can also occur. Thrombocytosis, thrombocytopenia, leucopenia, hypereosinophilia, neutropenia and agranulocytosis may also occur. Other adverse events that may occur are	New		K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bv gÄy K iv †h‡Z cv‡i	K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bvgÄy K iv nj

			lactamase positive) and *S. pyogenes*. Pharyngitis and Tonsillitis, caused by *S. pyogenes*. Acute Bronchitis and Acute Exacerbations of Chronic Bronchitis, caused by *Streptococcus pneumoniae* and *Haemophilus* influenzae (beta-lactamase positive and negative strains). Uncomplicated gonorrhea (cervical/ urethral), caused by *Neisseria gonorrhoeae* (penicillinase and non-penicillinase producing strains).	abdominal pain, abdominal cramps, flatulence, indigestion, headache, vaginitis, vulvar itch, rash, hives, itch, dysuria, chills, chest pain, shortness of breath, mouth ulcers, swollen tongue, sleepiness, thirst, anorexia.				
266.	Acme Laboratories Ltd.	Cefixime 100 mg + Clavulanic Acid 62.50mg Tablet Cefixime Trihydrate BP 112 mg (Eqv. to Cefixime 100 mg) + Diluted Potassium Clavulanate [As Potassium Clavulanate and Microcrystalline Cellulose (Avicel) 1:1] BP 149 mg (Eqv. to Clavulanic Acid 62.50 mg) Antibiotic (Cephalosporin + Beta Lactam Inhibitor)	-do-	-do-	New		K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bv gÄÿ K i v †h‡Z cv‡i |	K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bvgÄÿ K i v n j |
267.	Acme Laboratories Ltd.	Cefixime 200 mg + Clavulanic Acid 125 mg Tablet Cefixime Trihydrate BP 224mg (Eqv. to Cefixime 200 mg) + Diluted Potassium Clavulanate [As Potassium Clavulanate and Microcrystalline Cellulose (Avicel) 1:1] BP 298 mg (Eqv. to Clavulanic Acid 125 mg) Antibiotic (Cephalosporin + Beta Lactam Inhibitor)	-do-	-do-	New		K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bv gÄÿ K i v †h‡Z cv‡i |	K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bvgÄÿ K i v n j |
268.	Square Pharmaceuticals Ltd., Pabna Unit, Salgaria, Pabna	Moxifloxacin 5.00mg + Dexamethasone 1.0mg/ml Opthalmic Solution	Treatment of ocular infections caused by susceptible microorganisms and prevention of inflammation and	Hypersensitivity to other quinolones and any components, Patients with glaucoma and/or ocular disease causing thinning of the cornea	New		K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bv gÄÿ K i v †h‡Z	K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bvgÄÿ K i v n j |

		Moxifloxacin BP 5.450gm eq. to Moxifloxacin 5.00mg + Dexamethasone Sodium Phosphate BP 1.093mg eq. to Dexamethasone Phosphate 1.00mg/ml **Antibiotic + Anti-inflammatory**	bacterial infection that may occur after ocular surgery	or sclera, Inflammation in the eye caused by virus, fungi or mycobacteria, Epithelial herpes simplex keratitis (dendritic keratitis), vaccinia, varicella and many other viral diseases of the cornea, and conjunctiva. Mycobacterial infection of the eye, Fungal diseases of ocular structures.			cv‡i |	
269.	Incepta Pharmaceuticals LTd (Dhamrai Unit)	Clindamycin Phosphate 1.2gm + Benzoyl Peroxide 3.75gm/100gm Gel Clindamycin Phosphate BP 1.2gm eq. to Clindamycin 1.0gm + Benzoyl Peroxide BP 3.75gm/100gm **Antibiotic + Antiacne Antibacterial**	It is a combination of clindamycin phosphate (a lincosamide antibacterial) and benzoyl peroxide indicated for the topical treatment of acne vulgaris in patients 12 years of age and older.	Contraindications: Patients who have demonstrated hypersensitivity (e.g., anaphylaxis) to clindamycin, benzoyl peroxide, any components of the formulation, or lincomycin. Patients with a history of regional enteritis, ulcerative colitis, or antibiotic-associated colitis. Side effect: The most common adverse reactions are: burning sensation (0.4%); contact dermatitis (0.4%); pruritus (0.4%); and rash (0.4%).	New	USFDA	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |
270.	Opsonin Pharma Limited, Bagura Road, Barisal	Moxifloxacin Hydrochloride BP 0.5% + Bromfenac Sodium Sesquihydrate INN 0.09% Eye drops. Moxifloxacin Hydrochloride BP 0.5 gm + Bromfenac Sodium Sesquihydrate INN 0.09 gm/100ml Anti-infective & Anti-inflammatory (Non-steroid) Combination	Post-operative ocular Infection & Inflammation in case of steroid sensitive patients.	Contraindications: Hypersensitivity to any component of this product, **Side effects:** The most frequently ocular adverse reported with the use of moxifloxacin were conjunctivitis, decreased visual acuity, dry eye, ocular discomfort, ocular hyperemia, ocular pain and ocular pruritus, subconjunctival hemorrhage, and tearing. The most commonly reported adverse reactions following use of Bromfenac after cataract surgery include: abnormal sensation in eye, conjunctival hyperemia, eye irritation (including burning/stinging), eye pain, eye pruritus, eye redness, headache and iritis.	Moxifloxacin 0.5% Eye Drops Bromfenac 0.7% & 0.09% Eye Drops		K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i |	K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bvgÄyi Kiv nj |
271.	Popular Pharmaceuticals Limited	Azithromycin 250mg + Ambroxol Hydrochloride BP 75mg Extended Release Bi-Layer Film Coated Tablet Azithromycin USP 262mg eq. to Azithromycin 250mg + Ambroxol	It is indicated for the treatment of patients with mild to moderate infections caused by susceptible strains of the designated microorganisms in the specific conditions listed below. Upper Respiratory Tract Infections	**Contraindications:** Azithromycin Plus is contraindicated in patients with known hypersensitivity to ambroxol, azithromycin, erythromycin, or any macrolide antibiotic. **Side effects:** In clinical trials, most of the reported side effects with azithromycin	New		K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i |	K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bvgÄyi Kiv nj |

		Hydrochloride BP 75mg **Antibiotic + Expectorant**	Pharyngitis/Tonsillitis. Acute Bacterial Sinusitis Otitis Media Lower Respiratory Tract Infections Acute Bacterial Exacerbations of Chronic Obstructive Pulmonary Disease Community-Acquired Pneumonia [CAP]	were mild to moderate in severity and were related to the gastrointestinal tract, e.g., nausea, vomiting, diarrhea, or abdominal pain. Rarely but potentially serious side effects were angioedema and cholestatic jaundice. Ambroxol: The side effects on account of ambroxol include gastrointestinal side effects, skin rashes, headache, dizziness and sweating				
272.	Opsonin Pharma Limited, Bagura Road, Barisal	Moxifloxacin Hydrochloride BP 0.5% + Dexamethasone Sodium Phosphate BP 0.1% Eye drops. Moxifloxacin Hydrochloride BP 0.5gm + Dexamethasone Sodium Phosphate BP 0.1gm/100ml Antibiotic + Steroid combination	Coexisting ocular infection & inflammation or to prevent post-operative ocular infection & inflammation.	Contraindications: Hypersensitivity to other quinolones and any components of combination. Patients with glaucoma, infection in the eye caused by virus, fungi or mycobacteria. Side effects: The adverse reactions that may occur with use of corticosteroids are: Glaucoma with optic nerve damage, visual acuity defects and visual field defects; cataract formation. The most frequently ocular adverse reported with the use of moxifloxacin ophthalmic solution 0.5% were conjunctivitis, decreased visual acuity, dry eye, ocular discomfort, ocular hyperemia, ocular pain and ocular pruritus, subconjunctival hemorrhage, and tearing.	New		*K||¤‡bkb c‡qvRb ‡bB ||ea||q Av‡e`b bv gÄyi Kiv †h‡Z cv‡i |*	*K||¤‡bkb c‡qvRb ‡bB ||ea||q Av‡e`b bvgÄyi Kiv nj |*
273.	Popular Pharmaceuticals Limited	Azithromycin 500mg + Ambroxol Hydrochloride 75mg Extended Release Bi-Layer Film Coated Tablet Azithromycin USP 524mg eq. to Azithromycin 500mg + Ambroxol Hydrochloride BP 75mg **Antibiotic + Expectorant**	It is indicated for the treatment of patients with mild to moderate infections caused by susceptible strains of the designated microorganisms in the specific conditions listed below. Upper Respiratory Tract Infections Pharyngitis/Tonsillitis Acute Bacterial Sinusitis Otitis Media Lower Respiratory Tract Infections Acute Bacterial Exacerbations of Chronic Obstructive Pulmonary Disease Community-Acquired Pneumonia [CAP]	Contraindications: Azithromycin Plus is contraindicated in patients with known hypersensitivity to ambroxol, azithromycin, erythromycin, or any macrolide antibiotic. Side effects: In clinical trials, most of the reported side effects with azithromycin were mild to moderate in severity and were related to the gastrointestinal tract, e.g., nausea, vomiting, diarrhea, or abdominal pain. Rarely but potentially serious side effects were angioedema and cholestatic jaundice. Ambroxol: The side effects on account of ambroxol include gastrointestinal side effects, skin rashes, headache, dizziness and sweating	New		*K||¤‡bkb c‡qvRb ‡bB ||ea||q Av‡e`b bv gÄyi Kiv †h‡Z cv‡i |*	*K||¤‡bkb c‡qvRb ‡bB ||ea||q Av‡e`b bvgÄyi Kiv nj |*
274.	Ziska Pharmaceuticals Ltd.	Codeine Phosphate Anhydrous 54.3mg+ Chlorpheniramine Maleate 8 mg Tablet	It is a combination of codeine, an opiate agonist antitussive, and chlorpheniramine, a histamine-1 (H1) receptor antagonist indicated	Contraindications: Postoperative pain management of children undergoing tonsillectomy and/or adenoidectomy. Patients with known	Chlorpheniramine 4 mg Tablet & 2mg/ml Syrup	USFDA	*K||¤‡bkb c‡qvRb ‡bB ||ea||q Av‡e`b bv gÄyi Kiv †h‡Z cv‡i |*	*K||¤‡bkb c‡qvRb ‡bB ||ea||q Av‡e`b bvgÄyi Kiv nj |*

		Codeine Phosphate Anhydrous USP 54.30mg eq to 40.00 mg Codeine + Chlorpheniramine Maleate USP 8.0mg eq to 5.60 mg Chlorpheniramine **Anticholinergic and sedative**	for relief of cough and symptoms associated with upper respiratory allergies or a common cold. Limitation of Use: Not indicated for pediatric patients under 18 years of age.	hypersensitivity to codeine, chlorpheniramine, or any of the product components of this tablet. Side effects: Nausea and vomiting, constipation, abdominal distension, abdominal pain, blurred vision, diplopia, visual disturbances, confusion, dizziness, depression, drowsiness, sedation, headache, euphoria, facial dyskinesia, feeling faint, light-headedness, general feeling of discomfort or illness, excitability, nervousness, agitation, restlessness, somnolence, insomnia, dyskinesia, irritability, tremor				
275.	Square Pharmaceuticals Ltd., Pabna Unit, Salgaria, Pabna	Atorvastatin 10 mg + Ezetimibe 10 mg Tablet Atorvastatin Calcium USP 10.820 mg eq. to Atorvastatin 10 mg + Ezetimibe INN 10 mg Antihyperlipidemic	It contains a cholesterol absorption inhibitor and an HMG-CoA reductase inhibitor (statin), is indicated as adjunctive therapy to diet to: ➢ Reduce elevated total-C, LDL-C, Apo B, TG, and non-HDL-C, and to increase HDL-C in patients with primary (heterozygous familial and non-familial) hyperlipidemia or mixed hyperlipidemia. (1.1) ➢ Reduce elevated total-C and LDL-C in patients with homozygous familial hypercholesterolemia (HoFH), as an adjunct to other lipid-lowering treatments. Limitations of Use: No incremental benefit of this product on cardiovascular morbidity and mortality over and above that demonstrated for atorvastatin has been established. It has not been studied in Fredrickson Type I, III, IV, and V dyslipidemias.	Contraindications: ➢ Active liver disease or unexplained persistent elevations of hepatic transaminase levels. ➢ Hypersensitivity to any component of this product. ➢ Women who are pregnant or may become pregnant. ➢ Nursing mothers. Side Effects: Common adverse reactions (incidence ≥2% and greater than placebo) are: increased ALT, increased AST, and musculoskeletal pain.	Atorvastatin 10mg Tablet, Ezetimibe 10 mg Tablet	USFDA	K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bv gÄÿ K i‡V †h‡Z cv‡i |	K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bvgÄÿ K i‡V n‡j |
276.	Square Pharmaceuticals Ltd., Pabna Unit, Salgaria, Pabna	Atorvastatin 20 mg + Ezetimibe 10 mg Tablet Atorvastatin Calcium USP 21.640 mg eq. to Atorvastatin 20 mg +	-do-	-do-	Atorvastatin 10mg Tablet, Ezetimibe 10mg Tablet	USFDA	K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bv gÄÿ K i‡V †h‡Z cv‡i |	K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bvgÄÿ K i‡V n‡j |

		Ezetimibe INN 10 mg Antihyperlipidemic						
277.	Navana Pharmaceuticals Ltd. MedRx Life Science Ltd, BSCIC, Nandanpur, B. Baria	Sumatriptan Succinate 119mg eq. to Sumatriptan 85mg + Naproxen Sodium 500mg Tablet Sumatriptan Succinate BP 119mg Sumatriptan 85mg + Naproxen Sodium BP 500mg Antimigraine	Indicated for the acute treatment of migraine attacks with or without aura in adults.	Contraindications: Cardiac, Cerebrovascular, or Peripheral Vascular Disease, Uncontrolled Hypertension, Monoamine Oxidase-A Inhibitors, Ergotamine-Containing or Ergot-Type Medications, Other 5-HT1 Agonists, Hemiplegic or Basilar Migraine, Hepatic Impairment. Side Effects: Dizziness, drowsiness, Somnolence, Paresthesia, Nausea, Dyspepsia, dry mouth, chest pain or pressure, tight feeling in neck or jaw, pain spreading to arm or shoulder, sudden numbness or weakness, confusion, problems with vision, speech, or balance, bloody or tarry stools.	Naproxen Sodium 500mg Tablet	USFDA	*K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bv gÄy¨ K i v †h‡Z cv‡i |*	*K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bvgÄy¨ K i v n j |*
278.	Incepta Pharmaceuticals LTd (Dhamrai Unit)	Astaxanthin 6.0 mg + Blueberry extract 160mg+ Lutein 6mg Soft Gelatin Capsule. Astaxanthin Oleoresin 5% INN 120mg eq. to 6.0 mg Astaxanthin + Blueberry extract INN 160mg+ Lutein INN 6mg Antioxidant	• Strong antioxidant • Improves cardiovascular health (Atherosclerosis, reduce cholesterol). • Improves immune function. • Improves condition of skin • Protects skin from damage caused by sun (Reduce wrinkles, pimples and other signs of aging) • Improves recovery from central nervous system injuries • Protects from Parkinson 's disease, Dementia and Alzheimer's • Protects eyes from cataracts and macular degeneration. • Reduces inflammation (Arthritis) • Reduces risk of infertility Also Astaxnthin effectively reduce oxidative damage to DNA, decrease	Contraindication: Contraindicated for those with known allergies to Astaxanthin Side effect: No severe side effects have been reported yet for astaxanthin.	New		*K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bv gÄy¨ K i v †h‡Z cv‡i |*	*K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bvgÄy¨ K i v n j |*

			the risk for many types of cancer and stabilize blood sugar.					
279.	Incepta Pharmaceuticals LTd (Dhamrai Unit)	Astaxanthin 6.0 mg + DHA containing purified fish oil 240mg + Ginkgo Biloba Extract 120mg Soft Gelatin Capsule Astaxanthin Oleoresin 5% INN 120mg eq. to 6.0 mg Astaxanthin + DHA containing purified fish oil INN 240mg + Ginkgo Biloba Extract INN 120mg Antioxidant	• Strong antioxidant • Improves cardiovascular health (Atherosclerosis, reduce cholesterol). • Improves immune function. • Improves condition of skin • Protects skin from damage caused by sun (Reduce wrinkles, pimples and other signs of aging) • Improves recovery from central nervous system injuries • Protects from Parkinson 's disease, Dementia and Alzheimer's • Protects eyes from cataracts and macular degeneration. • Reduces inflammation (Arthritis) • Reduces risk of infertility Also Astaxnthin effectively reduce oxidative damage to DNA, decrease the risk for many types of cancer and stabilize blood sugar.	Contraindication: Contraindicated for those with known allergies to Astaxanthin Side effect: No severe side effects have been reported yet for astaxanthin. Possible side effects of ginkgo biloba include: • Nausea • Diarrhea • Dizziness • Headaches • Stomach ache • Restlessness • Vomiting.	New		*K¡¤‡bkb c‡qvRb ‡bB ¯ea¨vq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i |*	*K¡¤‡bkb c‡qvRb ‡bB ¯ea¨vq Av‡e`b bvgÄyi Kiv nj |*
280.	Incepta Pharmaceuticals LTd (Dhamrai Unit)	Astaxanthin 12.0 mg + Tocotrienol 20.25mg + Zinc 9mg Soft Gelatin Capsule Astaxanthin Oleoresin 5% INN 240mg eq. to 12.0 mg Astaxanthin + Tocotrienol INN 20.25mg + Zinc INN 9mg Antioxidant	• Strong antioxidant • Improves cardiovascular health (Atherosclerosis, reduce cholesterol). • Improves immune function. • Improves condition of skin • Protects skin from damage caused by sun (Reduce wrinkles, pimples and other signs of aging) • Improves recovery from central nervous system injuries • Protects from Parkinson 's disease, Dementia and Alzheimer's	Contraindication: Contraindicated for those with known allergies to Astaxanthin Side effect: No severe side effects have been reported yet for astaxanthin. They are sometimes prefixed by alpha-, beta-, gamma- or delta-tocotrienol. Tocotrienols emulate some of the antioxidant and disease-prevention properties of vitamin E and have some of the possible side effects of using vitamin E as well.	New		*K¡¤‡bkb c‡qvRb ‡bB ¯ea¨vq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i |*	*K¡¤‡bkb c‡qvRb ‡bB ¯ea¨vq Av‡e`b bvgÄyi Kiv nj |*

			• Protects eyes from cataracts and macular degeneration. • Reduces inflammation (Arthritis) • Reduces risk of infertility Also Astaxnthin effectively reduce oxidative damage to DNA, decrease the risk for many types of cancer and stabilize blood sugar.					
281.	Incepta Pharmaceuticals LTd (Dhamrai Unit)	Astaxanthin 4 mg + Natural Vitamin E 15mg + Ascorbic Acid (Vitamin C) 66mg Soft Gelatin Capsule Astaxanthin Oleoresin 5% INN 80mg eq. to 4 mg Astaxanthin + Natural Vitamin E INN 15mg + Ascorbic Acid (Vitamin C) BP 66.00 mg Antioxidant	Antioxidants to improve health and vitality.	Contraindication: Contraindicated for those with known allergies to Astaxanthin or any other component of the product. Side effect: No clear data found	New		*K‖¤‡bkb c‡qvRb ‡bB ‖eavq Avi‡e`b bv gÄyi Kiv †h‡Z cv‡i |*	*K‖¤‡bkb c‡qvRb ‡bB ‖eavq Avi‡e`b bvgÄyi Kiv nj |*
282.	Square Pharmaceuticals Ltd., Pabna Unit, Salgaria, Pabna	Olanzapine 3mg + Fluoxetine 25mg Capsule Olanzapine BP 3mg + Fluoxetine HCl BP 27.95 mg eq. to 25mg Fluoxetine Antipsychotic + Selective serotonin reuptake inhibitors	It combine olanzapine, an atypical antipsychotic and fluoxetine, a selective serotonin reuptake inhibitor, indicated for acute treatment of depressive Episodes Associated with Bipolar I Disorder and treatment Resistant Depression	Contraindication: MAOI: Because of the risk of serotonin syndrome, do not use MAOIs intended to treat psychiatric disorders with Ii or within 5 weeks of stopping treatment with it. Do not use it within 14 days of stopping an MAOI intended to treat psychiatric disorders. In addition, do not start it in a patient who is being treated with linezolid or intravenous methylene blue. Pimozide: Do not use. Risk of QT interval prolongation Thioridazine: Do not use. Risk of QT interval prolongation. Do not use thioridazine within 5weeks of discontinuing it. Adverse Reactions: Most common adverse reactions (≥5% and at least twice that for placebo) in adults: sedation, weight increased, appetite increased, dry mouth, fatigue, edema, tremor, disturbance in attention, blurred vision. Children and adolescents: sedation, weight increased, appetite increased, tremor, triglyceride increased, hepatic enzymes increased.	Olanzapine 5mg & , 10mg Tablet Fluoxetine 20mg Capsule	USFDA	*K‖¤‡bkb c‡qvRb ‡bB ‖eavq Avi‡e`b bv gÄyi Kiv †h‡Z cv‡i |*	*K‖¤‡bkb c‡qvRb ‡bB ‖eavq Avi‡e`b bvgÄyi Kiv nj |*

283.	Square Pharmaceuticals Ltd., Pabna Unit, Salgaria, Pabna	Olanzapine 6mg + Fluoxetine 25mg Capsule Olanzapine BP 6mg + Fluoxetine HCl BP 27.95mg eq. to 25mg Fluoxetine Antipsychotic + Selective serotonin reuptake inhibitors	It combine olanzapine, an atypical antipsychotic and fluoxetine, a selective serotonin reuptake inhibitor, indicated for acute treatment of depressive Episodes Associated with Bipolar I Disorder and treatment Resistant Depression	Contraindication: MAOI: Because of the risk of serotonin syndrome, do not use MAOIs intended to treat psychiatric disorders with Ii or within 5 weeks of stopping treatment with it. Do not use it within 14 days of stopping an MAOI intended to treat psychiatric disorders. In addition, do not start it in a patient who is being treated with linezolid or intravenous methylene blue. Pimozide: Do not use. Risk of QT interval prolongation Thioridazine: Do not use. Risk of QT interval prolongation. Do not use thioridazine within 5weeks of discontinuing it. Adverse Reactions: Most common adverse reactions (≥5% and at least twice that for placebo) in adults: sedation, weight increased, appetite increased, dry mouth, fatigue, edema, tremor, disturbance in attention, blurred vision. Children and adolescents: sedation, weight increased, appetite increased, tremor, triglyceride increased, hepatic enzymes increased.	Olanzapine 5mg & , 10mg Tablet Fluoxetine 20mg Capsule	USFDA	K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bv gÄÿ K i v †h‡Z cv‡i |	K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bvgÄÿ K i v n j |
284.	Eskayef Bangladesh Limited	Etoricoxib 1.0 gm + Methyl Salicylate 10gm/100gm Emul Gel Etoricoxib INN 1.0 gm + Methyl Salicylate BP 10gm/100gm Antirheumatics + Analgesic	It is used in the treatment, control, prevention and improvement of the following diseases, conditions and symptoms: Mild to moderate pain during dental operations Pain and swelling of joints, muscles due to rheumatoid arthritis, gout, low back pain.	Contraindications: Hypersensitivity Side effects: Rash, burning sensation	New		K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bv gÄÿ K i v †h‡Z cv‡i |	K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bvgÄÿ K i v n j |
285.	Eskayef Bangladesh Limited.	Silver Sulfadiazine 1% w/w + Chlorhexidine Gluconate 0.2% w/w Cream Silver Sulfadiazine USP 1.0 gm + Chlorhexidine Gluconate (20% Solution) BP 1.0gm eq. to Chlorhexidine Gluconate 0.2gm/100ml Antiseptic	For the treatment of leg ulcers, burns, skin grafts, incisions and other clean lesions, abrasions, minor cuts and wounds.	Contraindications: Pregnancy and Children: to increase the possibility of kernicterus. Side effects: Skin rash.	New		K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bv gÄÿ K i v †h‡Z cv‡i |	K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bvgÄÿ K i v n j |

286.	Incepta Pharmaceuticals Ltd.	Elvitegravir 150 mg , Cobicistat 150 mg, Emtricitabine 200 mg and Tenofovir 10mg Tablet Elvitegravir INN 150mg + Cobicistat INN 150mg + Emtricitabine INN 200mg + Tenofovir Alafenamide fumarate INN 11.2mg eq. to Tenofovir Alafenamide 10mg Antiviral	This product is a four-drug combination of elvitegravir, an HIV-1 integrase strand transfer inhibitor (INSTI), cobicistat, a CYP3A inhibitor, and emtricitabine and tenofovir alafenamide (TAF), both HIV1 nucleoside analog reverse transcriptase inhibitors (NRTIs) and is indicated as a complete regimen for the treatment of HIV-1 infection in adults and pediatric patients 12 years of age and older who have no antiretroviral treatment history or to replace the current antiretroviral regimen in those who are virologically-suppressed (HIV-1 RNA less than 50 copies per mL) on a stable antiretroviral regimen for at least 6 months with no history of treatment failure and no known substitutions associated with resistance to the individual components of this product.	Contraindication: Coadministration of Elvitegravir, Cobicistat, Emtricitabine and Tenofovir is contraindicated with drugs that: Are highly dependent on CYP3A for clearance and for which elevated plasma concentrations are associated with serious adverse events. Strongly induce CYP3A, which may lead to lower exposure of one or more components and loss of efficacy of Elvitegravir, Cobicistat, Emtricitabine and Tenofovir and possible resistance. Side effect: Most common adverse reaction (incidence greater than or equal to 10%, all grades) is nausea.	New	USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
287.	Incepta Pharmaceuticals Ltd. Beacon Pharmaceutical Ltd.,	Grazoprevir 100 mg + Elbasvir 50 mg Tablet Grazoprevir INN 100 mg + Elbasvir INN 50 mg Antiviral	Grazoprevir/Elbasvir is a fixed-dose combination product containing elbasvir, a hepatitis C virus (HCV) NS5A inhibitor, and grazoprevir, an HCV NS3/4A protease inhibitor, and is indicated with or without r bavirin for treatment of chronic HCV genotypes 1 or 4 infection in adults.	Contraindication: • Patients with moderate or severe hepatic impairment (Child-Pugh B or C). • OATP1B1/3 inhibitors, strong CYP3A inducers, and efavirenz. • If Grazoprevir/Elbasvir is administered with ribavirin, the contraindications to rbavirin also apply. Side effect: In subjects receiving Grazoprevir/Elbasvir for 12 weeks, the most commonly reported adverse reactions of all intensity (greater than or equal to 5% in placebo-controlled trials) were fatigue, headache, and nausea. In subjects receiving Grazoprevir/Elbasvir with ribavirin for 16 weeks, the most commonly reported adverse reactions of moderate or severe intensity (greater than or equal to 5%) were anemia and headache	New	USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

288.	Incepta Pharmaceuticals Ltd.	Sofosbuvir 400mg + Daclatasvir 60 mg Tablet Sofosbuvir INN 400 mg + Daclatasvir Dihydrpchloride INN 66mg eq. to 60 mg Daclatasvir Antiviral	It is indicated for the treatment of chronic hepatitis(CHC)infection	Contraindication: When Sofosbuvir is used in combination with ribavirin or peg-interferon alfa/ribavirin, the contraindications applicable to those agents are applicable to combination therapies. Refer to the prescribing information of peg-interferon alfa and ribavirin for a list of their contraindications. Sofosbuvir combination treatment with ribavirin or peg-interferon alfa/ribavirin is contraindicated in women who are pregnant or may become pregnant and men whose female partners are pregnant because of the risks for birth defects and fetal death associated with ribavirin The most common side effects reported with Daclatasvir in combination with sofosbuvir with or without ribavirin are fatigue (tiredness), nausea (feeling sick) and headache. For the full list of all side effects reported with Daclatasvir, see the package leaflet. Daclatasvir must not be used together with certain medicines that may reduce the effects of Daclatasvir. For more information on the medicines that should not be taken with Daclatasvir, see the package leaflet. Side effect: Adverse Reactions from Clinical Trials Experience Sofosbuvir should be administered with ribavirin or peg interferon alfa/ribavirin. Refer to the prescribing information of peg interferon alfa and ribavirin for a description of adverse reactions associated with their use. the clinical trials of a drug cannot be directly compared to rates in the clinical trials of another drug and may not reflect the rates observed in practice. The safety assessment of Sofosbuvir is based on pooled Phase 3 clinical trial data (both controlled and uncontrolled) including 650 subjects who received Sofosbuvir+ ribavirin (RBV) combination therapy for 12 weeks, 98 subjects who received	Sofosbuvir 400mg Tablet Daclatasvir 60 mg Tablet		*K¡¤‡bkb c‡q¡Rb ‡bB ¡ea¡q A¡‡e`b b¡ gÄy¨ K i¡ †h‡Z c¡‡i |*	*K¡¤‡bkb c‡q¡Rb ‡bB ¡ea¡q A¡‡e`b b¡gÄy¨ K i¡ n j |*

				Sofosbuvir+ ribavirin combination therapy for 16 weeks, 250 subjects who received Sofosbuvir+ ribavirin combination therapy for 24 weeks, 327 subjects who received SOFOSBUVIR+ peg interferon (Peg-IFN) alfa + ribavirin combination therapy for 12 weeks, 243 subjects who received peg interferon alfa + ribavirin for 24 weeks and 71 subjects who received placebo (PBO) for 12 weeks. The proportion of subjects who permanently discontinued treatment due to adverse events was 4% for subjects receiving placebo, 1% for subjects receiving SOFOSBUVIR+ ribavirin for 12 weeks, <1% for subjects receiving SOFOSBUVIR+ ribavirin for 24 weeks, 11% for subjects receiving peg interferon alfa + ribavirin for 24 weeks and 2% for subjects receiving SOFOSBUVIR+ peg interferon alfa + ribavirin for 12 weeks. Treatment-emergent adverse events observed in ≥15% of subjects in clinical trials are provided in Table 3. A side-by-side tabulation is to simplify presentation; direct comparison across trials should not be made due to differing trial designs. The most common adverse events (≥ 20%) for Sofosbuvir+ ribavirin combination therapy were fatigue and headache. The most common adverse events (≥ 20%) for Sofosbuvir + peginterferon alfa + ribavirin combination therapy were fatigue, headache, nausea, insomnia and anemia Daclatasvir is generally well tolerated with no known specific side-effects of its own. The most common side-effects seen in people taking daclatasvir with sofosbuvir are fatigue, nausea and headache. Taking daclatasvir with interferon and ribavirin can lead to additional side-effects including fever, muscle and joint aches, itching, depression, anemia (low haemoglobin level) and neutropenia (low white blood cell count). Ribavirin can also				

				cause birth defects, so it should not be used by pregnant women or their male partners				
289.	Incepta Pharmaceuticals Ltd. Healthcare Pharmaceutical ltd., Gazipur	Sofosbuvir 400mg + Velpatasvir 100mg Tablet Sofosbuvir INN 400mg + Velpatasvir INN 100mg Antiviral	Sofosbuvir + Velpatasvir Combination is indicated for the treatment of chronic hepatitis (CHC) all six genotypes infection.	Contraindication: When Sofosbuvir is used in combination with ribavirin or peg-interferon alfa/ribavirin, the contraindications applicable to those agents are applicable to combination therapies. Refer to the prescribing information of peg-interferon alfa and ribavirin for a list of their contraindications. Sofosbuvir combination treatment with ribavirin or peg-interferon alfa/ribavirin is contraindicated in women who are pregnant or may become pregnant and men whose female partners are pregnant because of the risks for birth defects and fetal death associated with ribavirin Side effect: The most common adverse events observed with Sofosbuvir in combination with velpatasvir were fatigue and headache. The most common adverse events observed with Sofosbuvir in combination with Peginterferon alfa and Ribavirin were fatigue, headache, nausea, insomnia and anemia.	Sofosbuvir 400mg Tablet		K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bv gÄyi K i‡V †h‡Z cv‡i	K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bvgÄyi K i‡V nj	
290.	Incepta Pharmaceuticals Ltd.	Tenofovir Alafenamide 10 mg Tablet Tenofovir Alafenamide Fumarate INN 11.2mg eq. to Tenofovir Alafenamide 10 mg Antiviral	Chronic Hepatitis B Human Immunodeficiency Virus (HIV)	Contraindication: No clear data found Side effect: In the Phase II study of adarunavir/cobicistat/emtricitabine/tenofovir alafenamide fixed-dose combination tablet discussed under the previous question (GS-US-299-0102), most side effects were mild to moderate in severity. Some side effects that were reported by participants receiving the darunavir/cobicistat/emtricitabine/tenofovir alafenamide tablet included diarrhea, upper respiratory tract infection, fatigue, nausea, and rash.	Tenofovir Disoproxil Fumarate 300 mg Tablet		K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bv gÄyi K i‡V †h‡Z cv‡i	K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bvgÄyi K i‡V nj	
291.	Aristopharma Ltd.	Tiotropium 2.5mcg + Olodaterol 2.5mcg/Puff Metered Dose Inhaler	It is a combination of tiotropium, an anticholinergic and olodaterol, a long-acting beta2-adrenergic	Contra-indication: All LABAs are contraindicated in patients with asthma without use of a long-term asthma control	New	USFDA	Abyjgv`b K i‡V †h‡Z cv‡i	Abyjgv`b K i‡V nj	

		Tiotropium Bromide Monohydrate BP 3.124 mcg eq.to Tiotropium 2.5 mcg + Olodaterol HCl INN 2.736mcg eq.to Olodaterol 2.5 mcg Bronchlodilator	agonist (LABA) indicated for: The long-term, once-daily maintenance treatment of airflow obstruction in patients with chronic obstructive pulmonary disease (COPD) Important limitations: It is NOT indicated to treat acute deterioration of COPD. It is NOT indicated to treat asthma.	medication. It is not indicated for the treatment of asthma. Hypersensitivity to tiotropium, ipratropium, olodaterol, or any component of this product. Side-effect: The most common adverse reactions (>3% incidence and more than an active control) were nasopharyngitis, cough, and back pain.				
292.	Acme Laboratories Ltd.	Indacaterol 110 mcg + Glycopyrronium 50 mcg Dry Powder for Inhalation Capsule Indacaterol Maleate (Superfine) INN 143 mcg Eq. to Indacaterol 110 mcg + Glycopyrronium Bromide (Superfine) BP 64 mcg Eq. to Glycopyrronium 50 mcg Bronchodilator	Indicated as a maintenance bronchodilator treatment to relieve symptoms in adult patients with chronic obstructive pulmonary disease (COPD).	Contraindication: Contraindicated in patients with known hypersensitivity to Indacaterol, Glycopyrronium or to any of the excipients of the product. Side effects: Side effects may include feeling of pressure or pain in the cheeks and forehead (possible symptoms of sinusitis), runny or stuffy nose, sneezing, dizziness, headache, cough, sore throat/or mouth, upset stomach, indigestion, cavities, pain in muscles, bones or joints, pain in extremities (e.g. arms or legs), fever , chest pain, problem falling asleep, tingling or numbness, nose bleeds, dry mouth, skin itching/rash, muscle spasm, tiredness, shakiness or trembling, nervousness, nausea, vomiting, diarrhea and abdominal pain (possible symptoms of gastroenteritis), high blood pressure. If any of these affects severely, consult with physician.	[Glycopyrronium Bromide INN 143 mcg eq. to Glycopyrronium 110 mcg (85mcg/actuation) + Indacaterol Maleate INN 64 mcg eq. to Indacaterol 50mcg (43mcg/actuation)]		*Abjgv` b Kiv †h‡Z cv‡i |*	*Abjgv` b Kiv nj |*
293.	Incepta Pharmaceuticals LTd (Dhamrai Unit)	Conjugated Estrogen 0.45mg + Bazedoxifene 20mg tablet Conjugated Estrogen BP 0.4500mg + Bazedoxifene INN 20mg Conjugated Estrogens + Antiestrogen	Conjugated estrogen & Bazedoxifene is a combination of conjugated estrogens with an estrogen agonist/antagonist indicated for treatment of the following conditions in women with a uterus: • Treatment of moderate to severe vasomotor symptoms associated with menopause • Prevention of postmenopausal osteoporosis *<u>Limitation of Use</u>:*	Contraindication: • Undiagnosed abnormal uterine bleeding Known, suspected, or past history of breast cancer • Known or suspected estrogen-dependent neoplasia • Active or past history of venous thromboembolism • Active or past history of arterial thromboembolism • Hypersensitivity (angioedema, anaphylaxis) to estrogens, bazedoxifene, or any ingredients Known hepatic impairment or disease	New	USFDA	*Abjgv` b Kiv †h‡Z cv‡i |*	*Abjgv` b Kiv nj |*

			Conjugated estrogen & bazedoxifene should be used for the shortest duration consistent with treatment goals and risks for the individual woman.	• Known protein C, protein S, or antithrombin deficiency or other known thrombophilic disorders Pregnancy, women who may become pregnant, and nursing mothers Side effect: In four prospective, randomized, placebo-controlled trials the common adverse reactions (incidence ≥5%) were muscle spasms, nausea, diarrhea, dyspepsia, abdominal pain upper, oropharyngeal pain, dizziness, and neck pain.				
294.	Globe Pharmaceuticals Ltd., Noakhali	Betamethasone 0.05% + Salicylic Acid 3% Ointment Betamethasone Dipropionate BP 0.064gm eq. to Betamethasone 0.05gm + Salicylic Acid BP 3.0gm/100gm Corticosteroid + Keratolytic	This combination is indicated for the treatment of hyperkeratotic and dry corticosteroid-responsive dermatoses where the cornified epithelium may resist penetration of the steroid. The salicylic acid constituent of this preparation, as a result of its descaling action, allows access of the dermis more rapidly than by applying steroid alone.	Contraindications: Rosacea, acne, perioral dermatitis, perianal and genital pruritus. Hypersensitivity to any of the ingredients of the Diprosalic presentations contra-indicates their use as does tuberculous and most viral lesions of the skin, particularly herpes simplex, vacinia, varicella. Diprosalic should not be used in napkin eruptions, fungal or bacterial skin infections without suitable concomitant anti-infective therapy. Side effects: Side effects that have been reported with the application of topical corticosteroids include: burning, itching, irritation, dryness, folliculities, hypertrichosis, hypopigmentation, perioral dermatitis and allergic contact dermatitis.	Betamethasone 0.05gm + Salicylic Acid 2.0gm/100ml Lotion	MHRA BNF 70 Page: 1025	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
295.	Square Pharmaceuticals Ltd., (Dhaka Unit) Kaliakoir, Gazipur	Mometasone Furoate 100mcg + Formoterol Fumarate Dihydrate 5mcg/ puff HFA Inhaler Mometasone Furoate BP 100mcg + Formoterol Fumarate Dihydrate BP 5mcg/puff Corticosteroid-Adrenorector agonist	This combination product containing a corticosteroid and a long acting beta2-adrenergic agonist indicated for the treatment of asthma in Patients of 12 years of age and older.	Contraindications: Status Asthmaticus It is contraindicated in the primary treatment of status asthmaticus or other acute episodes of asthma where intensive measures are required. Hypersensitivity: It is contraindicated in patients with known hypersensitivity to mometasone furoate, formoterol fumarate, or any of the ingredients in it. Side-effects: The most common side effects of this product include: inflammation of the nose and throat (nasopharyngitis), inflammation of the sinuses (sinusitis), Headache. Other side effects: Worsening	New	USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

				asthma or sudden asthma attacks have been reported with the use of inhaled mometasone Furoate.				
296.	Square Pharmaceuticals Ltd., Dhaka Unit, Kaliakoir, Gazipur	Mometasone Furoate 200 mcg + Formoterol Fumarate Dihydrate 5mcg/puff HFA Inhaler Mometasone Furoate BP 200mcg + Formoterol Fumarate Dihydrate BP 5mcg/puff Corticosteroid + Adrenorector agonist	-do-	-do-	New	USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
297.	Beximco Pharmaceuticals Ltd., Tongi ,Gazipur	Protease 1600 Units+ Lipase 40000 Units+ Amilase 25000 Units Capsule Pancreatin USP 400.00mg (Protease Activity USP 1600 Units+Lipase Activity USP 40,000 Units+Amilase Activity USP 25000 Units) **Pancreatic Enzyme**	The product is a used to treat people who cannot digest food normally because their pancreas does not make enough enzymes due to cystic fibrosis, swelling of the pancreas that lasts a long time (chronic pancreatitis), removal of some or the entire pancreas (pancreatectomy), or other conditions.	Contraindications: None. Side effects: Allergic reactions including skin rashes, swollen lips, painful joints, Gas, Headache etc.	Pancreatin 325 mg Tablet	BNF 70 Page-83	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
298.	Globe Pharmaceuticals Ltd., Noakhali	Dextromethorphan Hydrobromide 5mg + Doxylamine succinate 2mg + Acetaminophen 108mg/5ml Syrup Dextromethorphan Hydrobromide USP 5mg + Doxylamine succinate USP 2 mg + Acetaminophen BP/USP 108 mg/5ml Expectorant	This product is indicated for cough, sore throat, cold flu (fever, headache), sinus congestion, nasal congestion, runny nose, sneezing and rhinitis.	Contraindications: It is contraindicated if anyone has any allergy to one or any ingredients from this formulation or anyone have high blood pressure, severe heart blood vessel disease, rapid heartbeat, or severe heart problems. It's also contraindicated for asthmatic patients or those who are unable to urinate. Side effects: The common side effects are weakness, constipation, diarrhea, dizziness, drowsiness, excitability, headache, nausea, nervousness or anxiety.	New		Kv¤‡bkb c‡qvRb ‡bB wea\q Av‡e`b bv gÄyi Kiv †h‡Z cv‡i	Kv¤‡bkb c‡qvRb ‡bB wea\q Av‡e`b bvgÄyi Kiv nj
299.	Globe Pharmaceuticals Ltd.	Diphenhydramine HCl 14.0mg + L-menthol 2.0mg/5ml Syrup Diphenhydramine HCl USP 14.0mg + L-menthol USP 2.0 mg/5ml	It is indicated for the relief of cough and associated congestive symptoms (runny nose and sneezing), in the treatment of hay fever and other allergic conditions affecting the upper respiratory tract.	Contraindications: It is contraindicated in individuals with known hypersensitivity to the product or any of its constituents and also contraindicated in patients with chronic or persistent cough, such as occurs with asthma, or where cough is accompanied by excessive secretions,	New	MHRA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

		Expectorant		unless directed by the physician. It should not be administered to patients currently receiving monoamine oxidase inhibitors (MAOI) or those patients who have received treatment with MAOIs within the last two weeks Side effects: Side effects associated with the use of this syrup are uncommon. Diphenhydramine may cause drowsiness, dizziness, gastrointestinal disturbance, dry mouth, nose and throat; difficulty in urination or blurred vision.				
300.	Incepta Pharmaceuticals Ltd.	Human Fibrinogen 39.5mg + Human thrombin 349.5IU/Vial Powder for Topical Use Human Fibrinogen & Thrombin INN 0.50gm containing Human Fibrinogen 39.5 mg + Human thrombin 349.5IU/Vial Fibrin Sealant	It is a fibrin sealant indicated as an adjunct to hemostasis for mild to moderate bleeding in adults undergoing surgery when control of bleeding by standard surgical techniques (such as suture, ligature, and cautery) is ineffective or impractical. It is also used in conjunction with an absorbable gelatin sponge (USP) and is applied directly or using the Human fibrinogen and Human Thrombin Spray device.	Contraindication: Do not use: • Intravascularly. • For the treatment of severe or brisk arterial bleeding. • In patients known to have anaphylactic or severe systemic reactions to human blood products. Side effect: The most commonly reported adverse reactions (> 5 % subjects) were procedural pain, nausea, constipation, pyrexia, and hypotension.	New		K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bv gÄy K i v †h‡Z cv‡i |	K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bvgÄy K i v n j |
301.	Incepta Pharmaceuticals Ltd.	Human Fibrinogen 79.0 mg + Human thrombin 699IU/Vial Powder for Topical Use Human Fibrinogen & Thrombin INN 1.0 gm containing Human Fibrinogen 79.0 mg + Human thrombin 699IU/Vial Fibrin Sealant	It is a fibrin sealant indicated as an adjunct to hemostasis for mild to moderate bleeding in adults undergoing surgery when control of bleeding by standard surgical techniques (such as suture, ligature, and cautery) is ineffective or impractical. it is used in conjunction with an absorbable gelatin sponge (USP) and is applied directly or using the Human fibrinogen and Human Thrombin	Contraindication: Do not use: • Intravascularly. • For the treatment of severe or brisk arterial bleeding. • In patients known to have anaphylactic or severe systemic reactions to human blood products. Side effect: The most commonly reported adverse reactions (> 5 % subjects) were procedural pain, nausea, constipation, pyrexia, and hypotension.	New		K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bv gÄy K i v †h‡Z cv‡i |	K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bvgÄy K i v n j |
302.	Incepta Pharmaceuticals Ltd.	Human Fibrinogen 158.0 mg + Human Thrombin 1398 IU/Vial Powder for Topical Use Human Fibrinogen & Thrombin INN 2.0 gm containing Human Fibrinogen 158.0 mg + Human thrombin 1398IU/Vial	Human fibrinogen and Human Thrombin is a fibrin sealant indicated as an adjunct to hemostasis for mild to moderate bleeding in adults undergoing surgery when control of bleeding by standard surgical techniques (such as suture, ligature, and cautery) is	Contraindication: Do not use: • Intravascularly. • For the treatment of severe or brisk arterial bleeding. • In patients known to have anaphylactic or severe systemic reactions to human blood products.	New		K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bv gÄy K i v †h‡Z cv‡i |	K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bvgÄy K i v n j |

		Fibrin Sealant	ineffective or impractical. Human fibrinogen and Human Thrombin is used in conjunction with an absorbable gelatin sponge (USP) and is applied directly or using the Human fibrinogen and Human Thrombin Spray device.	Side effect: The most commonly reported adverse reactions (> 5 % subjects) were procedural pain, nausea, constipation, pyrexia, and hypotension.				
303.	Opsonin Pharma Limited, Bagura Road, Barisal	Levosulpiride 75 mg + Rabeprazole Sodium 20 mg Capsule Levosulpiride INN 75 mg + Rabeprazole Sodium INN 20 mg Gastro-prokinetics + Anti-ulcerant	Heartburn, GERD, IBS and gastritis, Dyspepsia etc.	Contraindication: Known hypersensitivity to Rabeprazole substituted benzimidazoles or any component of the formulation. Levosulpride-Phaeochromocytoma, epilepsy, manic states, hyperprolactinaemia, cardiac impairment etc. Side-effects: Dry mouth, nausea and vomiting, constipation, flatulence, abdominal pain, diarrhea, hypersensitivity reactions etc. Levosulpride: Genitourinary - Absence of menstrual period, breast enlargement in male, spontaneous milk secretion, and changes in libido. Potentially Fatal - Neuroleptic malignant syndrome.	New		K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i	K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bvgÄyi Kiv nj
304.	Renata Limited Mirpur, Dhaka	Drospirenone BP 0.5mg + Estradiol BP 1.0mg Tablet Drospirenone BP 0.5mg + Estradiol BP 1.0 mg Hormone	It is indicated in women with an intact uterus for the treatment of: ➢ Vasomotor symptoms due to menopause. ➢ Vulvar and vaginal atrophy symptoms due to menopause.	Contra-indication : Undiagnosed abnormal genital bleeding, Known, suspected, or history of cancer of the breast. Known or suspected estrogen-dependent neoplasia. Active DVT, PE, or a history of these conditions. Active arterial thromboembolic disease (for example, stroke and MI), or history of these conditions Renal impairment), Known liver impairment or disease, Adrenal insufficiency, Known or suspected pregnancy. Known protein C, protein S, or antithrombin deficiency, or other known thrombophilic disorders (4) Known anaphylactic reaction, angioedema, or hypersensitivity to Angeliq Side effects: The most common adverse reactions that occurred in at least 1 percent of users in clinical trials with Angeliq are gastrointestinal and abdominal pain, female genital tract bleeding, breast pain and discomfort, and headache.	Drospirenone BP 3mg + Ethinylestradiol BP 0.03 mg Tablet Drospirenone 3 mg + Ethinylestradiol 0.02mg Tablet	USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

305.	Renata Limited Mirpur, Dhaka	Drospirenone BP 2.0mg + Estradiol BP 1.0mg Tablet Drospirenone BP 2.0mg + Estradiol BP 1.0mg Hormone	It is used after menopause in women who have a uterus to reduce moderate to severe hot flashes.	Contra-indication : Progestogen/estrogens should not be used in individuals with any of the following conditions: 1. Undiagnosed abnormal genital bleeding. 2. Known, suspected, or history of cancer of the breast. 3. Known or suspected estrogen-dependent neoplasia. 4. Active deep vein thrombosis, pulmonary embolism or history of these conditions. 5. Active or recent (e.g., within the past year) arterial thromboembolic disease (e.g., stroke,myocardial infarction). 6. Renal insufficiency. 7. Liver dysfunction or disease. 8. Adrenal insufficiency. 9. Should not be used in patients with known hypersensitivity to its ingredients. 10. Known or suspected pregnancy. There is no indication in pregnancy. There appears to be little or no increased risk of birth defects in children born to women who have used estrogens and progestin's from oral contraceptives inadvertently during early pregnancy. Side effects: Undesirable effects which have been observed are: Between 1 and 10 in every 100 women are likely to experience the following: abdominal pain or bloating, or pain in your fingers or toes, feeling sick (nausea) or feeling unusually tired or weak, headache, mood swings, hot flashes, nervousness, enlarged or lumpy breasts,	Drospirenone 3mg + Ethinylestradiol 0.03mg Tablet Drospirenone 3 mg + Ethinylestradiol 0.02 mg Tablet	BNF-70 Page-652	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
306.	UniMed & UniHealth Mfg. Ltd., Gazipur	Salicylic Acid 16.70% + Lactic Acid 16.70% Solution Salicylic Acid BP 16.70gm + Lactic Acid BP 16.70gm/100ml Keratolytic	It is indicated the removal of warts, verrucas, corns and calluses.	Contra-indications: Significant peripheral neuropathy, patients with diabetes at risk of neuropathic ulcers; impaired peripheral circulation;protect surrounding skin and avoid broken skin ;not suitable for application to face, anogenital region, or large areas. Side-effects: Skin irritation, skin ulceration (with high concentrations)	New	BNF-70 Page-1058	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
307.	Globe Pharmaceuticals	Phytosterol esters 630mg +	This combinatiobn is indicated for	Contraindications: This drug is	New		K‖¤‡bkb c‡qvRb ‡bB ‖ea‖q	K‖¤‡bkb c‡qvRb ‡bB ‖ea‖q

	Ltd., Noakhali	Docosahexanoic Acid (DHA) 232.5mg + Eicosapentaenoic acid (EPA) 92.5 mg Soft Gelatin Capsule Phytosterol esters INN 630 mg + Docosahexanoic Acid (DHA) INN 232.5mg + Eicosapentaenoic acid (EPA) INN 92.5 mg Lipid Lowering Agent	the dietary management of hypertriglyceridemia.	contraindicated in patients with known hypersensitivity (e.g. anaphylactic reaction) to any of its components. It is not indicated for individuals with a condition known as phytosterolemia, also referred to as sitosterolemia. Side effects: It is well tolerated and no serious adverse events have been reported. The common adverse events reported are itching rash, fatigue and abdominal discomfort.			*Av‡e`b bvgÄÿ Kiv †h‡Z cv‡i |*	*Av‡e`b bvgÄÿ Kiv nj |*
308.	Incepta Pharmaceuticals Ltd.	Indacaterol 27.5 mcg + Glycopyrronium 15.6 mcg capsule for oral inhalation Indacaterol Maleate INN 35.6mcg eq. to Indacaterol 27.5 mcg + Glycopyrronium Bromide BP 19.5mcg eq. to Glycopyrronium 15.6 mcg Long-acting beta-2 Adrenergic Agonist + Nticholinergic	Indacaterol and Glycopyrronium is a combination of indacaterol, a long-acting beta2adrenergic agonist (LABA), and glycopyrrolate, an nticholinergic, indicated for the long term, maintenance treatment of airflow obstruction in patients with chronic obstructive pulmonary disease (COPD) Limitations of Use: Not indicated for the relief of acute bronchospasm or for the treatment of asthma.	Contraindication: All LABAs are contraindicated in patients with asthma without use of a long-term asthma controller medication. Indacaterol And Glycopyrronium is not indicated for the treatment of asthma. •History of known hypersensitivity to indacaterol, glycopyrrolate, or to any of the ingredients. Side effect: Most common adverse reactions (incidence ≥2% and higher than placebo) are nasopharyngitis and hypertension.	Indacaterol 50 mcg + Glycopyrronium 110mcg Hard Capsule for Inhalation	USFDA	*Abygv`b Kiv †h‡Z cv‡i |*	*Abygv`b Kiv nj |*
309.	Navana Pharmaceuticals Ltd.	Multivitamin & Multimineral Tablet for eye Vitamin A USP (As Vitamin A Acetate) 3500IU + Vitamin C USP (As Ascorbic Acid) 90 mg + Vitamin D USP (As *Cholecalciferol)* 400 IU + Vitamin E USP (As Vitamin E Acetate) 60 IU+ Vitamin K USP (As Dry Vitamin K_1) 25 mcg + Thiamin USP (As Thiamin	The most complete eye multivitamin and contains 10 mg of lutien and 2 mg of zeaxanthin to help support our eye health. Designed to provide key nutrients, helps support glare reduction, retinal function and healthy vision. With 25 other age-adjusted nutrients, this multivitamin also helps support our whole body health.	Contraindication: Long term intake of high level of vitamin A may increase the risk of osteoporosis in adults. Do not take this product if taking other vitamin A supplements. Side effect: Side effects has not found in PDR (Non prescription Drugs) 34 edition,2013	New		*Kwc¤‡bkb c‡qvRb ‡bB wea|q Av‡e`b bvgÄÿ Kiv †h‡Z cv‡i |*	*Kwc¤‡bkb c‡qvRb ‡bB wea|q Av‡e`b bvgÄÿ Kiv nj |*

		Mononitrate) 1.50mg + Riboflavin USP 1.70mg + Niacin USP 20mg+ Vitamin B_6 USP(As Pyridoxin hydrochloride) 3mg + Folic Acid USP 200mcg + Vitamin B_{12} USP (As Cyanocobalamin) 25mcg + Biotin BP (As D-Biotin) 30mcg + Pantothenic Acid USP (As Calcium Pantothenate) 5mg + Calcium USP (As Calcium Carbonate & Dibasic Calcium Phosphate Anhydrous) 200 mg + Phosphorous USP (As Dibasic Calcium Phosphate) 50mg + Iodine USP (As Potassium Iodide) 150mcg + Magnesium USP (As Magnesium Oxide) 100mg + Zinc USP (As Zinc Oxide) 15mg + Selenium USP (As Sodium Selenate Anhydrous) 55mcg + Copper USP (As Cupric Oxide) 0.5mg + Manganese USP (As Manganese Sulphate Monohydrate) 2.3mg + Chromium USP (As Chromic Chloride Hexahydrate) 35 mcg + Molybdenum BP (As Sodium Molybdate Dihydrate) 45 mcg + Chloride USP (As Potassium Chloride) 72 mg + Potassium USP (As Potassium Chloride) 80 mg + Lutien Ph.Gr (As Lutien) 10mg + Zeaxanthin INN 2mg Meniral + Vitamin							
310.	UniMed & UniHealth Mfg. Ltd., Gazipur	Calcium 110mg/5ml Syrup Calcium Gluconate USP 11.400gm Eqv. to 1.015gm Calcium + Calcium Lactobionate USP 24gm eq. to 1.20gm Calcium/100ml	For the Treatment calcium deficiency	Contra-indications : conditions associated with hypercalcaemia and hypercalciuria (e.g. some forms of malignant disease) Side-effects: rarely gastro-intestinal disturbances; with injection, bradycardia, arrhythmias, peripheral vasodilatation, fall in blood pressure, sweating, injectionsite reactions, severe tissue damage with	New		*K‖¤‡bkb c‡q‖Rb ‡bB ‖ea‖q Av‡e`b b‖ gÄy K i‖ †h‡Z c‖‡i |*	*K‖¤‡bkb c‡q‖Rb ‡bB ‖ea‖q Av‡e`b b‖gÄy K i‖ n j |*

		Mineral		extravasation				
311.	Incepta Pharmaceuticals Ltd.	Elemental Iron 60mg + Elemental Zinc 15mg + Folic Acid 1.0 mg + Methycobalamin 0.5mg Tablet Ferrous Bisglycinate INN eq. 300 mg to elemental Iron 60mg + Zinc Bisglycinate INN 75mg eq. to elemental Zinc 15mg + Folic Acid USP 1.0 mg + Methycobalamin USP 0.5mg Mineral + Vitamin	Treatment and prevention of iron deficiency anaemia.	Contraindication: In conditions of iron overload presentation or who are hypersensitive to any component of the product Side effect: Adverse reactions with iron therapy may include constipation, diarrhea, nausea, vomiting, dark stools and abdominal pain. Adverse reactions with iron therapy are usually transient. Allergic sensitization has been reported following both oral and parenteral administration of folic acid.	New		K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bv gÄy K iv †h‡Z cv‡i |	K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bvgÄy K iv nj |
312.	Incepta Pharmaceuticals Ltd.	Elemental Iron 1.0 gm + Elemental Zinc 0.275gm + Folic Acid 0.01gm + Cyanocobalamin 0.003gm + Lysine Hydrochloride 1.0gm/100ml Paediatric Drop. Ferrous Bisglycinate INN eq. 5gm to elemental Iron 1.0gm + Zinc Bisglycinate INN 1.75gm eq. to elemental Zinc 0.275gm + Folic Acid USP 0.01gm + Cyanocobalamin USP 0.003gm + Lysine Hydrochloride BP 1.0gm/100ml Mineral + Vitamin	Treatment and prevention of iron deficiency anaemia.	Contraindication: In conditions of iron overload presentation or who are hypersensitive to any component of the product Side effect: Adverse reactions with iron therapy may include constipation, diarrhea, nausea, vomiting, dark stools and abdominal pain. Adverse reactions with iron therapy are usually transient. Allergic sensitization has been reported following both oral and parenteral administration of folic acid.	New		K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bv gÄy K iv †h‡Z cv‡i |	K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bvgÄy K iv nj |
313.	Beacon Pharmaceutical Ltd.,	Iodine 150µgm + Folic acid 500 µgm Tablet Potassium Iodide USP 196.20µgm eq. to Iodine 150µgm + Folic acid USP 500 µgm Mineral + Vitamin	• Develop baby's brain, eye sight and hearing • Reduce the risk of birth defects of the brain and spinal cord such as spina bifida	Contraindication: Not to be taken with Folate, or other folic acid containing supplements. If suffering from a thyroid condition patient should consult with physician. Side effect: side effects include allergic reaction: hives; difficulty breathing; swelling of your face, lips, tongue, or throat, nausea; or upset stomach	New		K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bv gÄy K iv †h‡Z cv‡i |	K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bvgÄy K iv nj |
314.	MedRx Life Science Ltd, BSCIC, Nandanpur, B. Baria	Elemental Calcium 600mg + Vitamin D_3 4.12µg + Folic Acid BP 400IU Calcium carbonate BP 1500mg eq. to Calcium 600mg+	• Lack in Minerals Medications • Treatment to Prevent Mineral Deficiency Medications • Post-Menopausal Osteoporosis Prevention Medications	Contraindications: Not known Side Effects: Common: Gas, Swelling of the Abdomen, Infrequent: Feel Like Throwing Up, Incomplete or Infrequent Bowel Movements, Intense Abdominal Pain	Calcium 600 mg + Cholecalciferol (Vit. D3) 400 IU Tablet		K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bv gÄy K iv †h‡Z cv‡i |	K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bvgÄy K iv nj |

		Cholecalciferol (Vitamin D_3) BP 4.12µg + Folic Acid BP 400IU Minerals + Vitamin	• Treatment To Prevent Vitamin Deficiency Medications • Lack in Vitamins Medications	Rare: Bronchospasm, High Amount of Calcium in the Blood, Kidney Stone, Loss of Appetite, Confused, Depression, Easily Angered or Annoyed, Itching, Not Feeling Well, Over Excitement, Rash, Redness of Skin, Sleep Disorder, Taste Problems				
315.	Opsonin Pharma Limited, Bagura Road, Barisal	Vitamin A 1000 IU+ Vitamin C 15 mg + Vitamin D3 200 IU + Vitamin E 5 IU + Vitamin B6 500 mcg + Vitamin B1 500 + mcg + Vitamin B2 500 mcg + Nicotinamide 6mg + Cyanocobalamin 0.9 mcg + Pantothenic Acid 2 mg + L-Lysine 100mg/ 5ml Syrup Vitamin A BP 1000IU+ Vitamin C BP 15mg + Vitamin D3 BP 200IU + Vitamin E BP 5IU + Vitamin B6, BP 500mcg + Vitamin B1 BP 500mcg + Vitamin B2 BP 500 mcg + Nicotinamide BP 6mg + Cyanocobalamin BP 0.9mcg + Pantothenic Acid BP 2mg + L-Lysine USP 100mg/ 5 ml Multivitamin + Amino Acid	Promotes muscle growth, weight gain and calcium retention; Helps to enhance body height and weight gain; Ensures good eye sight; Necessary for the normal process in protein, fat carbohydrate metabolism; For RBC formation and correct functioning of nervous system; For proper food assimilation and cell functioning; Protect body cell from free radical	Contraindications: The products are contraindicated in patients with a known hypersensitivity to any of the ingredients of the products. Side effects: Generally well tolerated.	New		*Kw¤‡bkb c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i* |	*Kw¤‡bkb c‡qvRb ‡bB weavq Av‡e`b bvgÄyi Kiv nj* |
316.	Popular Pharmaceuticals Limited	Pregabalin 50mg + Mecobalamin 0.50mg Capsule Pregabalin INN 50mg + Mecobalamin INN 0.50mg **Neuropathic Pain Agent**	Pregabalin+Mecobalamin are prescribed for the treatment of spinal cord injury-related neuropathic pain, fibromyalgia, perioperative pain, migraine, chronic pain, peripheral neuropathy, Sciatica and other conditions. Pregabalin+Mecobalamin can improve energy, memory and learning, supports nervous and immune systems, promotes cardiovascular health, control plasma levels of homocysteine.	Contraindications It is contraindicated in patients who have demonstrated hypersensitivity to Pregabalin & Mecobalamin or to any of the excipients. Side effects: Generally well tolerated. Most common side effects are somnolence, dizziness, dry mouth and blurring of vision.	New		*Kw¤‡bkb c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i* |	*Kw¤‡bkb c‡qvRb ‡bB weavq Av‡e`b bvgÄyi Kiv nj* |
317.	Popular Pharmaceuticals Limited	Pregabalin 75mg + Mecobalamin 0.50mg Capsule Pregabalin INN 75mg +	It is indicated for the treatment of spinal cord injury-related neuropathic pain, fibromyalgia, perioperative pain, migraine, chronic	Contraindications It is contraindicated in patients who have demonstrated hypersensitivity to Pregabalin & Mecobalamin or to any of the	New		*Kw¤‡bkb c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i* |	*Kw¤‡bkb c‡qvRb ‡bB weavq Av‡e`b bvgÄyi Kiv nj* |

		Mecobalamin INN 0.50mg Neuropathic Pain Agent	pain, peripheral neuropathy, Sciatica and other conditions. It can improve energy, memory and learning, supports nervous and immune systems, promotes cardiovascular health, control plasma levels of homocysteine.	excipients. Side effects: Generally well tolerated. Most common side effects are somnolence, dizziness, dry mouth and blurring of vision.				
318.	Eskayef Bangladesh Limited	Naproxen Sodium 220mg + Dyphenhydramine HCl 25mg Tablet Naproxen Sodium BP 220 mg + Dyphenhydramine HCl BP 25mg **NSAID + Decongestant**	➢ for relief of occasional sleeplessness when associated with minor aches and pains ➢ helps you fall asleep and stay asleep	Contraindications: • Allergy alert • right before or after heart surgery • with any other product containing diphenhydramine, even one used on skin Side effects: • Stomach bleeding warning • redness or swelling is present in the painful area • have difficulty swallowing or the caplet feels stuck in throat get nervous, dizzy, or sleepless	New		*K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bvgÄy Kiv †h‡Z cv‡i |*	*K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bvgÄy Kiv nj |*
319.	Eskayef Bangladesh Limited.	Naproxen Sodium 220mg + Pseudoephedrine HCl 120mg Extended Release Tablet Naproxen Sodium USP 220mg + Pseudoephedrine Hydrochloride USP 120mg **NSAID + Decongestant**	temporarily relieves these cold, sinus, and flu symptoms: • sinus pressure • minor body aches and pains • headache, fever • nasal and sinus congestion (promotes sinus drainage and restores freer breathing through the nose)	Contraindications: Allergy alert, right before or after heart surgery, if you are now taking a prescription monoamine oxidase inhibitor (MAOI) Side effects: • Stomach bleeding warning • redness or swelling is present in the painful area • any new symptoms appear • fever gets worse or lasts more than 3 days • have difficulty swallowing or the caplet feels stuck in your throat • get nervous, dizzy, or sleepless nasal congestion lasts more than 7 days	New		*K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bvgÄy Kiv †h‡Z cv‡i |*	*K‖¤‡bkb c‡qvRb ‡bB ‖eavq Av‡e`b bvgÄy Kiv nj |*
320.	Globe Pharmaceuticals Ltd., Noakhali Ziska Pharmaceuticals Ltd.	Macrogol (3350) 6.563gm + Sodium Chloride 175.4mg + Sodium Bicarbonate 89.30mg + Potassium Chloride 25.10mg/Sachet Oral powder for solution (Pediatric formulation) Macrogol (3350) USP 6.563gm +	Chronic constipation/ prevention of Faecal impaction and Faecal impaction in children.	Contraindications: Cardiovascular impairment, renal impairment Side effects: Abdominal distension, abdominal pain, flatulence, nausea.	Macrogol 3350 13.125gm + Potassium Chloride 46.6mg + Sodium Bicarbonate 178.5mg + Sodium Chloride 350.7mg/Sachet	MHRA	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

		Sodium Chloride BP 175.40mg + Sodium Bicarbonate BP 89.30mg + Potassium Chloride BP 25.10mg/Sachet **Osmolytic laxative**						
321.	Acme Laboratories Ltd.	Saccharomycis Boulardii (Lyophilized) 2.5 Billion CFU + Lactic Acid Bacillus (Not less than 100 Million Spores) 25 mg + Elemental Zinc 10.0mg/Sachet Saccharomycis Boulardii (Lyophilized) Ph. Grade 141.25 mg (Eqv. to 2.5 Billion CFU) + Lactic Acid Bacillus (Not less than 100 Million Spores) Ph. Grade 25 mg + Zinc Sulfate Monohydrate USP 27.64 mg (Eqv. to 10 mg Elemental Zinc)/Sachet **Probiotics**	Acute infections/antibiotic induced diarrhoea, irritable bowel syndrome, diarrhoea in tube fed patients.	Contraindication: Patients with a known hypersensitivity to the content or any of its ingredients. Side effects: Probiotics are generally regarded as safe for human consumption. Few side effects have been reported. Some people experience excessive production of gas due to the corrective activity of probiotics in the colon. This is patientspecific and normally will decrease with use. Gradual increase of dosing over time is recommended to minimize this effect.	New		*K||¤‡bkb c‡qvRb ‡bB ||eavq Av‡e`b bv gÄy Kiv †h‡Z cv‡i |*	*K||¤‡bkb c‡qvRb ‡bB ||eavq Av‡e`b bvgÄy Kiv nj |*
322.	Opsonin Pharma Limited, Bagura Road, Barisal	Dapoxetine HCl INN 60mg + Tadalafil BP 20mg Tablet Dapoxetine HCl INN 60mg + Tadalafil BP 20mg **SSRI + Phosphodiesterase type 5 inhibitor**	Erectile dysfunction with premature ejaculation	Contraindicated: People with a history of cardiovascular or hepatic diseases should have a thorough consultation with the doctor before they are allowed to take Dapoxetine HCl INN 60mg + Tadalafil BP 20mg Tablet. Hypersensitivity to any of the drug's components is also a contraindication. Side-effects: Most common adverse effects include nausea, vomiting, flushes, head redness, body pain and dizziness. Patients are also disposed to the risk of heart problems and stomach ulcer. The most serious adverse effects of the drug manifest in case of misreading or ignoring precautions and instructions, and avoiding consultation with a doctor, or overdose as well. In this case, patients can face serious heart diseases, hepatotoxicity, diarrhea, hearing impairments etc.	New		*K||¤‡bkb c‡qvRb ‡bB ||eavq Av‡e`b bv gÄy Kiv †h‡Z cv‡i |*	*K||¤‡bkb c‡qvRb ‡bB ||eavq Av‡e`b bvgÄy Kiv nj |*

323.	ACI Ltd., Narayanganj	Diosmin 900 mg + Hespiridine 100 mg Tablet Diosmin 90% & Hespiridine 10% as Flavonoids glycoside INN 1040mg containing Diosmin 900 mg with 3.6% Overage + Hespiridine 100 mg with 0.4% Overage **Venotonic & Vascular Protector**	It is indicated for - •Treatment of the manifestations of chronic venous insufficiency. •Treatment of functional symptoms related to venous insufficiency of the hemorrhoidal plexus. •In the pre- and postoperative saphenous vein stripping (surgical removal of the saphenous vein) for relief of the signs and Symptoms resulting from this procedure.. •Relief of chronic pelvic pain associated with pelvic congestion syndrome	Contraindications: This combination is contraindicated in patient with known hypersensitivity to the active substances or to any of the excipients. Side Effects: The common side effects are diarrhea, dyspepsia (indigestion), nausea, vomiting, colitis, dizziness, headache, malaise, rash, pruritus and urticaria	Diosmin 450 mg + Hesperidin 50mg Tablet		*Kw¤‡bkb c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i|*	*Kw¤‡bkb c‡qvRb ‡bB weavq Av‡e`b bvgÄyi Kiv nj|*
324.	Ziska Pharmaceuticals Ltd.	Vitamin A 3500IU + Vitamin C 90 mg + Dry Vitamin D3 4 mg + Dry Vitamin E Acetate 120mg + Dry Vitamin K1 0.50mg + Thiamin Mononitrate 1.5mg+ Riboflavin 1.7mg+ Niacin 20mg+ Pyridoxine Hydrochloride (Vitamin B6) 3mg+ Folic Acid 0.20mg+ Cyanocobalamin 0.1% 25.0 mg + Biotin 0.03mg+ Calcium Pantothenate 5.435 mg + Calcium Carbonate 500mg + Phosphorus 50mg+ Potassium Iodate 150 mcg + Magnesium Oxide 100mg + Zinc 15mg +Sodium 50mcg + Copper 0.50mg + Manganese 2.3mg + Chromic 35mcg + Sodium Molybdate 96.60mcg + Chloride 72mg + Potassium 80mg + Lutein 10mg + Zeaxanthine 2mg Tablet ——— Beta-Carotene 20% USP 10.50 mg eq. to Vitamin A 3,500 IU + Vitamin C USP 90 mg + Dry Vitamin D3 USP 4.0 mg eq. to Vitamin D 400 IU + Dry Vitamin E Acetate USP 120.0 mg eq. to Vitamin E60 IU + Dry Vitamin K1 USP 0.50 mg eq. to Vitamin K_1 5 mcg+ Thiamin Mononitrate USP 1.5 mg+ Riboflavin USP 1.7 mg +	For improvement of vision	Contraindications: As with any supplement, if you are pregnant, nursing or taking medication, consult your doctor before use. Side effects: Long-term intake of high levels of vitamin A (excluding that sourced from beta-carotene) may increase the risk of osteoporosis in adults.	New		*Kw¤‡bkb c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i|*	*Kw¤‡bkb c‡qvRb ‡bB weavq Av‡e`b bvgÄyi Kiv nj|*

		Niacin USP 20mg + Pyridoxine Hydrochloride (Vitamin B6) USP 3 mg+ Folic Acid USP 0.20mg + Cyanocobalamin 0.1% USP 25.0mg eq. to Vitamin B_1 225 mcg+ Biotin USP 0.03 mg+ Calcium Pantothenate USP 5.435mg eq. to Pantothenic Acid 5mg + Calcium Carbonate BP 500.00 mg eq. to Calcium 200mg + Phosphorus USP 50mg + Potassium Iodate USP 150 mcg + Magnesium Oxide USP 184.23mg eq. to Magnesium100 mg + Zinc Oxide USP 18.6705 mg eq. to Zinc15 mg + Sodium Selenate Anhydrous Ph.Gr. 119.64mcg eq. to Selenium50 mcg + Cupric Oxide Ph.Gr. 0.626 mg eq. to Copper 0.50 mg + Manganese Sulfate USP 7.075mg eq. to Manganese 2.3mg + Chromic Chloride USP 0.107 mg eq. to Chromium 35mcg + Sodium Molybdate USP 96.60mcg eq. to Molybdenum 45mcg + Potassium Chloride USP 152.57mg eq. to Chloride 72mg + Potassium Chloride USP 152.57mg eq. to Potassium80 mg + Lutein USP 10 mg + Zeaxanthine Ph.Gr 2 mg **Vitamin**						
325.	Ziska Pharmaceuticals Ltd.	Vitamin A 2,500 IU + Vitamin C 92mg+ Dry Vitamin D3 4mg + Dry Vitamin E Acetate 70.0 mg + Dry Vitamin K1 0.60 mg + Thiamin Mononitrate 1.40mg + Riboflavin 1.40mg+ Niacin 18mg+ Pyridoxine Hydrochloride (Vitamin B6) 1.90 mg+ Folic Acid 0.80 mg+ Cyanocobalamin 0.1% 2.60mg + Biotin 0.03 mg+ Calcium Pantothenate 6.522mg +	For healthy heart.	Contraindications: As with any supplement, if you are pregnant, nursing or taking medication, consult your doctor before use. Side effects: Long-term intake of high levels of vitamin A (excluding that sourced from beta-carotene) may increase the risk of osteoporosis in adults.	New		*K‡¤‡bkb c‡qvRb ‡bB ‡ea‡q Av‡e`b bv gÄyi Kiv †h‡Z cv‡i |*	*K‡¤‡bkb c‡qvRb ‡bB ‡ea‡q Av‡e`b bvgÄyi Kiv nj |*

		Calcium Carbonate 625mg + Ferrous Sulfate 27.0mg+ Phosphorus 20mg+ Potassium Iodate 287.77mg + Magnesium Oxide 92.115mg + Zinc Oxide 13.693 mg + Sodium Selenate Anhydrous. 71.786 mcg + Cupric Oxide 1.126mg + Manganese Sulfate 6.152 mg + Chromic Chloride 0.153mg + Sodium Molybdate 107.33mcg + Chloride 72mg + Potassium 80.0mg + DHA 200 mg+ EPA 15 mg Tablet ---Beta-Carotene 20% USP 5.25 mg eq. to Vitamin A 1750 IU + Vitamin C USP 30mg+ Dry Vitamin D3 USP 2 mg eq. to Vitamin D 200IU+ Dry Vitamin E Acetate USP 30.0 mg eq. to Vitamin E 15IU+ Dry Vitamin K1 USP 0.25 mg eq. to Vitamin K 12.50mcg + Thiamin Mononitrate USP 0.75 mg+ Riboflavin USP 0.85 mg+ Niacin USP 10 mg+ Pyridoxine Hydrochloride (Vitamin B6) USP 2.5 mg+ Folic Acid USP 0.20 mg+ Cyanocobalamin 0.1% USP 100 mg eq. to Vitamin B12 100 mcg+ Biotin USP 0.015mg + Calcium Pantothenate USP 5.435 mg eq. to Pantothenic Acid 5mg+ Calcium Carbonate BP 135mg eq. to Calcium 54mg+ Ferrous Sulfate USP 3.0 mg+ Phosphorus USP 40 mg+ Potassium Iodate USP 98.105 mg eq. to Iodine 75mcg + Magnesium Oxide USP 36.846mg eq. to Magnesium 20 mg + Zinc Oxide USP 4.668mg eq. to Zinc 3.75mg + Sodium Selenate Anhydrous Ph.Gr. 0.024mg eq. to Selenium 10mcg + Cupric Oxide Ph.Gr. 0.438 mg						

		eq. to Copper 0.35mg+ Manganese Sulfate USP 3.076mg eq. to Manganese 1 mg + Chromic Chloride USP 0.183 mg eq. to Chromium 60mcg + Sodium Molybdate USP 80.50mcg eq. to Molybdenum 37.50mcg+ Potassium Chloride USP 61.452 mg eq. to Chloride 29.0mg + Potassium Chloride USP 61.028 mg eq. to Potassium 32mg + Boron Citrate USP 295.75 mcg eq. to Boron 16mcg +Nickel Sulfate USP 6.591mcg eq. to Nickel 2.5mcg + Stanous fluoride USP 5.0 mcg+ Sodium Metavendate USP 11.965 mcg eq. to Vanadium 5.0 mcg+ Phytosterols USP 400 mg **Vitamin**						
326.	Ziska Pharmaceuticals Ltd.	Vitamin A 1750 IU + Vitamin C 30mg+ Dry Vitamin D3 2 mg + Dry Vitamin E Acetate 30.0 mg + Dry Vitamin K1 0.25 mg + Thiamin Mononitrate 0.75mg+ Riboflavin 0.85mg+ Niacin 10 mg+ Pyridoxine Hydrochloride (Vitamin B6) 2.5 mg+ Folic Acid 0.20mg+ Cyanocobalamin 0.1% 100mg + Biotin 0.015 mg+ Calcium Pantothenate 5.435mg + Calcium Carbonate 135mg + Ferrous Sulfate 3.0mg+ Phosphorus 40mg+ Potassium Iodate 98.105 mg + Magnesium Oxide 36.846mg +Zinc Oxide 4.668mg +Sodium Selenate Anhydrous 0.024mg + Cupric Oxide 0.438 mg + Manganese Sulfate 3.076 mg + Chromic Chloride 0.183mg + Sodium Molybdate 80.50mcg + Chloride 29 mg + Potassium 32 mg + Boron Citrate 295.75 mcg +Nickel Sulfate 6.591mcg + Stanous	For prenatal stage	Contraindications: As with any supplement, if you are pregnant, nursing or taking medication, consult your doctor before use. Side effects: Long-term intake of high levels of vitamin A (excluding that sourced from beta-carotene) may increase the risk of osteoporosis in adults.	New		*Kv¤‡bkb c‡qvRb ‡bB wea vq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i |*	*Kv¤‡bkb c‡qvRb ‡bB wea vq Av‡e`b bvgÄyi Kiv nj |*

		fluoride 5.0mcg+ Sodium Metavendate 11.965 mcg + Phytosterols USP 400 mg Tablet Beta-Carotene 20% USP 7.50 mg eq. to Vitamin A 2,500 IU + Vitamin C USP 92mg + Dry Vitamin D3 USP 4 mg eq. to Vitamin D 400 IU + Dry Vitamin E Acetate USP 70.0 mg eq. to Vitamin E 35IU + Dry Vitamin K1 USP 0.60mg eq. to Vitamin K 30mcg + Thiamin Mononitrate USP 1.40 mg+ Riboflavin USP 1.40 mg+ Niacin USP 18 mg + Pyridoxine Hydrochloride (Vitamin B6) USP 1.90mg+ Folic Acid USP 0.80 mg+ Cyanocobalamin 0.1% USP 2.60 mg eq. to Vitamin B12 2.60 mcg + Biotin USP 0.03mg + Calcium Pantothenate USP 6.522mg eq. to Pantothenic Acid 6 mg + Calcium Carbonate BP 625 mg eq. to Calcium 250mg+ Ferrous Sulfate USP 27mg+ Phosphorus USP 20mg+ Potassium Iodate USP 287.77 mg eq. to Iodine 220mcg + Magnesium Oxide USP 92.115mg eq. to Magnesium 50mg +Zinc Oxide USP 13.693 mg eq. to Zinc 11mg + Sodium Selenate Anhydrous Ph.Gr. 71.786 mcg eq. to Selenium 30 mcg + Cupric Oxide Ph.Gr. 1.126mg eq. to Copper 0.90 mg + Manganese Sulfate USP 6.152 mg eq. to Manganese 2 mg + Chromic Chloride USP 0.153mg eq. to Chromium 30mcg + Sodium Molybdate USP 107.33 mcg eq. to Molybdenum 50 mcg+ Potassium Chloride USP 152.57 mg eq. to Chloride 72mg + Potassium Chloride USP						

		152.57mg eq. to Potassium 80mg + DHA (Docosahexaenoic Acid) Ph.Gr. 200mg + EPA (Eicosapentaenoic Acid) Ph.Gr.15 mg.						
327.	Incepta Pharmaceuticals LTd (Dhamrai Unit)	Fish oil 250 mg + Vitamin D_3 0.52 mg + Natural Vitamin E 9.85mg + Ascorbic Acid (Vitamin C) 41.30 mg + Astaxanthin 2 mg Soft Gelatin Capsule. Fish oil (rich in Omega-3Acids) BP 250 mg contaning EPA 280mg/gm & DHA 190mg/gm + Vitamin D_3 BP 0.52 mg + Natural Vitamin E INN 9.85mg + Ascorbic Acid (Vitamin C) BP 41.30 mg + Astaxanthin Oleoresin 5% INN 40mg eq. to 2 mg Astaxanthin Vitamin	It is indicated for improvement of muscle function and edurance. Daily intake of this effective formulation contributes to the maintenance of normal heart, muscle and bone health. As a food supplement combination of antioxidants and fish oil to improve health and vitality.	Contraindication: Contraindicated for those with known allergies to Astaxanthin or any other component of the product. Side effect: No clear data found	New		Kw¤‡bkb c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i|	Kw¤‡bkb c‡qvRb ‡bB weavq Av‡e`b bvgÄyi Kiv nj|
328.	Ziska Pharmaceuticals Ltd.	L-Cystine HCl 200 mg + Methionine 200 mg + Ascorbic Acid (Vitamin C) 60mg + Pantothenic Acid(Vit B5) 6mg+ Pyridoxine HCl (Vit B6) 2mg+ Riboflavin 1.6mg+ Biotin .15 mg+ Vit E 10mg+ Ferrous Sulfate 14mg+ Zinc Oxide 14mg+ Magnesium Oxide 45mg Capsule L-Cystine HCl Ph.Gr 200mg + L-Methionine Ph.Gr 200 mg + Ascorbic Acid (Vitamin C) BP 60mg + Pantothenic Acid (Vit B5) BP 6mg+ Pyridoxine HCl (Vit B6) BP 2 mg + Riboflavin BP 1.6mg+ Biotin BP 0.15mg + Vit E as Alphatocopherol Acetate BP 10mg+ Ferrous Sulfate BP 14mg+ Zinc Oxide BP 14mg+ Magnesium Oxide BP 45mgVitamin	Hair and nails fragility. Brittle nails, change in hair quality, temporary hair loss. In alopecia of various origins, and low fractional hair growth disorders and hair structure. In brittle, splitting and nail dystrophy.	Contraindications: None Side-effects: None	New		Kw¤‡bkb c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i|	Kw¤‡bkb c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv nj|
329.	Globe Pharmaceuticals Ltd., Noakhali	L-methylfolate calcium 3mg + Pyridoxal-5-Phosphate 35 mg + Methylcobalamin 2mg FC Tablet L-methylfolate calcium INN 3 mg +Pyridoxal-5-Phosphate INN 35 mg +	This combination is indicated for the treatment of endothelial dysfunction in patients with loss of protective sensation and neuropathic pain associated with diabetic peripheral neuropathy and alsocauses hyperhomocysteinemia who present with lower extremity ulceration.	Contraindications: This combination is contraindicated in patients who have hypersensitivity to any of the components of this product. Side effects: Paresthesia, somnolence, nausea, headaches, mild transient diarrhea,	New		Kw¤‡bkb c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i|	Kw¤‡bkb c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv nj|

		Methylcobalamin INN 2mg Vitamin		polycythemia vera, itching and feeling of swelling of the entire body have been reported.				
330.	Incepta Pharmaceuticals LTd (Dhamrai Unit) Cod Liver Oil Soft Gelatin Capsule	Cod Liver Oil Soft Gelatin Capsule Pure Cod Liver oil BP 320mg containing EPA 26mg + DHA 24mg + Vitamin A 670IU + Vitamin D3 67IU + Vitamin E 0.3 IU Vitamin	Treatment of combined vitamins A, D and E deficiencies. Combined vitamins A, D and E and Polyunsaturates supplement to the diet during childhood and in adults with very poor dietary status. Traditional remedy in the symptomatic relief of muscular and joint stiffness and aches.	Contraindication: - Hypersensitivity to the active substances or to any of the excipients. - Hypercalcaemia - Hypervitaminoses A, D and E Side effect: Vitamin excess can be harmful but a very large overdose of this product would be needed to produce ill effects. As most undesirable effects are based on post-marketing spontaneous reporting, precise frequency estimation is not possible. - Gastrointestinal disorders, particularly at high doses, e.g. eructation, fishy after-taste, nausea, vomiting, abdominal pain, constipation, diarrhoea - Moderate increases in hepatic transaminases have been reported in patients with hypertriglyceridaemia - Skin reactions, e.g. acne, eczema and rash Reporting of suspected adverse reactions	New		K¤‡bkb c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i |	K¤‡bkb c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv nj |
331.	Incepta Pharmaceuticals Ltd.	Bromelain 50mg eq. to 20,000 Units +Trypsin 2500 USP Units Tablet Bromelain INN 50mg eq. to 20,000 Units +Trypsin USP 1.0mg eq. to Trypsin 2500 USP Units	Natural Medicines Comprehensive Database rates effectiveness based on scientific evidence according to the following scale: Effective, Likely Effective, Possibly Effective, Possibly Ineffective, Likely Ineffective, Ineffective, and Insufficient Evidence to Rate. The effectiveness ratings for BROMELAIN are as follows: Possibly effective for Arthritis (osteoarthritis) pain and knee function when used in combination with trypsin and rutin (Combination of Bromelain, trypsin). This combination seems to be about as effective as some prescription painkillers. Possibly ineffective for.. Preventing muscle soreness (myalgia) after exercise. Taking bromelain orally, immediately following intense exercise does not seem to delay onset of muscle	Contraindication: Bromelain can raise the risk of bleeding. Make sure to stop taking it at least two weeks before surgery. Check with a doctor before using bromelain if you have any health conditions, such as a bleeding disorder, asthma, heart problems, liver or kidney disease, or stomach ulcers. Side effect: Bromelain is Possibly Safe for most people when taken in appropriate amounts. Bromelain may cause some side effects, such as diarrhea and stomach and intestinal discomfort. Bromelain may also cause allergic reactions, especially in people who have other allergies. If you have allergies, be sure to check with your healthcare provider before taking bromelain.	New		K¤‡bkb c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i |	K¤‡bkb c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv हल|

			soreness and has no effect on pain, flexibility, or skeletal weakness. Insufficient evidence to rate effectiveness for Knee pain. There's some evidence that taking bromelain by mouth can reduce mild acute knee pain that's lasted for less than three months in otherwise healthy people. Reducing swelling after surgery or injury. Some research suggests bromelain might reduce swelling and pain after surgery or injury. Interestingly, bromelain 02.doesn't seem to reduce swelling after mouth surgery. Severe burns.Inflammation. Improving antibiotic absorption. Hay fever. Preventing cancer. Shortening of labor. Ulcerative colitis. Other conditions. More evidence is needed to rate the effectiveness of bromelain for these uses.					
332.	Libra Infusions Limited	20% Fat Emulsion Infusion Refine Soyabean Oil BP 20.0gm + Egg Yolk Phospholipids Ph. Grade 1.20gm + Glycerol BP 2.25gm/100ml	It is indicated as a source of calories source OF calories and essential fatty acids for patients requiring parenteral nutrition for extended periods of time (usually for more than 5 days) and as a source of essential fatty acids for prevention of efad.	Contra-indications: The administration of 20% fat emulsion is contraindicated in patients with disturbances of normal fat metabolism such as pathologic hyperlipemia, lipoid nephrosis or acute pancreatits if accompanied by hyperlipidem Side effects: Those more frequently encountered are due: either to contamination of the intravenous catheter and result in sepsis, or to vein irritation by concurrently infused hypertonic solutions and may result in thrombophlebitis. These adverse reactions are inseparable from the hyper-alimentation procedure with or without 20% i.v. fat emulsion. 2. Less frequent reactions more directly related to 10% i.v fat emulsion) are: a) immediate or early adverse reactions, each of which has been reported to occur in clinical trials, in an incidence of less than 1 %; dyspnea, cyanosis, allergic reactions, hyperlipemia, hypercoagulability, nausea,	10% Fat Emulsion Infusion	USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

				vomiting, headache, flush-ing, increase in temperature, sweating, sleepiness, pain in the chest and back, slight pressure over the eyes, dizziness, and irritation at the site of infusion, and, rarely, thrombocytopenia in neonates; b) delayed adverse reactions such as hepatomegaly, jaundice due to central lobular cholestasis, splenomegaly, thrombocytopenia, leukopenia, transient increases in liver function tests, and overloading syndrome (focal seizures, fever, leukocytosis, hepatomegaly, splenomegaly and shock).				
333.	Libra Infusions Limited	15% Amino acid with 20% Dextrose Infusion Chamber-1: L-Isoleucine USP 0.990gm + L-Leucine USP 1.50 gm + L-Lysine Hydrochloride USP 1.575gm + L-Methionine USP 0.258gm + L-Phenyalanine USP 0.447gm + L-Threonine USP 0.60gm + L-Tryptophan USP 0.30gm + L-Valine USP 0.750gm + L-Alanine USP 1.490gm + L-Arginine USP 1.527gm + L-Aspartic Acid USP 1.050gm + L-Glutamic Acid USP 1.107gm + L-Histidine USP 0.450gm+ L-Proline USP 1.083gm + L-Serine USP 0.795gm +L-N-Acetyl-L-Tyrosine USP 0.405gm + L-Glysine USP 0.750gm/100ml; Chamber-2: Dextrose USP 20gm/100ml Mix two chambers Before using.	Amino acid solution infused with dextrose by peripheral vein infusion is indicated as a source of nitrogen in the nutritional support of patients with adequate stores of body fat, in whom, for short periods of time, oral nutrition cannot be tolerated, is undesirable, or inadequate. amino acid can be administered peripherally with dilute dextrose solution. This form of nutritional support can help to preserve protein and reduce catabolism in stress conditions where oral intake is inadequate. Amino acid is also indicated for central vein infusion to prevent or reverse negative nitrogen balance in patients where the alimentary tract, by the oral, gastrostomy or jejunostomy route cannot or should not be used and gastrointestinal absorption of protein is impaired. Products are indicated as a source of amino acids and carbohydrate calories in clinical conditions where enteral nutritional supply is or is expected to be insufficient or impossible in order to offset or prevent nitrogen loss or negative nitrogen balance.	Contra-indications: • This preparation should not be used in patients with hepatic coma or metabolic disorders involving impaired nitrogen utilization. • Known allergy to corn or corn products since the products contain corn-derived dextrose • Patients with acute renal failure and without undergoing renal replacement therapy. • Patients with severe liver failure or hepatic coma • Congenital abnormality of amino acid metabolism • Severe hyperglycemia (glucose concentration greater than 180 mg/dL or 10 mmol/L) Side effects: Hyperosmolar syndrome, resulting from excessively rapid administration of concentrated dextrose may cause mental confusion and/or loss of consciousness.Reactions which may occur because of the solution or the technique of administration include febrile response, infection at the site of injection, venous thrombosis or phlebitis extending from the site of injection, extravasation and hypervolemia. Generalized flushing, fever and nausea also have been reported during peripheral infusions of amino acid solutions.	New		*Abjgv`b Kiv †h‡Z cv‡i |*	*Abjgv`b Kiv nj |*

334.	Libra Infusions Limited	5.2% Amino acid Infusion L-Isoleucine USP 0.462gm + L-Leucine USP 0.726 gm + L-Lysine Hydrochloride USP 0.535gm + L-Methionine USP 0.726gm + L-Phenyalanine USP 0.726gm + L-Threonine USP 0.330gm + L-Tryptophan USP 0.165gm + L-Valine USP 0.528gm + L-Arginine USP 0.60gm + L-Histidine USP 0.429gm/100ml	Amino acid-RF 5.2 %(an amino acid injection — renal formula) is indicated only as an adjunct to management of patients with potentially reversible acute renal failure who are unable to eat. When infused with hypertonic dextrose as a source of calories and with added appropriate electrolytes and vitamins, Amino acid-RF 5.2% is suitable as an intravenous source of protein in a parenteral nutritional regimen for such patients.	-do-	New	USFDA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
335.	Libra Infusions Limited	15% Amino acid Infusion L-Isoleucine USP 0.990gm + L-Leucine USP 1.50 gm + L-Lysine Hydrochloride USP 1.575gm + L-Methionine USP 0.258gm + L-Phenyalanine USP 0.447gm + L-Threonine USP 0.60gm + L-Tryptophan USP 0.30gm + L-Valine USP 0.750gm + L-Alanine USP 1.490gm + L-Arginine USP 1.527gm + L-Aspartic Acid USP 1.050gm + L-Glutamic Acid USP 1.107gm + L-Histidine USP 0.450gm+ L-Proline USP 1.083gm + L-Serine USP 0.795gm +L-N-Acetyl-L-Tyrosine USP 0.405gm +L-Glysin USP 0.750gm/100ml	Products are indicated as a source of amino acids and carbohydrate calories in clinical conditions where enteral nutritional supply is or is expected to be insufficient or impossible in order to offset or prevent nitrogen loss or negative nitrogen balance.	-do-	New	USFDA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj

Annex-B

1.2 Proposed Product for Import (Human):

bs	***c¯ZKvi‡Ki bvg***	***JI‡ai bvg I †RMbwiK bvg***	***wb‡`kbv***	**Contraindication & Side-effect**	**FSC/CPP**	**Status (New Molecule/ Existing)**	***‡UKwbK¨vj mve-KwgwUi 64 Zg mfvi wm×vš***	***mfvi wm×vš***
01.	Savoy Laboratories (International) Ltd. **Local agent:** Zas Corporation, Outer circular Extention Road . Banglamotor . Dhaka	Burn Relief Spray Benzocaine/Chlorhexidine Diacetate **Anesthetics**	For treatment of minor burns and scalds, where the skin remain largely intact.	**Contraindications:** Do not use when there is extensive damage to the skin. Do not uses in the skin remain largely intact. **Side-effects:** May produce local irritation, numbness in the treated area	U.K	New	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
02.	Savoy Laboratories (International) Ltd. **Local agent:** Zas Corporation, Outer circular Extention Road . Banglamotor . Dhaka	First aid Spray Benzocaine/Chlorhexidine Diacetate/Luviskol VA64 (copoviidone Solution 50%) **Anesthetics**	Protective applications for the treatment of cuts, scratches, abrasions, minor burns and scalds.	**Contraindications:** Do not use when there is extensive damage to the skin. Do not uses in the skin remain largely intact. **Side-effects:** May produce local irritation, numbness in the treated area	U.K	New	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

03.	Manufactured by : Silver Spring Sagl, 6850 Mendrisio Switzerland **Local Agent:** Global Life Care Services& Consultation Centre, Hashim Tower, H:205/1-A, (6th Floor, Unit-F) Tejgaon-Gulshan Link Road, Tejgaon, Dhaka-1208, Bangladesh)	Goleic Injection Glycoprotein Macrophage Activating Factor (GcMAF) **Anti Cancer**	Its an anticancer and immune boosting agent that helps in destruction of cancer cells and success can be achieved with all tumor cancers including breast, lung, prostrate, pancreatic and melanoma. Treatment with Goleic Injection (GcMAF) shows improvement in 85% cases of autism and 15% total cure, as it improves neuronal metabolic activity through cAMP signaling It also help cure the diseases causing immune dysfunction.	**Contraindication:** Should not be given during pregnancy and lactation. Low-dose naltrexone, Externally administered heparin. Aspartame, All kinds of Corticosteroids (Prednisolon, etaprednisolon, Solu-Medrol etc) Anti-inflammatory drugs should be avoided. (NSAIDs like Ibuprofen, Diclofenac, Aspirin etc if necessary, should be taken in moderation.) Cytotoxic medications Cyclophosphamide, Etoposide, Methotrexate, should be taken carefully Morphine and analogues Tramadole, codeine, Fentanyl Oxycodon should be avoided. **Side effects :** Goleic Injection (GcMAF) has shown no side effects of its own in some cases give you minor side effects like fatigue and minor weight loss, headache ,occasional mild muscular pain, joint pain and the symptoms of a fever (3-5 hours of hot flushes)due to rebuild of immune system.	FSC-Federal Republic of Germany, Switzerland and Czech Republic		*c‡qvRb ‡bB wea‡q Av‡e`b bvgÄyb Kiv †h‡Z cv‡i |*	*c‡qvRb ‡bB wea‡q Av‡e`b bvgÄyb Kiv nj |*

04.	Manufactured by : Silver Spring Sagl, 6850 Mendrisio Switzerland **Local Agent:** Global Life Care Services& Consultation Centre, Hashim Tower, H:205/1-A, (6th Floor, Unit-F) Tejgaon-Gulshan Link Road, Tejgaon, Dhaka-1208, Bangladesh)	MicroBioMax Suppository Glycoprotein Macrophage Activating Factor (GcMAF) **Anti Cancer**	It's an anticancer and immune boosting agent that helps in destruction of cancer cells and success can be achieved with all tumor cancers including breast, lung, prostrate pancreatic and melanoma. Treatment with Microbiomax Suppositories (GcMAF) shows improvement in 85% cases of autism and 15% total cure, as it improves neuronal metabolic activity through cAMP signaling. It also helps cure the diseases causing immune dysfuction.	**Contraindication:** Should not be given during pregnancy and lactation. Low-dose naltrexone,Externally administered heparin.Aspartame ,All kinds of Corticosteroids (Prednisolon, Betaprednisolon, Solu-Medrol etc) Anti-inflammatory drugs should be avoided. (NSAIDs like Ibuprofen, Diclofenac, Aspirin etc if necessary, should be taken in moderation.) Cytotoxic medications Cyclophosphamide, Etoposide, Methotrexate, should be taken carefully Morphine and analogues Tramadole, codeine, Fentanyl Oxycodon should be avoided. **Side effects**: MIcrobiomax Suppositories (GcMAF) has shown no side effects of its own.In some cases give minor side effects like fatigue and minor weight loss, headache, occasional mild muscular pain, joint pain and the symptoms of a fever (3-5 hours of hot flushes)due to rebuild of immune system.	FSC-Federal Republic of Germany, Switzerland and other European countries.		*c‡qvRb ‡bB wea|q Av‡e`b bvgÄyi Kiv †h‡Z cv‡i |*	*c‡qvRb ‡bB wea|q Av‡e`b bvgÄyi Kiv nj |*

05.	Manufactured by : Silver Spring Sagl, 6850 Mendrisio Switzerland **Local Agent:** Global Life Care Services& Consultation Centre, Hashim Tower, H:205/1-A, (6th Floor, Unit-F) Tejgaon-Gulshan Link Road, Tejgaon, Dhaka-1208, Bangladesh)	MicroBioMax Immune Blend Glycoprotein Macrophage Activating Factor (GcMAF**)** **Anti Cancer**	Its' an anticancer and immune boosting agent that helps in destruction of cancer cells and success can be achieved with all tumor cancers including breast, lung, prostrate, pancreatic and melanoma. Treatment with Microbiomax Immune Blend (GcMAF) shows improvement in 85% cases of autism and 15% total cure, as it improves neuronal metabolic activity through cAMP signaling. It also help cure the diseases causing immune dysfunction.	**Contraindication:** Should not be given during pregnancy and lactation. Low-dose naltrexone, Externally administered heparin. Aspartame, All kinds of Corticosteroids (Prednisolon, Betaprednisolon, Solu-Medrol etc) Anti-inflammatory drugs should be avoided. (NSAIDs like Ibuprofen, Diclofenac, Aspirin etc if necessary, should be taken in moderation.) Cytotoxic medications Cyclophosphamide, Etoposide, Methotrexate, should be taken carefully Morphine and analogues Tramadole, codeine, Fentanyl Oxycodon should be avoided. **Side effects**: Microbiomax immune blend (GcMAF) has shown no side effects of its own. in some cases give you minor side effects like fatigue and minor weight loss, headache, occasional mild muscular pain, joint pain and the symptoms of a fever (3-5 hours of hot flushes)due to rebuild of immune system.	FSC-European Community and in Switzerland		*c‡qvRb ‡bB wea‡q Av‡e`b bvgÄyi Kiv †h‡Z cv‡i |*	*c‡qvRb ‡bB wea‡q Av‡e`b bvgÄyi Kiv nj |*

06.	**Manufacturer:**Novartis Pharma AG, Switzerland (Novartis Bangladesh Ltd)	Zykadia 150 mg Hard Capsule Ceritinib INN 150mg Hard Capsule **Anti Cancer**	ZYKADIA is indicated for the treatment of patients with locally advanced or metastatic non-small cell lung cancer (NSCLC) that is anaplastic lymphoma kinase (ALK)-positive.	**Contraindication:** Hypersensitivity to either of the active substances or to any of the excipients. **Side Effects:** Anemia, decreased appetite, hyperglycemia, hypophosphatemia, vision disorder, pericarditis, bradycardia, diarrhea, nausea, vomiting, abdominal pain, constipation, esophageal disorder, rash, fatigue, liver laboratory test abnormalities, blood creatinine increased	Switzerland	New	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
07.	Octa pharma, France **Local agent:** Discount Pharma. 30, Bir Uttam K.M. Shafiullah Road. Dhaka.	OCTAGAM 10% Solution for Infusion Liquid Intravenous Immunoglobulin-G Solution2gm/20ml Vial **Anti Cancer**	OCTAGAM 10% is indicated for the treatment of Primary Immunodeficiency Syndrome, Replacement Therapy in myelama or chronic lymphatic Leukemia, Replacement Therapy in children with AIDS, chronic immune thrombocytopenic purpura (ITP), Guillain- Barre' Syndrome (GBS), Kawasaki Disease, Allogenic Bone Marrow Transplantation.	**Contraindications:** History of an allergic reaction to any Human immunoglobulin preparation or to any constituent of Octagam 10%. Hyper sensitivity to homologous immunoglobulins, especially in very rare cases of immunoglobulin A (IgA) difeciency, when the patient has antibodies against IgA. **Side Effects:** In general, various allergic and hypersensitivity type of reactions and headache, chills, back pain, chest pain, fever, cutaneous reactions, vomiting, arthralgia, low blood pressure and nausea may occasionally occur. Reactions to IV Ig tend to be related to the dose and rate of infusion.	France	BNF-70; Nornal Immunoglobulin; Page: 1064	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj

08.	Octa pharma, France **<u>Local agent:</u>** Discount Pharma. 30, Bir Uttam K.M. Shafiullah Road. Dhaka.	OCTAGAM 10% Solution for Infusion Liquid Intravenous Immunoglobulin-G Solution 5gm/50ml Vial **Anti Cancer**	OCTAGAM 10% is indicated for the treatment of Primary Immunodeficiency Syndrome, Replacement Therapy in myelama or chronic lymphatic Leukemia , Replacement Therapy in children with AIDS, chronic immune thrombocytopenic purpura (ITP), Guillain- Barre' Syndrome (GBS), Kawasaki Disease, Allogenic Bone Marrow Transplantation.	**Contraindications:** History of an allergic reaction to any Human immunoglobulin preparation or to any constituent of Octagam 10%. Hyper sensitivity to homologous immunoglobulins, especially in very rare cases of immunoglobulin A(IgA) deficiency, when the patient has antibodies against IgA. **Side Effects:** In general, various allergic and hypersensitivity type of reactions and headache, chills, back pain, chest pain, fever, cutaneous reactions, vomiting, arthralgia, low blood pressure and nausea may occasionally occur. Reactions to IVIg tend to be related to the dose and rate of infusion.	France	BNF-70; Nornal Immunoglobulin; Page: 1064	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
09.	Octa pharma, France **<u>Local agent:</u>** Discount Pharma. 30, Bir Uttam K.M. Shafiullah Road. Dhaka.	OCTAGAM 10% Solution for Infusion Liquid Intravenous Immunoglobulin-G Solution 10gm/100ml Vial **Anti Cancer**	OCTAGAM 10% is indicated for the treatment of Primary Immunodeficiency Syndrome, Replacement Therapy in myelama or chronic lymphatic Leukemia , Replacement Therapy in children with AIDS, chronic immune thrombocytopenic purpura (ITP), Guillain- Barre' Syndrome (GBS), Kawasaki Disease, Allogenic Bone Marrow Transplantation.	**Contraindications:** History of an allergic reaction to any Human immunoglobulin preparation or to any constituent of Octagam 10%. Hyper sensitivity to homologous immunoglobulins, especially in very rare cases of immunoglobulin A(IgA) difeciency, when the patient has antibodies against IgA. **Side Effects:** In general, various allergic and hypersensitivity type of reactions and headache, chills, back pain, chest pain, fever, cutaneous reactions, vomiting, arthralgia, low blood pressure and nausea may occasionally occur. Reactions to IVIg tend to be related to the dose and rate of infusion.	France	BNF-70; Nornal Immunoglobulin; Page: 1064	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj

10.	Octa pharma, France **Local agent:** Discount Pharma. 30, Bir Uttam K.M. Shafiullah Road. Dhaka.	OCTAGAM 10% Solution for Infusion Liquid Intravenous Immunoglobulin-G Solution 20gm/200ml Vial **Anti Cancer**	OCTAGAM 10% is indicated for the treatment of Primary Immunodeficiency Syndrome, Replacement Therapy in myelama or chronic lymphatic Leukemia , Replacement Therapy in children with AIDS, chronic immune thrombocytopenic purpura (ITP), Guillain- Barre' Syndrome (GBS), Kawasaki Disease, Allogenic Bone Marrow Transplantation.	**Contraindications:** History of an allergic reaction to any Human immunoglobulin preparation or to any constituent of Octagam 10%. Hyper sensitivity to homologous immunoglobulins, especially in very rare cases of immunoglobulin A (IgA) difeciency, when the patient has antibodies against IgA. **Side Effects:** In general, various allergic and hypersensitivity type of reactions and headache, chills, back pain, chest pain, fever, cutaneous reactions, vomiting, arthralgia, low blood pressure and nausea may occasionally occur. Reactions to IV Ig tend to be related to the dose and rate of infusion.	France	BNF-70; Nornal Immunoglobulin; Page: 1064	Abjgv`b Kiv †h‡Z cv‡i |	Abjgv`b Kiv nj |
11.	Thymoorgan Pharmazie GmbH, Germany **Local agent:** Zas Corporation, Outer circular Extention Road . Banglamotor . Dhaka	IdarubicinPhaRes 1mg/ml Solution for Injection in 5mg/5ml Vial Idarubicin HCl 5mg/5ml **Anti Cancer**	To treat Leukemia	**Contraindications:** Hypersensitivity reactions to idarubicin or other drugs formulated in Cremophor EL (polyoxyethylated castor oil) , patients with solid tumors . **Side Effects:** Nausea, vomiting, abdominal cramps, diarrhea, and headache, hair loss may occur	Germany	New Molecule	Abjgv`b Kiv †h‡Z cv‡i |	Abjgv`b Kiv nj |
12.	Thymoorgan Pharmazie GmbH, Germany **Local agent:** Zas Corporation, Outer circular Extention Road . Banglamotor . Dhaka	IdarubicinPhaRes 1mg/ml Solution for Injection in 10mg/10ml Vial Idarubicin HCl 10mg/10ml **Anti Cancer**	To treat Leukemia	**Contraindications:** Hypersensitivity reactions to idarubicin, patients with solid tumors. **Side Effects:** Nausea, vomiting, abdominal cramps, diarrhea, and headache, hair loss may occur	Germany	New Molecule	Abjgv`b Kiv †h‡Z cv‡i |	Abjgv`b Kiv nj |

13.	Jenssen Cilag, SpA, Vai C Janssen 04010,Borgo San Michele Italy. Importer: Janata Traders	Invokana 300mg Film coated Tablet Canagliflozin 300mg **Anti Diabetic**	Invokana (Canagliflozin) is indicated in adults aged 18 years and older with type two diabetes mellitus to improve glycemic control as: a) Monotherapy: When died and exercise along do not provide adequate glycemic control in patients for whom the use of metformine is considered inappropriate due to intolerance or contraindication.	**Contraindication**s: Hypersentivity to the active substance or to any of the excipients. **Side effect:** Constipation, Nausea, thirst, polyurea urinary tract infection and vulbovaginal candidiasis.	Italy, EMA		c‡qvRb ‡bB wea©vq Av‡e`b bvgÄyi Kiv †h‡Z cv‡i	c‡qvRb ‡bB wea©vq Av‡e`b bvgÄyi Kiv nj
14.	Sanofi-Aventis Deutschland GmbH, Germany (Sanofi Bangladesh Ltd.)	LYXUMIA 10µg Solution for injection Lixisenatide 10µg/0.2ml Pre-Filled Pen **Anti Diabetic**	LYXUMIA is indicated for the treatment of adults with type 2 diabetes mellitus to achieve glycaemic control in combination with metformin, metformin and sulphonylurea, basal insulin and metformin, basal insulin and sulphonyl urea when these, together with diet and exercise, do not provide adequate glycaemic Control.	**Contraindications:** Lyxumia is contraindicated in patients with known hypersensitivity to lixisenatide or to any of the inactive ingredients in the formulation. **Side effects:** Very common adverse reactions are Hypoglycemia, Headache, Nausea, vomiting, Diarrhea and the common adverse reactions are Influenza, upper respiratory tract infection, cystitis, viral infection, dizziness, somnolence, dyspepsia, Back pain, Injection site pruritus, urticaria.	EMA,		Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

15.	Sanofi-Aventis Deutschland GmbH, Germany (Sanofi Bangladesh Ltd.)	LYXUMIA 20µg Solution for injection Lixisenatide 20µg/0.2ml Pre-Filled Pen **Anti Diabetic**	LYXUMIA is indicated for the treatment of adults with type 2 diabetes mellitus to achieve glycaemic control in combination with metformin, metformin and sulphonylurea, basal insulin and metformin, basal insulin and sulphonyl urea when these, together with diet and exercise, do not provide adequate glycaemic control.	**Contraindications:** Lyxumia is contraindicated in patients with known hypersensitivity to lixisenatide or to any of the inactive ingredients in the formulation. **Side effects:** Very common adverse reactions are Hypoglycemia, Headache, Nausea, vomiting, Diarrhea and the common adverse reactions are Influenza, upper respiratory tract infection, cystitis, viral infection, dizziness, somnolence, dyspepsia ,Back pain, Injection site pruritus, urticaria.	EMA		Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj

16.	Novo Nordisk A/S Novo Allé DK-2880 Bagsværd Denmark **Local agent:** Novo Nordisk Pharma (Private) Limited Nina Kabbo, Level-9, 227/A, Gulshan-Tejgaon Link Road, Tejgaon, Dhaka-1208, Bangladesh	Xultophy®; Solution for injection 1 ml solution contains 100units Insulin Degludec & 3.6mg Liraglutide **(Antidiabetic)**	Xultophy® is indicated for the treatment of adults with type-2 diabetes mellitus to improve glycaemic control in combination with oral glucose-lowering medicinal products when these alone or combined with GLP-1 receptor agonist or basal insulin do not provide adequate glycaemic control.	**Contra-indication:** Hypersensitivity to either or both active substances or to any of the excipients. **Undesirable effects** Summary of the safety profile The most frequently reported adverse reactions during treatment with Xultophy® were hypoglycaemia &gastrointestinal adverse reactions. Adverse reactions Adverse reactions associated with Xultophy® are given below by frequency. Frequency categories are defined as: Very common (≥1/10); common (≥1/100 to <1/10); uncommon (≥1/1,000 to <1/100); rare (≥1/10,000 to<1/1,000); very rare (<1/10,000) and not known (cannot be estimated from the available data). **Very common:** Hypoglycaemia. **Common:** Decreased appetite, nausea, diarrhoea, vomiting, constipation, dyspepsia, gastritis, abdominal pain, flatulence, gastroesophageal reflux disease, abdominal distension, injection site reaction. **Uncommon:** Urticaria, dehydration, rash, pruritus, increased heart rate. **Rare:** Hypersensitivity, lipodystrophy acquired. **Unknown:** Anaphylactic reaction, pancreatitis (including necrotising pancreatitis), peripheral oedema.	EMA CPP (Country of Origin, Denmark)	New	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj

17	Ferrer International, Spain. **Local agent:** Zas Corporation, Outer circular Extention Road . Banglamotor . Dhaka	Zalain Cream Sertaconazole 2% **Anti Fungal**	To treat interdigital tinea pedis in immunocompetent patients 12 years of age and older, caused by: Trichophyton rubrum, Trichophyton mentagrophytes, and Epidermophyton floccosum.	**Contraindications**: Allergic to other azole antifungals **Side Effect:** Burning, swelling, irritation, tenderness, discoloration, or dry skin may occur	UK	New	*Abyjgv` b Kiv †h‡Z cv‡i |*	*Abyjgv` b Kiv nj |*
18.	Ferrer International, Spain. **Local agent:** Zas Corporation, Outer circular Extention Road . Banglamotor . Dhaka	Eneas 10mg/20mg Enalapril maleate 10mg Nitrendipine 20mg **Antihypertensive**	Treatment of essential arterial hypertension in patients whose blood pressure is not adequately controlled on enalapril or nitrendipine alone	**Contra-indication:** Patients with history of angioedema, Second and third trimesters of pregnancy, unstable haemodynamic conditions, especially cardiovascular shock, acute heart failure, acute coronary syndrome, acute stroke. **Side Effect:** Usual: headache, facial redness, cough and swollen legs. Uncommon: dizziness, tachycardia, erythematous rash, nausea, dyspepsia and hypotension.	U.K	New	*c‡qvRb ‡bB wea‡q Av‡e`b bvgÄyi Kiv †h‡Z cv‡i |*	*c‡qvRb ‡bB wea‡q Av‡e`b bvgÄyi Kiv nj |*
19	**Manufacturer:** Novartis Farmaceutica SA, Spain. **Local Agent:** Novartis Bangladesh Ltd	Exforge HCT Film-coated tablet Amlodipine Besilate BP 6.94 mg (eqv. To 5 mg Amlodipine) + Valsartan USP 160 mg + Hydrochlorothiazide USP 12.5 mg) **Antihypertensive**	Treatment of essential hypertension as substitution therapy in adult patients whose blood pressure is adequately controlled on the combination of amlodipine, valsartan and hydrochlorothiazide taken either as three single-component formulation or as a dual-component formulations or as dual-component and a single-component formulation.	**Contraindication:** Hypersensitivity to amlodipine, valsartan, HCTZ, other sulfonamides or to any of the excipients. Exforge HCT is contraindicated in pregnancy. **Side effects:**Headache, fatigue, oedema, flushing.	EMA	Amlodipine 5 mg + Valsartan 160mg Tablet	*c‡qvRb ‡bB wea‡q Av‡e`b bvgÄyi Kiv †h‡Z cv‡i |*	*c‡qvRb ‡bB wea‡q Av‡e`b bvgÄyi Kiv nj |*

20.	**Manufacturer:** Novartis Farmaceutica SA, Spain. **Local Agent:** **Novartis Bangladesh Ltd**	Exforge HCT Film-coated tablet Amlodipine Besilate BP 13.87 mg (eqv. To 10 mg Amlodipine) + Valsartan USP 160mg + Hydrochlorothiazide USP 12.5 mg) **Anti hypertensive**	Treatment of essential hypertension as substitution therapy in adult patients whose blood pressure is adequately controlled on the combination of amlodipine, valsartan and hydrochlorothiazide taken either as three single-component formulation or as a dual-component formulations or as dual-component and a single-component formulation.	**Contraindication:** Hypersensitivity to amlodipine, valsartan, HCTZ, other sulfonamides or to any of the excipients. Exforge HCT is contraindicated in pregnancy. **Side effects:**Headache, fatigue, oedema, flushing.	EMA	Amlodipine 10mg + Valsartan 160mg Tablet	*c‡qvRb ‡bB wea‡q Av‡e`b bvgÄy Kiv †h‡Z cv‡i |*	*c‡qvRb ‡bB wea‡q Av‡e`b bvgÄy Kiv nj |*
21.	**Manufacturer:** Novartis Farmaceutica SA, Spain. **Local Agent:** **Novartis Bangladesh Ltd**	Exforge HCT Film-coated tablet Amlodipine Besilate BP 6.94 mg (eqv. To 5 mg Amlodipine) + Valsartan USP 160 mg + Hydrochlorothiazide USP 25 mg) **Antihypertensive**	-do-	-do-	EMA	Amlodipine 5 mg + Valsartan 160mg Tablet	*c‡qvRb ‡bB wea‡q Av‡e`b bvgÄy Kiv †h‡Z cv‡i |*	*c‡qvRb ‡bB wea‡q Av‡e`b bvgÄy Kiv nj |*
22.	**Manufacturer:** Novartis Farmaceutica SA, Spain. **Local Agent:** **Novartis Bangladesh Ltd**	Exforge HCT Film-coated tablet Amlodipine Besilate BP 13.87 mg (eqv. To 10 mg Amlodipine) + Valsartan USP 160 mg + Hydrochlorothiazide USP 25 mg) **Antihypertensive**	-do-	-do-	EMA	Amlodipine 10mg + Valsartan 160mg Tablet	*c‡qvRb ‡bB wea‡q Av‡e`b bvgÄy Kiv †h‡Z cv‡i |*	*c‡qvRb ‡bB wea‡q Av‡e`b bvgÄy Kiv nj |*

23.	**Manufacturer:** Novartis Singapore Pharmaceutical Manufacturing Pte. Ltd **Local Agent:** Novartis Bangladesh Ltd	Entresto 50 mg FC Tablet Sacubitril/Valsartan free anhydrous acid 50 mg Eq. to Sacubitril INN 24.3 mg and Valsartan USP 25.7 mg **Antihypertensive**	Entresto is indicated to reduce the risk of cardiovascular mortality and morbidity in adult patients with systolic heart failure (NYHA class II-IV, LVEF ≤40%). Entresto is administered in combination with other heart failure therapies (e.g. beta blockers, diuretics and mineralocorticoid antagonists) as appropriate, in place of an ACE inhibitor or ARB.	**Contraindication** • Hypersensitivity to either of the active substances, sacubitril or valsartan, or to any of the excipients • Concomitant use of an ACE inhibitor. Entresto must not be administered until 36 hours after discontinuing ACE inhibitor therapy. • Known history of angiooedema related to previous ACE inhibitor or ARB therapy. • Concomitant use of Entresto with aliskiren-containing products in patients with diabetes mellitus or patients with renal impairment (GFR <60 ml/min/1.73m2) • Severe renal impairment with an eGFR<10 ml/min/1.73 m2 owing to a lack of data. • Pregnancy **Side Effects:** Hyperkalemia, hypokalaemia, dizziness, headache,Vertigo, hypotension, diarrhea, cough, renal impairment, fatigue, asthenia & nausea.	Swiss and USA	New	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj

24.	**Manufacturer:** Novartis Singapore Pharmaceutical Manufacturing Pte. Ltd **Local Agent:** **Novartis Bangladesh Ltd**	Entresto 100 mg FC Tablet Sacubitril/Valsartan free anhydrous acid 100 mg Eq. to Sacubitril INN 48.6mg and Valsartan USP 51.4 mg **Antihypertensive**	Entresto is indicated to reduce the risk of cardiovascular mortality and morbidity in adult patients with systolic heart failure (NYHA class II-IV, LVEF ≤40%). Entresto is administered in combination with other heart failure therapies (e.g. beta blockers, diuretics and mineralocorticoid antagonists) as appropriate, in place of an ACE inhibitor or ARB.	**Contraindication** • Hypersensitivity to either of the active substances, sacubitril or valsartan, or to any of the excipients • Concomitant use of an ACE inhibitor. Entresto must not be administered until 36 hours after discontinuing ACE inhibitor therapy. • Known history of angiooedema related to previous ACE inhibitor or ARB therapy. • Concomitant use of Entresto with aliskiren-containing products in patients with diabetes mellitus or patients with renal impairment (GFR <60 ml/min/1.73m2) • Severe renal impairment with an eGFR<10 ml/min/1.73 m2 owing to a lack of data. • Pregnancy **Side Effects:** Hyperkalemia, hypokalaemia, dizziness, headache,Vertigo, hypotension, diarrhea, cough, renal impairment, fatigue, asthenia & nausea.	-do-	New	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
25.	**Manufacturer:** Novartis Singapore Pharmaceutical Manufacturing Pte. Ltd **Local Agent:** **Novartis Bangladesh Ltd**	Entresto 200 mg FC Tablet Sacubitril/Valsartan free anhydrous acid 200 mg Eq. to Sacubitril INN 97.2mg and Valsartan USP 102.8mg **Antihypertensive**	-do-	-do-	-do-	New	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj

26.	**Ferring International** Center SA Chemin De la Vergognausaz 50, 1162 Saint-Prex, Switzerland **Local agent:** Radiant Export Import Enterprise. Uttara, Dhaka	Pentasa Prolonged Release Granules 1.0gm/sachet Mesalazine Ph. Eur. Prolonged Release Granules 1.0gm/sachet **Aminosalicylates**	Treatment of mild to moderate ulcerative colitis and maintenance of remission.	**Contraindication**: Aminosalicylates should be avoided in salicylate hypersensitivity **Side effects**: Side-effects of the aminosalicylates include diarrhoea, nausea, vomiting, abdominal pain, exacerbation of symptoms of colitis, headache, hypersensitivity reactions (including rash and urticaria); side effects that occur rarely include acute pancreatitis, hepatitis, myocarditis, pericarditis, lung disorders (including eosinophilia and fibrosing alveolitis), peripheral neuropathy, blood disorders (including agranulocytosis, aplastic anaemia, leucopenia, methaemoglobinaemia, neutropenia, and thrombocytopenia—see also recommendation above), renal dysfunction (interstitial nephritis, nephrotic syndrome), myalgia, arthralgia, skin reactions (including lupus erythematosus-like syndrome, Stevens Johnson syndrome), alopecia. **Cautions:** Renal function should be monitored before starting an oral aminosalicylate, at 3 months of treatment, and then annually during treatment (more frequently in renal impairment). Blood disorders can occur with aminosalicylates. **Hepatic impairment:** avoid in severe impairment **Renal impairment:** use with caution; avoid if eGFR less than 20 mL/minute/1.73 m^2 **Adverse effects:** The most frequent adverse reactions seen in clinical trials are diarrhoea (3%), nausea (3%), abdominal pain (3%), headache (3%), vomiting (1%), and rash (1%).	Switzerland	New	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj

27.	**Ferring International** Center SA Chemin De la Vergognausaz 50, 1162 Saint-Prex, Switzerland **Local agent:** Radiant Export Import Enterprise. Uttara, Dhaka	Pentasa 2.0gm/Sachet Prolonged Release Granules Mesalazine Ph. Eur. Prolonged Release Granules 2.0 gm/Sachet **Aminosalicylates**	Treatment of mild to moderate ulcerative colitis and maintenance of remission.	-do-	Switzerland	New	*Abjgv`b Kiv †h‡Z cv‡i |*	*Abjgv`b Kiv nj |*
28.	Ferrer International, Spain. **Local agent:** Zas Corporation, Outer circular Extention Road . Banglamotor . Dhaka	Nadixa 1% Cream Nadifloxacine **Antibiotics**	Topical treatment of mild or moderate infl¬ammatory forms of acne vulgaris (papulopustular acne, grade I-II)	**Contra-indication:** Nadixa is contraindicated in cases of known hypersensitivity to Nadi¬oxacin or any of the excipients of the formulation **Side-effects**: Pruritus, burning sensation, erythema, contact dermatitis and urticaria. Skin hypopigmentation is also described in rare reports.	UK	New	*c‡qvRb ‡bB wea‡q Av‡e`b bvgÄyi Kiv †h‡Z cv‡i |*	*c‡qvRb ‡bB wea‡q Av‡e`b bvgÄyi Kiv nj |*
29.	**Ferring GmbH** Wittland 11, 24109 Kiel, Germany **Local agent:** Radiant Export Import Enterprise. Uttara, Dhaka	Minirin Nasal Spray 0.1mg/ml (1 Vial of 2.5ml) Desmopressin Ph. Eur Nasal Spray 0.1mg/ml **Antidiuretic**	Treatment of Diabetes insipidus & primary nocturnal enuresis. Vasopressin (antidiuretic hormone, ADH) is used in the treatment of pituitary ('cranial') diabetes insipidus as is its analogue desmopressin. Desmopressin is also used to boost factor VIII concentration in mild to moderate haemophilia and in von Willebrand's disease; it is also used to test fibrinolytic response.	**Contraindication**: Cardiac insufficiency and other conditions treated with diuretics; psychogenic polydipsia and polydipsia in alcohol dependence. **Renal impairment:** use with caution; antidiuretic effect may be reduced **Side effects**: Fluid retention, and hyponatraemia (in more serious cases with convulsions) on administration without restricting fluid intake; stomach pain, headache, nausea, vomiting, allergic reactions, and emotional disturbance in children also reported; epistaxis, nasal congestion, rhinitis with nasal spray.	Germany	New	*Abjgv`b Kiv †h‡Z cv‡i |*	*Abjgv`b Kiv nj |*

30.	**Ferring International** Center SA Chemin De la Vergognausaz 50, 1162 Saint-Prex, Switzerland **Local agent:** Radiant Export Import Enterprise. Uttara, Dhaka	Minirin 0.1 mg Tablets Desmopressin Acetate Ph. Eur. 0.1mg **Antidiuretic**	Treatment of Diabetes insipidus & primary nocturnal enuresis. Vasopressin (antidiuretic hormone, ADH) is used in the treatment of pituitary ('cranial') diabetes insipidus as is its analogue desmopressin. Desmopressin is also used to boost factor VIII concentration in mild to moderate haemophilia and in von Willebrand's disease; it is also used to test fibrinolytic response.	**Contraindication**: Cardiac insufficiency and other conditions treated with diuretics; psychogenic polydipsia and polydipsia in alcohol dependence. **Renal impairment:** use with caution; antidiuretic effect may be reduced **Side effects**: Fluid retention, and hyponatraemia (in more serious cases with convulsions) on administration without restricting fluid intake; stomach pain, headache, nausea, vomiting, allergic reactions, and emotional disturbance in children also reported; epistaxis, nasal congestion, rhinitis with nasal spray.	Switzerland	New	*Abjgv`b Kiv †h‡Z cv‡i |*	*Abjgv`b Kiv nj |*
31.	**Ferring International** Center SA Chemin De la Vergognausaz 50, 1162 Saint-Prex, Switzerland **Local agent:** Radiant Export Import Enterprise. Uttara, Dhaka	Minirin 0.2 mg tablet Desmopressin Acetate Ph. Eur. 0.2mg **Antidiuretic**	Treatment of Diabetes insipidus & primary nocturnal enuresis. Vasopressin (antidiuretic hormone, ADH) is used in the treatment of pituitary ('cranial') diabetes insipidus as is its analogue desmopressin. Desmopressin is also used to boost factor VIII concentration in mild to moderate haemophilia and in von Willebrand's disease; it is also used to test fibrinolytic response.	-do-	Switzerland	New	*Abjgv`b Kiv †h‡Z cv‡i |*	*Abjgv`b Kiv nj |*

32.	**Manufacturer:** **Catalent UK Swindon Zydis Ltd, UK** **License Holder:** **Ferring AG,** Switzerland **Local agent:** Radiant Export Import Enterprise. Uttara, Dhaka	Minirin Melt 60 mcg Sublingual Tablet Desmopressin Ph. Eur Oral Lyophilisate 60 mcg **Antidiuretic**	Treatment of Diabetes insipidus & primary nocturnal enuresis. Vasopressin (antidiuretic hormone, ADH) is used in the treatment of pituitary ('cranial') diabetes insipidus as is its analogue desmopressin. Desmopressin is also used to boost factor VIII concentration in mild to moderate haemophilia and in von Willebrand's disease; it is also used to test fibrinolytic response.	-do-	Switzerland	New	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
33.	**Manufacturer:** **Catalent UK Swindon Zydis Ltd, UK** **License Holder:** **Ferring AG,** Switzerland **Local agent:** Radiant Export Import Enterprise. Uttara, Dhaka	Minirin Melt 120 mcg Sublingual Tablet Desmopressin Ph. Eur Oral Lyophilisate 120 mcg **Antidiuretic**	Treatment of Diabetes insipidus & primary nocturnal enuresis. Vasopressin (antidiuretic hormone, ADH) is used in the treatment of pituitary ('cranial') diabetes insipidus as is its analogue desmopressin. Desmopressin is also used to boost factor VIII concentration in mild to moderate haemophilia and in von Willebrand's disease; it is also used to test fibrinolytic response.	-do-	Switzerland	New	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj

34.	River Pharma Srl,, Italy **Local Agent:** Benvue International, 14/1 Joy Tower, Chattesory Road, Joy nagor R/A, Chittagonj	Nevralip 600 Retard Tablet Alpha Lipoic Acid 600 mg **Antioxidant**	Alpha Lipoic Acid (ALA) is indicated for management of neuropathic pain (Diabetic polyneuropathy), Alzheimer & Parkinson Disease, Diabetic Cardiomyopathy	**Contra-indications:** Alpha Lipoic acid (ALA) is contraindicated in patients with known hypersensitivity to Alpha Lipoic acid (ALA) or any of its components. **Side effects:** Alpha Lipoic acidic well tolerated but is generally rare. Low doses of lipoic acid show no side effects. But higher doses could cause skin rash, nausea or stomach upset, along with nervousness, fatigue and insomnia.	Italy		c‡qvRb ‡bB wea‡q Av‡e`b bvgÄyK iv †h‡Z cv‡i	c‡qvRb ‡bB wea‡q Av‡e`b bvgÄyK iv nj
35.	River Pharma Srl,, Italy **Local Agent:** Benvue International, 14/1 Joy Tower, Chattesory Road, Joy nagor R/A, Chittagonj	Syalox 150 Capsule Hyaluronic Acid 150mg **Cartilaginous Defect Repair Agent**	Hyaluronic Acid (HA) is indicated for degenerative & connective tissue disorders due to impaired hyaluronic acid level in the joints.	**Contra-indications:** Hyaluronic Acid (HA) is contraindicated in patients with known hypersensitivity to Hyaluronic Acid (HA) or any of its components. **Side effects:** Hyaluronic Acid (HA) is well tolerated with few side effects like dyspepsia, nausea, abdominal pain etc.	Italy		c‡qvRb ‡bB wea‡q Av‡e`b bvgÄyK iv †h‡Z cv‡i	c‡qvRb ‡bB wea‡q Av‡e`b bvgÄyK iv nj

36.	**Ferring GmbH** Wittland 11, 24109 Kiel, Germany **Local agent:** Radiant Export Import Enterprise. Uttara, Dhaka	Firmagon 80mg/Vial Powder and solvent for solution for injection. (For Subcutaneous use Only) Degarelix (as acetate) INN 80mg /Vial **Gonadotrophin-releasing hormone antagonist**	Degarelix is a gonadotrophin-releasing hormone antagonist used to treat advanced hormone-dependent prostate cancer. It does not induce a testosterone surge or tumour 'flare', therefore anti-androgen therapy is not required.	**Contraindication**: Hypersensitivity to the active substance or to any of the excipients. It is not indicated in women or paediatric patients. In addition, due to its pharmacological effects may cause foetal harm if administered to a pregnant woman. **Side effects**: nausea; dizziness, headache, drowsiness, insomnia, asthenia; influenza-like symptoms; hot flushes, sweating (including night sweats), weight gain; injection-site reactions; less commonly diarrhoea, vomiting, abdominal discomfort, dry mouth, constipation, anorexia, atrio-ventricular block, QT-interval prolongation, fainting, hypertension, hypersensitivity reactions, depression, anxiety, oedema, gynaecomastia, micturition urgency, renal impairment, sexual dysfunction, pelvic pain, prostatitis, testicular pain, anaemia, musculoskeletal pain, tinnitus, urticaria, alopecia, and rash. **Cautions**: susceptibility to QT-interval prolongation (avoid concomitant use of drugs that prolong QT interval); monitor bone density; diabetes **Hepatic impairment:** manufacturer advises caution in severe impairment—no information available **Renal impairment:** manufacturer advises caution in severe impairment—no information available. **Adverse effects:** The most frequently reported adverse reactions at the injection sites were pain (28%), erythema (17%), swelling (6%), induration (4%) and nodule (3%). These adverse reactions were mostly transient, of mild to moderate intensity, occurred primarily with the starting dose and led to few discontinuations (<1%).	Germany	**New**	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj

37.	**Ferring GmbH** Wittland 11, 24109 Kiel, Germany **Local agent:** Radiant Export Import Enterprise. Uttara, Dhaka	Firmagon 120mg/Vial Powder and solvent for suspension for injection (For Subcutaneous use Only) Degarelix (as acetate) INN 120 mg /Vial **Gonadotrophin-releasing hormone antagonist**	Degarelix is a gonadotrophin-releasing hormone antagonist used to treat advanced hormone-dependent prostate cancer. It does not induce a testosterone surge or tumour 'flare', therefore anti-androgen therapy is not required.	-do-	Germany	**New**	*Abyjgv` b Kiv †h‡Z cv‡i |*	*Abyjgv` b Kiv nj |*

38.	Mearck Sharp & Dohme Corp. **Importer:** Janata Traders	Varivax Vaccine Vericella Virus Vaccine **Vaccine**	Vaccine is indicated for active immunization for the prevention of varicella in individuals 12 month of age and older. Varivax can be administrated to infants from 9 month of age under special circumstances, such as to conform with national vaccination schedules or in outbreak situation. Varivax may also be administration to susceptible individuals who have been exposed varicella. Vaccination within 3 days of exposure may prevent a clinically apparent infection or modify the course of the infection .In addition ,there are limited data that indicate that indicate that vaccination up to 5 days after exposure may modify the course of the infection.	**Contraindication**: • History of hypersensitivity to any vericella vaccine ,to any of the excipients or to gelatin or neomycin . • Blood dyscrasias eukaemia,lymphomas, of any type or other malignant neoplasm affecting the hemic and lymphatic systems. • Individuals receiving immunosuppressive therapy . • severe hormonal or cellular immunodeficiency, agammglobulinemia and AIDS or symtomatic HIV infection or an age specific CD4+T-lymphocyte percentage in children bellow 12 month CD4+<25%,Children between 12-35 months;CD4+<20%children between 36-59 months.CD4<15% . • Individual with a family history of congenital or hereditary immunodeficiency ,unless the immune competence of the potential vaccine recipient is demonstrated. **Side effect:** Common: Upper respiratory infection , Rash.Measles/rubella-like rash, ,Injection , Varicella- like rash,injection site erythema,pain/ ,Skin and subcutaneous tissue disorder,tenderness/Soreness/,Swelling Irritability. **Very Common :** Fever	EMA		Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj

39.	**Merck Sharp & Dohme Corp. USA** **Local agent :** Janata Traders TCB Bhabon , 1 kawran Bazar,Dhaka **Importer:** Janata Traders	**VAQTA 0.5ml** Hepatitis A Vaccine 0.5ml Inactivated **Vaccine**	**VAQTA** is indicated for active pre-exposure prophylaxis against disease caused by Hepatitis-A virus.	CONTRAINDICATION Do not administer VAQTA to individuals with a history of immediate and/or severe allergic or hyper sensitivity reactions , (anaphylaxis) after a previous dose of any hepatitis A vaccine or with an Anaphylactic reaction to neomycin. Possible side effects Like all medicines and vaccines, this vaccine can cause side effects, although not everybody gets them. As with all vaccines, allergic reactions, in rare cases leading to shock, may Occur. These reactions may include: Hives, Difficulty in breathing , Swelling of the face, tongue and throat, Dizziness, collapse.	USA		Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
40.	**Merck Sharp & Dohme Corp. USA** **Local agent :** Janata Traders TCB Bhabon , 1 kawran Bazar,Dhaka **Importer:** Janata Traders	**VAQTA 1ml** Hepatitis A Vaccine 1ml Inactivated. **Vaccine**	VAQTA is indicated for active pre-exposure prophylaxis against disease caused by Hepatitis-A virus.	-do-	USA		Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj

41.	**Ferring Leciva a.s.** 252 42 Jesenice u Prahy, Czech Republic. **Local agent:** Radiant Export Import Enterprise. Uttara, Dhaka.	Glypressin 1mg/8.5ml Solution for injection. Terlipressin (as acetate) INN Injection. **Vasoconstrictor**	Bleeding from oesophageal varices. .	**Contraindication**: Contraindicated in pregnancy. Hypersensitivity to terlipressin or any other excipients of the product. **Renal impairment:** use with caution in chronic renal failure **Side effects**: abdominal cramps, diarrhoea, hypertension, hypotension, peripheral ischaemia, pallor, arrhythmia, bradycardia, headache; less commonly nausea, vomiting, hot flushes, angina, myocardial infarction, tachycardia, intestinal ischaemia, bronchospasm, respiratory failure, pulmonary oedema, convulsions, hyponatraemia; rarely dyspnoea; very rarely stroke, hyperglycaemia; also reported heart failure, skin necrosis. **Adverse effects:** Headache, Bradycardia, Peripheral vasoconstriction, Peripheral ischaemia, Facial pall or Hypertension, Transient abdominal cramps, Transient diarrhea.	Czech Republic & Australia	New	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj

42.	Manufactturer : Baxter AG, Vienna, Austria. Impoter : Swadesh , 30, Bijoy Nagaar Road (3rd Fl.) Dhaka-1000.	Tisseel Lyo-Powders and solvents for sealant Component-1: **Sealer Protein Solution** Sealer Protein Concentrate, Lyophilized (Tissel poeder), reconstituted with aprotinin Solution. Component-2: **Thrombin Solution** Thrombin, Lyophilized, reconsitituted with Calcium chloride solution	Supportive treatment where standard surgical techniques are insufficient - for improvement of haemostasis - as a tissue glue to promote adhesion/sealing or as suture support: - in gastrointestinal anastomoses -in neurosurgery where contact with cerebro-spinal fluid or dura mater can occur - For mesh fixation in hernia repair, as an alternative or adjunct to sutures or staples.	**Contraindications:** TISSEEL Lyo must not be applied intravascularly. Hypersensitivity to the active substances or to any of the excipients. TISSEEL Lyo alone is not indicated for the treatment of active or spurting arterial or venous bleeding which is not controlled by conventional surgical techniques. **Side Effects:** Hypersensitivity or allergic reactions (which may include angioedema, burning and stinging at the application site, bradycardia, bronchospasm, chills, dyspnoea, flushing, generalized urticaria, headache, hives, hypotension, lethargy, nausea, pruritus, restlessness, tachycardia, tightness of the chest, tingling, vomiting, wheezing) may occur in rare cases in patients treated with fibrin sealants / haemostatics. In isolated cases, these reactions have progressed to severe anaphylaxis. Such reactions may especially be seen, if the preparation is applied repeatedly, or administered to patients known to be hypersensitive to aprotinin or any other constituents of the product. Even if a second treatment with TISSEEL Lyo was well tolerated, a subsequent administration of TISSEEL Lyo or systemic administration of aprotinin may result in severe anaphylactic reactions. In the event of hypersensitivity reactions the administration has to be discontinued immediately. Soft tissue injection of TISSEEL Lyo carries the risk of an anaphylactoid reaction and / or local tissue damage. Reactions to antibodies against components of fibrin sealant / haemostatic products may occur rarely.	Austria		Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj

43.	Norma Chemicals Ltd UK **Local agent:** Zas Corporation, Outer circular Extention Road . Banglamotor . Dhaka	Dubam Spray Ethyl salicylate/glycol salicylate/methyl nicotinate/methyl salicylate	Symptomatic relief of muscular aches and pains. For external use only.	**Contraindications:** Do not use when there is extensive damage to the skin. Do not uses in the skin remain largely intact**.** **Side-effects:** May produce local irritation, numbness in the treated area	U.K	New	*c‡qvRb ‡bB wea‡q Av‡e`b bvgÄyK iv †h‡Z cv‡i |*	*c‡qvRb ‡bB wea‡q Av‡e`b bvgÄyK iv nj |*

Annex-C

Proposed Product for locally manufacture (Veterinary Products):

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †Rvlbv‡iK bvg	wb‡`kbv	Contra-indication & Side-effect	Status (New Molecule/ Existing)	Av‡e`bKvix c`Ë †idv‡iÝ	‡UKwbK¨vj mve-Kvgvwi 64 Zg mfvi wm×vš	mfvi wm×vš
01.	Al-Madina Pharmaceuticals Ltd.	Ammonioum Chloride 17.06 gm + Didecyl Dimethyl Benzyl Ammonium Chloride 7.80 gm + Isopropanol 14.625 gm + Gluteraldehyde 10.725 gm + Pine Oil 2.0 gm + Propylene Glycol 0.10 gm100 ml Liquid Ammonioum Chloride INN 17.06 gm + Didecyl Dimethyl Benzyl Ammonium Chloride INN 7.80 gm + Isopropanol BP 14.625 gm + Gluteraldehyde BP 10.725 gm + Pine Oil BP 2.0 gm + Propylene Glycol BP 0.10 gm100 ml **Antibacterial**	It is used to destroy bacterial, virus, fungi, algae, protozoa, mycoplasma and other microorganisms including sopre.	Contra-indication : N/A Side Effect : At high dosage is laying hens egg drop and in broilers growth inhibition and polyneuritis can occur	New		‡idv‡iÝmn cÖvPxiv‡q Dc¯vc‡bi Rb¨ ejv ‡h‡Z cv‡i|	‡idv‡iÝmn cÖvPxiv‡q Dc¯vc‡bi Rb¨ ejv হল|
02.	ACI Limited	Doxycycline Hyclate 20.00g + Tylosin Tartrate 10.00g + Colistin Sulphate 2.632g + Bromhexine HCl 0.50g + Amantadine HCl 4.00g/100 Powder (Vet) (For veterinary use only) Doxycycline Hyclate BP 20.00g + Tylosin Tartrate BP 10.00g + Colistin Sulphate BP2.632g + Bromhexine HCl BP 0.50g + Amantadine HCl BP 4.00g/100 **Antibiotic**	For the prevention and treatment of respiratory, CRD & gastrointestinal tract infection of large Animal & Poultry caused by Mycoplasma, Pasteurella, Streptococcus, Hemophilus, Rickettsia, Camphylobacter, Staphylococcus, Bordetilla, Salmonella, Clamydia etc and control of Avian Influenza in Poultry.	**Contra-indications**: Animal hypersensitivity to Enramycin or any other ingredient of the product. Concurrent administration of penicillin's, Cephalosporin's, & Quinolones is contraindicated. **Adverse Effects:** Diarrhoea may occur in some cases. Discoloration of teeth in young animals.	New		‡idv‡iÝmn cÖvPxiv‡q Dc¯vc‡bi Rb¨ ejv ‡h‡Z cv‡i|	‡idv‡iÝmn cÖvPxiv‡q Dc¯vc‡bi Rb¨ ejv হল|

03	ACI Limited	Colistin Sulphate 0.4g + Neomycin Sulphate 7.00g + Clortetracycline HCl 8.00g/100g Powder (Vet) (For Veterinary use only) Colistin Sulphate BP 0.4g + Neomycin Sulphate BP 7.00g + Clortetracycline HCl BP 8.00g/100g **Antibiotic**	Early chick's mortality, CRD, CCRD, Fowl Cholera, Bacterial enteritis, Blue Comb, Coli-bacillosis, Synovitis, Septicemia and Omphalitis.	**Contra-indications:** Hypersensitivity to any of the ingredients. Administration to animals with a serious impaired renal function and/or liver function. **Side Effects:** Hypersensitivity reactions	New		‡id‡iÝmn clyivq Dc¯vc‡bi Rb¨ ejv ‡h‡Z cv‡i	‡id‡iÝmn clyivq Dc¯vc‡bi Rb¨ ejv হল।
04.	ACI Limited	Erythromycin 5.0 gm+ Colistin Sulphate 0.25gm + Bromhexine HCl 0.375g/100gm Powder (Vet) (For veterinary use only) Erythromycin Thiocyanate BP 5.403g (eqv. to 5g Erythromycin) + Colistin Sulphate BP 0.25g (Eqv. to 5,000,000IU Colitin Sulphate) + Bromhexine Hydrochloride BP 0.375g/100g **Antibiotic**	Mainly effective against Mycoplasma gallisepticum, E. Coli, infectious Coryza, Infectious sinusitis and synovitis in chicken, Very effective in therapy of chronic respiratory disease (CRD) Effective in stress conditions due to vaccination, debeaking, transporting, rehousing, changes in feed and weather.	**Contra-indications:** Hypersensitivity to any of the ingredients. **Side Effects:** Hypersensitivity reactions	New		‡id‡iÝmn clyivq Dc¯vc‡bi Rb¨ ejv ‡h‡Z cv‡i	‡id‡iÝmn clyivq Dc¯vc‡bi Rb¨ ejv হল।
05.	Acme Laboratories Ltd.	Lincomycin 22.200 gm + Spectinomycin 44.400 gm/100 gm Water Soluble Powder (WSP) [For Veterinary Use Only Lincomycin Hydrochloride USP 25.177gm eq. to Lincomycin 22.200gm + Spectinomycin Sulphate Tetrahydrate BP 67.100gm eq. to 44.40gm Spectinomycin /100gm **Antibiotic**	The combination of Lincomycin and Spectinomycin WSP is indicated for the treatment of diseases caused by micro-organisms susceptible to Lincomycin and Spectinomycin. Poultry: Treatment of chronic respiratory diseases (CRD) caused by Mycoplasma	**Contraindication:** Lincomycin and Spectinomycin is contraindicated in animals known to have hypersensitivity to Lincomycin and/or Spectinomycin or any of its components. It should not administer to animals with a serious impaired renal function. It should not use in layers producing eggs for human	New	USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

			infections as well as any infection due to *E. coli* bacteria sensitive to Lincomycin and Spectinomycin. Pigs: Gastrointestinal diseases (e.g. swine dysentery caused by Treponema (Spirochates, Vibrio) spp., bacterial enteritis as well as complications caused by *Escherichia coli*, Salmonella spp., Streptococcus and Staphylococcus, inflammation in the joints (infectious arthritis caused by Streptococcus, Staphylococcus and Mycoplasma).	consumption. **Side effects:** Hypersensitivity reactions may occur.				
6.	**Bridge Pharmaceuticals Ltd.**	(Levofloxacin 10gm + Colistin 1.5gm + Lactic Acid 32ml + Acetic Acid 11.5ml)/ 100ml Oral Solution Levofloxacin Hemihydrate USP 10.2492gm eq. to 10gm Levofloxacin + Colistin Sulphate BP 1.6243gm eq. to 1.5gm Colistin + Lactic Acid BP 32 ml+ Acetic Acid 11.5ml/ 100ml **Antibiotic**	Levofloxacin is a third generation fluoroquinolone antibiotic. It is twice as active as it's isomer ofloxacin, effective against number of Gram-positive, Gram-negative and specifically effective against the organisms that cause atypical pneumonia. Levofloxacin is one of the so called respiratory quinilones. Levofloxacin is to be	**Contraindication**: None **Side-effect:** There have been no reports of custaneous drug reaction resulting from fluoroquinolones usage in the veterianary.	New		‡id‡iÝmn cÖvPwq Dc¯'vc‡bi Rb¨ ejv ‡h‡Z c‡i|	‡id‡iÝmn cÖvPwq Dc¯'vc‡bi Rb¨ ejv হল|

			considered a drug of last resort when all other antibiotics have failed. To control Gram-Positive, GHram-Negative bacteria and most spp. of mycoplasma. It controls infections effectively against drug resistant bacteria causing severe infections like CRD, CCRD, Callisepticaemia, Salmonellosis, infections coryza, Fowl cholera, Necrotic Enteritis, Gangrenous Dermatitis, Non Specific & Secondary bacterial infections during viral diseases etc. It is a ideal combination for controlling E-coll as both fluoroquinolone and colistin sulphate are highly effective against E-coll and approach E.Coli in two ways, i.e. fluoroquinolone systemically and colistin through out.					
7.	Acme Laboratories Ltd.	Tylvalosin 62.50gm/100gm Water Soluble Powder (WSP) (For Veterinary Use Only) Tylvalosin Tartrate INN 71.500 g eq. to 62.50gm Tylvalosin)/ 100gm	<u>Poultry</u> Tylvalosin is indicated for the prevention and treatment of Mycoplasmosis and necrotic enteritis, enteritis resulting in wet litter syndrome and cholangio- hepatitis in chickens, replacement pullets	**Contraindication:** Animals hypersensitive to Tylvalosin are contraindicated. **Side effects**: No adverse reactions related to the drug were observed	New	USFDA	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

		Antibiotics	and turkeys. It is also effective against diseases caused by *Ornithobacterium rhinotracheale* (ORT) of poultry. Swine Tylvalosin is indicated for the treatment and prevention of Swine Enzootic Pneumonia, Ileitis (Porcine Proliferative Enteropathy and Swine Dysentery). Tylvalosin is also active against colitis in pigs.	during clinical or target animal safety trials.				
8.	Acme Laboratories Ltd.	Ceftiofur 0.500gm/Vial Powder for Injection (PFI) (For Veterinary Use Only) Ceftiofur Sodium Sterile Powder INN 0.532mg eq. to 0.500gm Ceftiofur/ Vial **Antibiotics**	Cattle Indicated for the treatment of bovine respiratory disease (shipping fever, pneumonia) associated with *Mannheimia haemolytica*, *Pasteurella multocida* and *Histophilus somni*. It is also indicated for the treatment of acute bovine interdigital necrobacillosis (foot rot, pododarmatitis) associated with *Fusobacterium necrophorum* and *Bacteroides melaninogenicus*. Day-old Chicks Indicated for the control of early mortality, associated with *E. coli* organism susceptible to Ceftiofur, in day-old chicks.	**Contraindication:** Contraindicated in animals previously found to be hypersensitive to the drug. **Side effects:** The use of Ceftiofur may result in some signs of immediate and translent local pain to the animal. Results from a five day tolerance study in normal feeder calves indicated that formulated Ceftiofur was well tolerated at 25 times.	1 gm/vial Injection	USFDA	*Abjgv`b Kiv †h‡Z cv‡i |*	*Abjgv`b Kiv nj |*
09.	Acme Laboratories Ltd.	Gamithromycin 1.50gm/10ml Injection Vial [For Veterinary Use Only] Gamithromycin INN 1.50gm/10 ml Vial **Antibiotics**	Gamithromycin is indicated for the treatment of bovine respiratory disease (BRD) associated with *Mannheimia haemolytica, Pasteurella multocida, Histophilus somni* and *Mycoplasma bovis* in beef and non-lactating dairy cattle. Gamithromycin is also indicated for the control of	**Contraindication:** Gamithromycin Injection is contraindicated in animals known to have hypersensitivity to the drug or any of its components. **Side effects**: Transient animal	New Molecule	USFDA	*Abjgv`b Kiv †h‡Z cv‡i |*	*Abjgv`b Kiv nj |*

			respiratory disease in beef and non-lactating dairy cattle at high risk of developing BRD associated with *Mannheimia haemolytica* and *Pasteurella multocida.*	discomfort and mild to moderate injection site swelling may be seen in cattle treated with Gamithromycin.				
10.	Acme Laboratories Ltd.	Gamithromycin 4.50gm/30ml Injection Vial [For Veterinary Use Only] Gamithromycin INN 4.50gm/30ml Vial **Antibiotics**	Gamithromycin is indicated for the treatment of bovine respiratory disease (BRD) associated with *Mannheimia haemolytica, Pasteurella multocida, Histophilus somni* and *Mycoplasma bovis* in beef and non-lactating dairy cattle. Gamithromycin is also indicated for the control of respiratory disease in beef and non-lactating dairy cattle at high risk of developing BRD associated with *Mannheimia haemolytica* and *Pasteurella multocida.*	**Contraindication:** Gamithromycin Injection is contraindicated in animals known to have hypersensitivity to the drug or any of its components. **Side effects**: Transient animal discomfort and mild to moderate injection site swelling may be seen cattle treated with Gamithromycin.	New Molecule	USFDA	*Abjgv`b Kiv †h‡Z cv‡i |*	*Abjgv`b Kiv nj |*
11.	Acme Laboratories Ltd.	Menbutone (Genabilic Acid) 1.0 gm/10 ml Injection Vial [For Veterinary Use Only] Menbutone (Genabilic Acid) INN 1.0 gm/10 ml Vial **Choleretic agent**	Menbutone is indicated in situations where stimulation of digestive secretions is required such as: Cattle: Indigestion, toxemia, ketosis, anorexia, liver and pancreatic insufficiency. Pigs: Indigestion, anorexia, poisoning, liver and pancreatic insufficiency.	**Contraindication:** Menbutone Injection is contraindicated in animals known to have hypersensitivity to the drug or any of its components. It should not administer to animals with cardiac disorders, hyperthermia or blockage in the bile ducts. Menbutone Injection should not administer to cats in any case. **Side effects:** After intravenous administration, salivation, lachrymosity, tremors,	New Molecule	Veterinary Formulary, 5th Edition, P: 328	*Abjgv`b Kiv †h‡Z cv‡i |*	*Abjgv`b Kiv nj |*

				spontaneous urination and defecation may occur. After intramuscular administration, reaction at the injection site (oedema, haemorrhage, necrosis) may occur.				
12.	Acme Laboratories Ltd.	Menbutone (Genabilic Acid) 3 g/30 ml Injection Vial [For Veterinary Use Only] Menbutone (Genabilic Acid) INN 3 g/30 ml Vial **Choleretic agent**	Menbutone is indicated in situations where stimulation of digestive secretions is required such as: Cattle: Indigestion, toxemia, ketosis, anorexia, liver and pancreatic insufficiency. Pigs: Indigestion, anorexia, poisoning, liver and pancreatic insufficiency.	**Contraindication:** Hypersensitivity to Bromhexine. Animal with an impaired liver function. **Side effects:** No undesirable effects are to be expected when the prescribed dosage regimen is followed.	New Molecule	Veterinary Formulary, 5th Edition, P: 328	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
13.	Acme Laboratories Ltd.	Bromhexine HCl 100mg Bolus (For Veterinary Use Only) Bromhexine HCl BP 100mg **Expectorant**	Respiratory disease where excess tenacious mucus is present.	**Contraindication:** Hypersensitivity to Bromhexine. Animal with an impaired liver function. **Side effects:** No undesirable effects are to be expected when the prescribed dosage regimen is followed.	10 mg/gm Powder	Veterinary Formulary, 5th Edition, P: 355-356	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
14.	Acme Laboratories Ltd.	Bromhexine HCl 0.08gm/100ml Oral Solution (For Veterinary Use Only) Bromhexine HCl BP 0.08gm/100ml **Expectorant**	Respiratory disease where excess tenacious mucus is present.	**Contraindication:** Hypersensitivity to Bromhexine. Animal with an impaired liver function. **Side effects:** No undesirable effects are to be expected when the prescribed dosage regimen is followed.	10 mg/gm Powder	Veterinary Formulary, 5th Edition, P: 355-356	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
15.	Al-Madina Pharmaceuticals Ltd.	Zinc Bolus Zinc Sulphate Monohydrate USP 552mg eq. to elemental Zinc 200mg **Mineral**	Treatment of Zinc deficiency. Recurrent respiratory tract infections, diarrhea, loss of appetite, severe growth retardation.	**Contraindication:** Known case of hypersensitivity to zinc or zinc product. **Side- Effect:** Mild side-effects have been observed with zinc therapy, such as abdominal pain and dyspepsia, gastric Ulcer,			***‡idv‡iÝmn cÖvivq Dc¯vcb‡bi Rb¨ ejv ‡h‡Z cv‡i |***	***‡idv‡iÝmn cÖvivq Dc¯vcb‡bi Rb¨ ejv হল|***

				pancreatitis,				
16.	Al-Madina Pharmaceuticals Ltd.	Zinc Sulphate Monohydrate 2.0gm/100ml Liquid Zinc Sulphate Monohydrate BP 2gm/100ml **Mineral**	Treatment of Zinc deficiency. Quick growth of Bone, to avoid bony deformities, quick weight gain, increase egg production and fertility, increase immunity development of birds, secondary treatment in case diarrhea.	**Contraindication:** No recognized side effect is identified in recommended dose. **Side effect:** It has no side effect at recommended dose.	500 ml & 1000 ml Liquid		*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
17.	Square Pharmaceuticals Ltd. (Agrovet Division), Dhaka Unit, Kaliakoir, Gazipur	Calcium 2.14gm/100ml Injection Calcium Gluconate USP 23.00gm eq. to 2.14gm Calcium/100ml **Electrolyte Supliment**	use in cattle as an aid in treating uncomplicated milk fever (parturient paresis)	**Contra-indication:** Calcium Gluconate Injection is contraindicated in animals hypersensitive to any of the active ingredients. **Side effect:** Nausea, vomiting, anorexia, abdominal pain, muscle weakness	New	Calcium Gluconate 23% Solution Nova- Tech, Inc. Grandland, USA	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
18.	Al-Madina Pharmaceuticals Ltd.	Zinc Sulphate Monohydrate 20.00 mg + Thiamine HCl 5.0mg + Riboflavin Sodium-5 Phosphate 1.50 mg + Pyridoxine HCl 8.0mg + Nicotinamide 4.0 mg/ml Oral Liquid Zinc Sulphate Monohydrate BP 20.00 mg + Thiamine HCl BP 5.0mg + Riboflavin Sodium-5 Phosphate BP 1.50 mg + Pyridoxine HCl BP 8.0mg + Nicotinamide BP 4.0 mg/ml **Minerals+ Vitamin**	Treatment of Zinc deficiency. Recurrent respiratory tract infections, diarrhea, loss of appetite, severe growth retardation, deformed bone formation, impaired immunological status	**Contraindication:** It is contraindicated for the birds hypersensitive to any of its contents. **Side effect:** It has no side effect at recommended dose.	500 ml & 1000 ml Liquid		*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
19.	Al-Madina Pharmaceuticals Ltd.	Sodium Chloride 26.6 gm + Sodium Bicarbonate 50 gm + Dextrose Anhydrous 18 gm + Potassium Chloride 5 gm + Vitamin A Acetate 0.40 gm/100 gm Powder	For the treatment of water and electrolytes imbalance as in Cholera,gastroenteriis, diarrhoeas, water & electrolytes	**Side effect:** It has no side effect at recommended dose	500.0 gm & 1.0 kg powder		*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

		Sodium Chloride BP 26.6 gm + Sodium Bicarbonate BP 50 gm + Dextrose Anhydrous BP 18gm + Potassium Chloride BP 5 gm + Vitamin A Acetate BP 0.40gm/100 gm **Minerals+ Vitamin**	loss, fever,hyperventilation, white bowel in animals					
20.	Al-Madina Pharmaceuticals Ltd.	Acetyl Salicylic Acid 6.70 gm + Vitamin C 20 gm/100 gm Powder Acetyl Salicylic Acid BP 6.70 gm + Vitamin C USP 20 gm/100 gm **NSAID**	In Broilers, prevents of heart stock, counters heat stress mortality, improves growth performanance, enhances disease resistance, and relieves pain in inflammation. In case of breeders and layers, protect from heat stress, increase of egg production, improves egg shellquality, restore fertility and hatchability, and enhances disease resistance	**Contraindications:** There is no contraindication to the administration of Acetyl salicylic acid **Side-Effects:** It has no side effect at recommended dose.	100 gm & 500 gm &1 kg		*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
21.	Al-Madina Pharmaceuticals Ltd.	Vitamin -C 2.0 gm + Glucose 98.0 gm/100 gm Powder Vitamin -C BP 2.0 gm + Glucose BP 98.0 gm/100 gm **Vitamin**	Supplement of Vitamin C & glucose deficiency, Prevention and treatment of scurvey, Adjunct in the treatment of wounds, infections, trauma, fractures, burns, cold exposure, stress, infertility, develop Hatching	**Contraindications:** There is no contraindication to the administration of vitamin-c. **Side-Effects:** Generally ascorbic acid is well tolerated. However, few side-effects including stomach upset, diarrhea, mouth sores.	100 gm & 500 gm powder		*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
22.	Al-Madina Pharmaceuticals Ltd.	Vitamin E Acetate 10.0gm + Sodium Selenite 0.05gm/100ml Liquid Vitamin E Acetate BP 10.0gm + Sodium Selenite BP 0.05gm/100ml	Vitamin E is indicated in the treatment &prevention of vitamin E deficiencyin various conditions. To increase the growth &	**Contrainctiontera:** Known hypersensivity to vitamin-E. **Side-Effect:** Vitamin-E is usually well tolerated. Large doses may cause	100 ml, 500ml & 1.0 Liquid		*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

		Vitamin	productivity of poultry.	diarrhea,abdominal pain &othergastro-intestinaldisturbances.				
23.	Al-Madina Pharmaceuticals Ltd.	Vitamin – C (Ascorbic Acid) BP 1000 mg Powder Vitamin – C (Ascorbic Acid) BP 1000 mg **Vitamin**	Supplement of Vitamin C deficiency. Helps to recover from stress conditions like heat stress, transport stress, vaccination stress, debeaking stress, stress in different diseased conditions. Increases immunity of the body. Helps to produce steroid hormone	**Contraindications:** There is no contraindication to the administration of vitamin - c. **Side-Effects:** No recognized side effect is identified in recommended dose.	100 gm & 500 gm		Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
24.	Al-Madina Pharmaceuticals Ltd.	Ascorbic Acid 0.50 gm + Riboflavin Sodium-5 Phosphate 0.40gm +Thiamine HCl 0.20 gm+ Pyridoxine HCl 0.150 gm + Nicotinamide 0.75gm + Calcium D Pantothenate 0.45 gm + Vitamin B_{12} (0.1 %) 0.001gm + Folic Acid 0.025 gm + Biotin (2%) 0.0015gm/100gm Powder Ascorbic Acid BP 0.50 gm + Riboflavin Sodium-5 Phosphate BP 0.40gm +Thiamine HCl BP 0.20 gm+ Pyridoxine HCl BP 0.150 gm + Nicotinamide BP 0.75gm + Calcium D Pantothenate BP 0.45 gm + Vitamin B_{12} 0.1 % BP 0.001gm + Folic Acid BP 0.025 gm + Biotin 2% BP 0.0015gm/100gm **Vitamin**	Curl Toe Paralysis, Dermatitis, Diarrhea. All types of stress like heat stress, adverse environmental conditions, transportation, shed changing, overcrowding, culling, vaccination, antibiotics and anthelmintic use etc. To prevent thin-shelled egg formation	**Contraindications:** It is contraindicated for the birds hypersensitive to any of its contents. **Side effect :** It has no side effect at recommended dose. But mega-doses of vitamin C may cause diarrhea, abdominal bloating, iron-over absorption.	100 gm & 500gm powder		Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
25.	Al-Madina Pharmaceuticals Ltd.	Thiamine Hydrochloride 6.0 gm+ Riboflavin	It helps to make over the deficiency of Thiamine	**Contraindications:** naphylactic reaction**,** with levodopa.	100 gm & 500		Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

		1.10 gm + Pyridoxine Hydrochloride 0.61 gm/100 gm Powder Thiamine Hydrochloride BP 6.0 gm+ Riboflavin BP 1.10 gm + Pyridoxine Hydrochloride BP 0.61 gm/100 gm **Vitamin**	Hydrochloride, Riboflavin and Pyridoxine Hydrochloride like diarrhea, Curl Toe Paralysis, loss of appetite, lame, increase digestive power.	**Side effect:** It has no side effect at recommended dose.	gm			
26.	Acme Laboratories Ltd.	Zilpaterol Hydrochloride 4.800 g + Melengestrol Acetate 0.110gm/100gm Water Soluble Powder (WSP) [For Veterinary Use Only] Zilpaterol Hydrochloride INN 4.800 g + Melengestrol Acetate USP 0.110 g/100 gm	For increased rate of weight gain, improved feed efficiency, increased carcass leanness, and suppression of estrus (heat) in heifers fed in confinement for slaughter during the last 20 to 40 days on feed.	**Contraindication:** Contraindicated in animals known to have hypersensitivity to Zilpaterol Hydrochloride and/or Melengestrol Acetate or any of the components of the product. **Side effects:** No undesirable effects have been found at recommended doses.	New Molecule	USFDA	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
27.	Al-Madina Pharmaceuticals Ltd.	Ammonium Bicarbonate BP 25.0 gm + Nuxvomica BP 7.0 gm + Sodium Bicarbonate BP 65.0 gm + Gentian Powder BP 1.5 gm + Ginger Powder BP 1.5 gm/100 gm powder Ammonium Bicarbonate BP 25.0 gm + Nuxvomica BP 7.0 gm + Sodium Bicarbonate BP 65.0 gm + Gentian Powder BP 1.5 gm + Ginger Powder BP 1.5 gm	Inappetance, indigestion, gastritis, bloat or tympany of animals.	**Contraindications: N/A** **Side effect : N/A**	20 gm,500gm powder		*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

Annex-D

Proposed Product for Import (Veterinary):

bs	c¯ZKvi‡Ki bvg	JI‡ai bvg I †RvlbviK bvg	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	Status (New Molecule/ Existing)	‡UKwbK¨vj mve-KvgvUi 64 Zg mfvi wm×vš	mfvi wm×vš
1.	Manufacturer: Merck Saude Animal Ltda., Brazil **Owner/Distributor.:** MERIAL Ltd., Georgia (Advance Animal Science Co. Ltd.; 2/10 Block-B, Lalmatia, Dhaka-1207)	Ivomec Super Injection Ivermectin In House 1gm + Clorsulon In House 10gm/100ml Anthelmintic	For the treatment and control of Gastrointestinal roundworms, lungworms, adult Liver fluke, eyeworm, warbles, mites and lice of beef and non-lacting dairy cattle.	**Contraindications:** This product is not to be used intramuscularly or intravenously. IVOMEC Super Injection for Cattle is a low-volume product registered for use in cattle. It should not be used in other species as severe adverse reactions, including fatalities in dogs, may occur. **Side effects:** Transitory discomfort has been observed in some cattle following subcutaneous administration. A low incidence of soft tissue swelling at the injection site has been observed. These reactions disappeared without treatment.	Brazil USFDA	New	Abjgv`b Kiv †h‡Z cv‡i |	Abjgv`b Kiv nj |
2.	Manufacturer: Merck Sharpe and Dohme de Puerto Rico Inc., USA Owner/Distributor.: MERIAL Ltd., Georgia (Advance Animal Science Co. Ltd.; 2/10 Block-B, Lalmatia, Dhaka-1207)	Heartgard® Plus Chewable Tablet Ivermectin 68mcg + Pyrantel Pamoate 57mg Anthelmintic	For use in dogs to prevent canine heartworm disease by eliminating the tissue stage of heartworm larvae (Dirofilaria immitis) for a month (30 days) after infection and for the treatment and control of ascarids (Toxocara canis, Toxascaris leonina) and hookworms (Ancylostoma caninum, Uncinaria stenocephala, Ancylostoma braziliense).	**Contraindications: None** **Adverse Reaction:** In clinical field trials with Heartgard Plus, vomiting or diarrhea within 24 hours of dosing was rarely observed (1.1% of administered doses). The following adverse reactions have been reported following the use of HEARTGARD: Depression/lethargy, vomiting, anorexia, diarrhea, mydriasis, ataxia, staggering, convulsions and hypersalivation.	Brazil USFDA	New	Abjgv`b Kiv †h‡Z cv‡i |	Abjgv`b Kiv nj |

3.	**Manufacturer:** M/S Shinil Bio-gen Co. Ltd., **Old Address:** 254-10,Dugok-rl, Shinam-myeon,Yeasan-gum, Choongchungnam-do, Korea. New Address: 235-18,Chusa-ro, Choongchungnam-do, Korea. **Importer**: M/S Pharma & Firm 3/2, City Heart Building, 67, Naya Paltan, Dhaka-1000.	Tia-Forte sol Tiamulin Hydrogen Fumarate 125gm /Litre Antibacterial	For the prevention and treatment of disease caused by pathogens susceptible to Tiamulin. **Chicken** : Prevention and treatment of Chronic respiratory disease(CRD) and Arthritis. Swine : Prevention and treatment of pneumonia, Swine dysentery, Proliferative ileitis And Arthritis	**Contraindication** -None **Side-effect** -Redness of the skin -Salivation -vomiting -CNS depression (For Pigs) **Withdraw Period:7 days**	Korea	New	*Abyjgvw` b Kiv †h‡Z cv‡i |*	*Abyjgvw` b Kiv nj |*
4.	**Manufacturer:** M/S Shinil Bio-gen Co. Ltd., **Old Address:** 254-10,Dugok-rl, Shinam-myeon,Yeasan-gum, Choongchungnam-do, Korea. New Address: 235-18,Chusa-ro, Choongchungnam-do, Korea. **Importer**: M/S Pharma & Firm 3/2, City Heart Building, 67, Naya Paltan, Dhaka-1000.	**Flotec 200 sol** **Each L Contains,** Florfenicol. ……..200g Antibacterials	For the treatment of diseases caused by pathogens susceptible to Florfenicol. **-Poultry** : Salmonellosis and Colibacillosis. **-Swine** : Respiratory disease caused by Actinobacillus pleuropneurnonia, Pasteurella multocida and Mycoplasma hyopneumoniae, Streptococcosis and Salmonellosis.	**Contraindication** **-**None **Side-effect** -None **Withdrawal Period**- Swine 16 days, Poultry-5 days	Korea	New	*Abyjgvw` b Kiv †h‡Z cv‡i |*	*Abyjgvw` b Kiv nj |*

5.	**Choong Ang Biotech Co., Ltd., Korea** **Locan agent: SHAMIM PHARMA** 112/A, 3rd Floor, Senpara Parbata, Mirpur 10, Dhaka-1216, Bangladesh	Cefalexin Hydrate, Colistin Sulfate Cefalexin Hydrate 100 gm + Colistin Sulfate – 250,000000 IU/kg Antibiotic	**Poultry:** Cefaxosil is highly effective active against infections caused by Gram positive and Gram negative Bacteria and Mycoplasma. Cefaxosil is indicated for treatment of Enteritis, Colibacillosis, Salmonellosis, Prevention of early Chick mortality, Chronic respiratory disease (CRD), Fowl Typhoid, Infectious coryza. **Cattle, Foal, Sheep, Goat:** For the treatment of following diseases Enteritis, Bacterial Pneumonia, Colibacillosis, Salmonellosis.	**Contraindication :** Cefalexin is contraindicated in patients with known allergy to the cephalosporins group of antibiotics. Cefalexin should be given cautiously to patients who have shown hypersensitivity to other drugs. Cefalexin is contraindicated in patients with acute porphyria. Colistin act synergistically when combined with potentiated sulfonamides, tetracyclines, and some other antibacterials; they also reduce the activity of endotoxins in body fluids and may be beneficial in endotoxemia. **Side effects:** The most common adverse effects of Cefalexin, like other oral Cephalosporins, are gastrointestinal (stomach area) disturbances and hypersensitivity reactions. Gastrointestinal disturbances include nausea, vomiting, and diarrhea. Colistin; The main toxicities described with intravenous treatment are nephrotoxicity (damage to the kidneys) and neurotoxicity (damage to the nerves), but this may reflect the very high doses given.	Korea	New	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
6.	Choong Ang Biotech Co., Ltd., Korea	Amoxicillin Hydrate, Erythromycin Thiocyanate, Colistin Sulfate	**Poultry:** Ery Moxol is highly effective active against infections caused by Gram positive and Gram	**Contraindication:** A history of allergic reaction to any of the penicillins is a contraindication. β-	Korea	New	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

	Local Agent: NEON PHARMA 184 (2nd Floor), Arambagh, Motijheel Dhaka-1000,	Amoxicillin Hydrate 100gm + Erythromycin Thiocyanate 50gm + Colistin Sulfate 10,000,000 IU/kg Antibiotic	negative Bacteria and Mycoplasma. Ery Moxol is indicated for treatment of Colibacillosis, Salmonellosis, Mycoplasmosis, Chronic respiratory disease (CRD), Infectious coryza, Streptococcosis, Staphylococcosis in poultry. **Cattle, Foal, Sheep, Goat:** For the treatment of following diseases Bacterial Pneumonia, Colibacillosis, Salmonellosis.	lactams in general interact chemically with the aminoglycosides and should not be mixed in vitro. Macrolide preparations for parenteral administration are incompatible with many other pharmaceutical preparations. Erythromycin and troleandomycin are microsomal enzyme inhibitors that depress the metabolism of some drugs. Colistin act synergistically when combined with potentiated sulfonamides, tetracyclines, and some other antibacterials; they also reduce the activity of endotoxins in body fluids and may be beneficial in endotoxemia. **Side effects:** Amoxicillin is similar to those for other β-lactam antibiotics, including nausea, vomiting, rashes, and antibiotic-associated colitis. Loose bowel movements (diarrhea) may also occur. Erythromycin estolate may be hepatotoxic and cause cholestasis; it may also induce vomiting and diarrhea, particularly when high doses are administered. Colistin; The main toxicities described with intravenous treatment are nephrotoxicity (damage to the kidneys) and neurotoxicity (damage to the nerves), but this may reflect the very high doses given.				
7.	Choong Ang Biotech Co., Ltd., Korea **Local Agent:** NEON PHARMA 184 (2nd Floor), Arambagh,	Egg and Chick Tonic Each kg contains: Oxytetracyclin HCl 55,000 gm+ Vitamin A- 6,500,000 IU, Vitamin D_3- 1,200,000 IU, Vitamin E- 2,500 IU,	Poultry: Helps on stress condition (stress of movement, rehousing, vaccination and change of feed). To improve feed utilization and absorption of nutrients. To increase egg production. To improve egg	**Contraindication:** Methoxyflurane anesthesia combined with tetracycline therapy may be nephrotoxic. The tetracyclines are less active in alkaline urine, and urine acidification can increase their	Korea	New	*Abyôgv` b Kiv †h‡Z cv‡i |*	*Abyôgv` b Kiv nj |*

	Motijheel Dhaka-1000,	Vitamin K_{3}- 1,000 mg, Vitamin B_{1}- 400 mg, Vitamin B_{2}- 3,000 mg, Vitamin B_{6}- 1,000 mg, Vitamin B_{12}- 3 mg, Calcium D pantothenate- 5,000 mg, Folic acid- 200 mg, Nicotinic acid- 6,000 mg, DL-Methionie- 16,000 mg, Glucose- q.s. Antibiotic	shell quality and carcass quality. Prevent and reduce Early chicks mortality and ensure healthy growth.	antimicrobial efficacy. The tetracyclines are also potentially nephrotoxic and are contraindicated (except for doxycycline) in renal insufficiency. Except the higher dosages, Vitamin, Minerals and Amino acids has no notable contra-indication. **Side effects:** Because several diverse effects may result from the administration of the tetracyclines, caution should be exercised. Severe and even fatal diarrhea can occur in horses receiving tetracyclines, especially if the animals are severely stressed or critically ill. In large doses, some vitamins have documented side-effects that tend to be more severe with a larger dosage. At high enough dosages, some vitamins cause side-effects such as nausea, diarrhea, and vomiting.				
8.	Eagle Vet. Tech Co., Ltd. Korea (ACI Ltd.)	Marbores 10% Inj. Injectable solution 20ml, 50ml, 100ml, 200ml & 250ml Marbofloxacine 100mg/ml Antibiotic	Treatment of cattle for respiratory disease and acute mastitis caused by bacteria which is susceptible to Marbofloxacin – Pasteurella multocida, Pasteurella haemolytica, Mycoplasma bovis, E. coli Acute mastitis.	**Contraindication**: Do not use for animals having hepatic & renal impairment. **Side Effect**: Gastrointestinal disorders, Central nervous system disorders may be appeared.	korea	New	*Abyôgv` b Kiv †h‡Z cv‡i |*	*Abyôgv` b Kiv nj |*
9.	Ewhapharmtek Corp., Korea (Fahat Trade International, 12/2 Purana Paltan Line, Dhaka-1000	Cephaxin 15% Water Soluble Powder Cefalexin Hydrate150gm/1kg Antibiotic	Cefalexin has a broad antibacterial action against both gram and negative bacteria.Prevent early chick mortality. Treatment of Coryza, Fowl typhoid, Fowl cholera, Bacillary white diarrhea and E.Coli. To prevent of secondary infections associated with viral diseases.	**Contraindication:** Cephalexin is contraindicated in patients with known allergy to cephalosporin group of antibiotics **Side effect:** Fever, Colitisanemia, hemorrhage, renal dysfunction and toxic nephropathy.	korea		*Abyôgv` b Kiv †h‡Z cv‡i |*	*Abyôgv` b Kiv nj |*

10.	Ewhapharmtek Corp., Korea (Fahat Trade International, 12/2 Purana Paltan Line, Dhaka-1000	Sunflox 15% Oral Solution Ciprofloxacin USP 150gm/Liter Antibiotic	For the treatment of the diseases caused by Mycoplasma, Pasteurella, Haemophilus, Staphylococcus, E. Coli, and Salmonella that are sensitive to Ciprofloxacin like CRD, CCRD, Enteritis, Colibacillosis, Salmonellosis, Fowl Cholera, Infectious Coryza, and Staphylococcosis.	Contraindications: Do not administer to animals with hypersensitivity to the preparation, as well as to animals with their liver of kidneys affected, during pregnancy period. Do not combine with non steroidal anti-inflammatory drug (NSAID), macrolide antibiotics, and tetracycline. -Co administration of ciprofloxacin with other drugs primarily metabolized by CYP1A2 result in increased plasma concentrations of these drugs and could lead to clinically significant adverse events of the co administered drug. -Concomitant administration with tizanidine is contraindicated. -Ciprofloxacin is contraindicated in persons with a history of hypersensitivity to ciprofloxacin, any member of the quinolone class of antimicrobial agents, or any of the product components. -Local I.V site reactions are more frequent if the infusion time is 30minutes or less. These may appear as local skin reactions that resolve rapidly upon completion of the infusion. Subsequent intravenous is not contraindicated unless the reactions recur or worsen. Side effect: Don't administer to animals with hypersensitivity to the preparation, as well as to animals with their liver and kidney affected,during pregnancy period.	korea		Abjgv`b Kiv †h‡Z cv‡i |	Abjgv`b Kiv nj |
11.	Green Cross Veterinary Products Co. Ltd., Korea (ACI Ltd.)	Cipro 20% Solution 100ml, 500ml, 1L, 5L & 20L Ciprofloxacin 200g /L	Treatment and of CRD, CCRD, Colibacillosis, Salmonellosis, Fowl cholera, Infectious Coryza, Staphylococcosis of Poultry.	**Contraindication**: Do not use in poultry producing eggs for human consumption. **Side effect**: Unknown.	KOREA		Abjgv`b Kiv †h‡Z cv‡i |	Abjgv`b Kiv nj |

		Antibiotic						
12.	Inter chemie Werken de Adelaar B. V. The Netherlands **(Importer:** C. P. Bangladesh Co., Ltd.; House-28, Alaol Avenue, Sector-6, Uttara, Dhaka-1230**)**	Amprolin-300 WS Powder Amprolium HCl BP 300mg/gm Antibiotic	Amprolin-300 WS is indicated for coccidiosis caused by coccidia susceptible to amprolium (Eimeria spp.) or gastrointestinal infections for which it is therapeutically or prophylactically indicated to administer amprolium in calves, goats, sheep and poultry.	**Contra-indications:** The use of amprolium is prohibited from a laying age onwards. Do not administer to poultry whose eggs are intended for human consumption. Do not administer to animals with impaired hepatic and/or renal functions. Do not administer to turkeys before the age of 8 to 10 weeks., or animals with impaired liver and/or renal functions. **Side-effect:** Overdosage of amprolium can suppress weight gain in broilers and cause polyneuritis. Long-term administration of amprolium in high doses may result in thiamine (vitamin B1) deficiency in the host. To treat amprolium overdose, thiamine should be administered parenterally or orally.	USA	EUDRALEX Volume 4 and GMP+B1	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
13.	Inter chemie Werken de Adelaar B. V. The Netherlands **(Importer:** C. P. Bangladesh Co., Ltd.; House-28, Alaol Avenue, Sector-6, Uttara, Dhaka-1230**)**	COLEXIL Oral Solution Enrofloxacin 100mg + Colistin Sulphate 1,200,000IU/ml Antibiotic	Colexil Oral solution is indicated for gastrointestinal, respiratory and urinary tract infections caused by colistin and enrofloxacin sensitive micro-organisms like Campylobacter, E. coli, Haemophilus, Mycoplasma, Pasteurella and Salmonella spp. in poultry and swine.	**Contra-indications:** Hypersensitivity to colistin and/or enrofloxacin or to any of the excipients. Administration to animals with seriously impaired renal and/or hepatic functions. Cases of resistance against quinolones and/or colistin. Administration to poultry producing eggs for human consumption. Administration to pregnant or lactating animals. Administration of Colexil Oral in subtherapeutic doses or for prevention. **Side-effect:** All members of the quinolone family of antibiotics have the ability to cause articular lesions in young animals. Digestive alterations may appear, such as intestinal dysbiosis, accumulation of gases, mild diarrhoea or vommiting. Side-effects for quinolones like rash and central nervous system disturbance may occur. During a period of rapid growth, enrofloxacin may affect joint cartilage.	The Netherlands	EUDRALEX Volume 4 and GMP+B1	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
14.	Inter chemie Werken de Adelaar B. V. The Netherlands	NEMOVIT WS Powder Neomycin Sulphate Ph.Eur. 40 mg + Oxytetracycline HCl USP 60mg + Vitamin A, Retinol Acetate 7500 IU +	Nemovit WS is a highly effective combination of broad-spectrum antibiotics and vitamins. The product stimulates egg production, increases growth, improves feed conversion and	**Contra-indications:** Hypersensitivity to tetracyclines or aminoglycocides. Administration to animals with a seriously impaired renal and/or hepatic function. Concurrent	The Netherlands	EUDRALEX Volume 4 and GMP+B1	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

	(Importer: C. P. Bangladesh Co., Ltd.; House-28, Alaol Avenue, Sector-6, Uttara, Dhaka-1230**)**	Cholecalciferol (Vitamin D3) 1500 IU + α-tocopherol acetate (Vitamin E) 5.0 mg + Thiamine Hydrochloride (Vitamin B1) Ph. Eur. 1 mg + Riboflavin (Vitamin B2) Ph. Eur. 2 mg + Pyridoxine HCl (Vitamin B6) Ph. Eur. 2 mg + Cyanocobalamin (Vitamin B12) Ph. Eur. 7.5 µg + Ascorbic acid (Vitamin C) Ph. Eur. 25 mg + Ca-pantothenate USP 7.5 mg + Menadione Sodium Bisulphite (Vitamin K3) 5 mg + Nicotinamide Ph.Eur. 15 mg + Folic Acid USP 0.3 mg + DL- Methionine Ph.Eur. 30 mg + Lysine HCl USP 50mg/gm Antibiotic	is used as a vitamin supplement during periods of diseases and stress. Gastrointestinal, respiratory and urinary infections caused by oxytetracycline and neomycin sensitive micro-organisms, like Bordetella, Campylobacter, Chlamydia, E. coli, Haemophilus, Klebsiella, Mycoplasma, Pasteurella, Rickettsia, Salmonella, Staphylococcus and Streptococcus spp. in calves, goats, sheep, poultry and swine.	administration of bactericidal agents like penicillins. Administration to animals with an active microbial digestion. **Side-effect:** Hypersensitivity reactions may occur.				
15.	Inter chemie Werken de Adelaar B. V. The Netherlands **(Importer:** C. P. Bangladesh Co., Ltd.; House-28, Alaol Avenue, Sector-6, Uttara, Dhaka-1230**)**	**Aliseryl WS Powder** Erythromycin Thiocyanate 35mg + Oxytetracycline HCl USP 50mg + Streptomycin Sulphate Ph. Eur. 35mg + Colistin Sulphate Ph. Eur. 200000Iu+ Vitamin A, Retinol Acetate 3000 IU + Cholecalciferol (Vitamin D3) 1500 IU + α-tocopherol acetate (Vitamin E) 2.0 mg + Thiamine Hydrochloride (Vitamin B1) Ph. Eur. 2.0mg + Riboflavin (Vitamin B2) Ph. Eur. 4 mg + Pyridoxine HCl Ph. Eur. 2 mg + Cyanocobalamin Ph. Eur. 10µg + Ascorbic acid Ph. Eur. 20 mg + Ca-pantothenate USP 10mg + Menadione Sodium Bisulphite 2.0mg + Nicotinamide Ph.Eur. 20mg + Inositol	Aliseryl WS is a highly effective combination of broad-spectrum antibiotics and vitamins. The product stimulates egg production, increases growth, improves feed conversion and is used as vitamin supplement during periods of diseases and stress. It is effective against gastrointestinal, respiratory and urinary infections caused by colistin, oxytetracycline, erythromycin and streptomycin sensitive micro-organisms, like Bordetella, Campylobacter, Chlamydia, E. coli, Haemophilus, Klebsiella, Mycoplasma, Pasteurella, Rickettsia, Salmonella, Staphylococcus and Streptococcus spp. in calves, goats, sheep, poultry	**Contra-indications:** Hypersensitivity to tetracyclines, macrolides, colistin or aminoglycocides. Administration to animals with a seriously impaired renal and/or hepatic function. Concurrent administration of penicillins, cephalosporins, quinolones and cycloserine. Administration to animals with an active microbial digestion. **Side Effects:** No undesirable effects are to be expected when the prescribed dosage regimen is followed.	The Netherland	EUDRALEX Volume 4 and GMP+B1	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

		1.0mg/gm Antibiotic	and swine.					
16.	Laboratorios Calier, S.A., Spain **Local Agent:** NEXUS Distriutor Mymensingh	YODIMASPEN Powder and solvent for suspension for injection (Vet) (15 ml Vial) Penethamate hydriodide 5.0gm/Vial (Equivalent to 5.000.000 UI of benzylpenicillin) Antibiotic	Cattle: Treatment of mastitis caused by *Streptococcus uberis*, *Streptococcus dysgalactiae*, *Streptococcus agalactiae* and *Staphylococcus aureus* (non-betalactamase-producing bacteria) sensible to benzylpenicillin.	Do not administer to animals presenting a hypersensitivity record to penicillin, cephalosporins and/or some of the excipients.	SPAIN		*Abjgv`b Kiv †h‡Z cv‡i |*	*Abjgv`b Kiv nj |*
17.	**Manufacturer:** M/S Shinil Bio-gen Co. Ltd., **Old Address:** 254-10,Dugok-rl, Shinam-myeon,Yeasan-gum, Choongchungnam-do, Korea. New Address: 235-18,Chusa-ro, Choongchungnam-do, Korea. **Importer**: M/S Pharma & Firm 3/2, City Heart Building, 67, Naya Paltan, Dhaka-1000.	**Cipryl Power solution** Ciprofloxacin 100g + Bromhexine Hydrochloride 10g/Litre Antibiotic	For the treatment of diease caused by parhogens (e.g., E. coli, Salmonella spp., Haemophilus paragallinarum, Mycoplasma spp.) susceptible to Ciprofloxacin. -Chicken and Duck: Colibacillosis, Enteritis, Salmonellosis, Chronic respiratory disease(CRD), Complicated chronic respiratory diesase(CCRD) and Infectious coryza.	**Contraindication** -None **Side-effect** **-**GI stress(Vomiting, Anorexia) -Headache, Dizziness, Sweating and Skin rashes. **Withdraw Period:10 days**	Korea	New	*Abjgv`b Kiv †h‡Z cv‡i |*	*Abjgv`b Kiv nj |*
18.	**Manufacturer:** M/S Shinil Bio-gen Co. Ltd., **Old Address:**	**Hytil sol** Each L Contains:	For the Treatment of the bacterial diseases caused by the below pathogens susceptible to Tilmicosin	**Contraindication** -None **Side-effect**	Korea	New	*Abjgv`b Kiv †h‡Z cv‡i |*	*Abjgv`b Kiv nj |*

	254-10,Dugok-rl, Shinam-myeon,Yeasan-gum, Choongchungnam-do, Korea. New Address: 235-18,Chusa-ro, Choongchungnam-do, Korea. **Importer**: M/S Pharma & Firm 3/2, City Heart Building, 67, Naya Paltan, Dhaka-1000.	Tilmicosin Phosphate 250g(Potency) Antibiotic	Chicken: Mycoplasmosis (Mycoplasma gallisepticum and Mycoplassma synoviae) Swine: Pasteurellosis (Pasteurella multocida), Pleuropneumonia (Actinobacillus pleuropneumoniae), Mycoplasmal pneumonia (Mycoplasma hyopneumoniae).	- Trim loss - Edema **Withdrawal Period**-10 days				
19.	**Manufacturer:** M/S Shinil Bio-gen Co. Ltd., **Old Address:** 254-10,Dugok-rl, Shinam-myeon,Yeasan-gum, Choongchungnam-do, Korea. New Address: 235-18,Chusa-ro, Choongchungnam-do, Korea. **Importer**: M/S Pharma & Firm 3/2, City Heart Building, 67, Naya Paltan, Dhaka-1000.	**FTD injection** **Each mL Contains,** Florfenicol.....50 mg Tylosin Tartrate….25 mg (Activity) Dexamethasone Disodium Phosphate……..330µg Antibiotic	For the treatment of diseases caused by pathogens sensitive to Florfenico and Tylosin. **-Swine** : Respiratory disease like pneumonia and bronchitis caused by Actinobacillus pleuropneumonia, Pasteurella multocida, Mycoolasma hyopneumonia, Haemophilus parasuis. Bacterial enteritis caused by Salmonella spp., Eacherchia coli. -**Bovine and Ovine**: Respiratory disease like pneumonia, Bronchitis and Bacterial enteritis caused by Salmonella spp., Escheria Coli.	**Contraindication:** None **Side-effect :** None **Withdrawal Period**- 28 days	Korea	New	*Abjgv` b Kiv †h‡Z cv¢i |*	*Abjgv` b Kiv nj |*
20.	**Manufacturer:** M/S Shinil Bio-gen Co. Ltd., **Old Address:** 254-10,Dugok-rl, Shinam-myeon,Yeasan-gum, Choongchungnam-do,	**AMIMOX inj**. **Each mL Contains,** Amoxicillin Trihydrate …….150 mg(Potency) Gentamycin Sulfate....40	**Calves** :For the treatment and prenvention of the pneumonia and diarrhea. **Piglets** : For the treatment and prenvention of the pneumonia, colibacillosis and diarrhea.	**Contraindication** **-None** **Side-effect** - **None** **Withdrawal Period**-	Korea	New	*Abjgv` b Kiv †h‡Z cv¢i |*	*Abjgv` b Kiv nj |*

	Korea. New Address: 235-18,Chusa-ro, Choongchungnam-do, Korea. **Importer**: M/S Pharma & Firm 3/2, City Heart Building, 67, Naya Paltan, Dhaka-1000.	mg(Potency) Antibiotic		Meat:15 days, Milk : 2 days				
21.	**Manufacturer:** M/S Shinil Bio-gen Co. Ltd., **Old Address:** 254-10,Dugok-rl, Shinam-myeon,Yeasan-gum, Choongchungnam-do, Korea. New Address: 235-18,Chusa-ro, Choongchungnam-do, Korea. **Importer**: M/S Pharma & Firm 3/2, City Heart Building, 67, Naya Paltan, Dhaka-1000.	**Pen-M injection** **Each mL Contains,** Procaine Penicillin G… ...200,000 I.U Dihydrostreptomycin Sulfate--250 mg Chlorpheniramine Maleate .....10mg Dxamethasone Disodium Phosphate ..... .0.5mg Antibiotic	For the treatment of the bacterial infections caused by pathogens susceptible to Penicillin and Streptomycin. - Bronchial Pneumonia, Cystitis, Swine Erysipelas, Mastitis and Endometritis. For the prevention of secondary infection by viral diseases.	**Contraindication** **-None** **Side-effect** - **None** **Withdrawal Period**- Cattle,Swin :30 days, Milk : 3 days	Korea	New	*Abyjgv` b Kiv †h‡Z cv‡i |*	*Abyjgv` b Kiv nj |*
22.	**Manufacturer:** M/S Shinil Bio-gen Co. Ltd., **Old Address:** 254-10,Dugok-rl, Shinam-myeon,Yeasan-gum, Choongchungnam-do, Korea. New Address: 235-18,Chusa-ro,	**Cefa4 Injection** **Each mL Contains,** Cefquinome Sulfate.... 25mg(Potency) Antibiotic	For the treatment of disease caused by pathogens to Cefquinome, -**Cattle, Calves** : Respiratory disease caused by Pasteurelle multocida and Pasteurella haemolytica. **- Swine** :pneumonia and respiratory disease caused by Pasteurelle multocida, Haemophilus parasuis,	**Contraindication** **-None** **Side-effect** - **None** **Withdrawal Period**- Cattle : 5 days, Swin : 3 days Milk : 12 hours	Korea	New	*Abyjgv` b Kiv †h‡Z cv‡i |*	*Abyjgv` b Kiv nj |*

	Choongchungnam-do, Korea. **Importer**: M/S Pharma & Firm 3/2, City Heart Building, 67, Naya Paltan, Dhaka-1000.		Actibobacillus pleuropnemonia and Streptococcus. MMA (Mastitis-Metritis-Agalactia syndrome) caused by E.coli, Staphylococcus spp. and Streptococcus spp.					
23.	**Manufacturer:** M/S Komipharma International CO. LTD. 1236-6, Chongwang- Dong, Shihung-SI, Kyonggi-DO, The Republic of Korea. Address: 593, Seonggok-ri, Gyeolseong-Myeon, Hongseong-Gun, Chungcheongnam-Do, Korea **Importer**: M/S. Rafique Medicine 55/B, Hazi zahir Bhaban, Sobahanbag, Saver, Dhaka.	**Tydohexin Powder** Doxycycline hyclate 200 gm + Tylosin Tartrate 100 gm + Bromhexine HCI (as Bromhexine) 20 gm/1 Kg Antibiotic	For the prevention and treatment of infectious disease, especially digestive and respiratory disease, causative of pathogens susceptible to Tylosin and doxycycline.	**Contra-indication:** i) Those with fever or serious nutrional disorder. ii) Those with infectious disease, parasitic infection, or stress. iii) Those with weakened immunity due to mold bacterial toxin (aflatoxin or toxin produced by E. coli or Salmonella spp.). iv) Those with hypersensitiveness to this kind of vaccine. **Side-effect:** Those administered with this produced may show hypersensitive reaction i.e. anorexia, vomiting, skin rash, despondency, convulsions and others. As this case requires to massage or administer epinephrine, according to the veterinarian's instruction.	Korea	New	*Abyjgv` b Kiv †h‡Z cv‡i |*	*Abyjgv` b Kiv nj |*
24.	**Manufacturer:** M/S Han Dong Co. Ltd., Address: Han Dong BLDG, 535,Ogeum-ro, songpa-gu, Seoul, Korea.	**Sentilo Sol. 200** Tilmicosin Phosphate 200gm/ liter Antibiotic	For treatment of bacterial diseases susceptive to tilmicosin - Pig : Pasteurella multocide, Actinobacillus pleuropneumoniae and Mycoplasma hyopneumoniae - Chicken : Mycoplasma	**Contraindication** - Animals who have shown shock or hypersensitivity to macrolide - Do not use lincosamide and macrolide **Side-effect**	Korea	New	*Abyjgv` b Kiv †h‡Z cv‡i |*	*Abyjgv` b Kiv nj |*

	Importer: M/S AVON POULTRY House 123 (Ground floor) Road 7 Block-B Section 12 Pallabi Mirpur Dhaka-1223.		hyopneumoniae and Mycoplasma synoviae	- None to our present knowledge. **Withdraw Period: Swine 7 days, Chicken 10 days**				
25.	**Manufacturer:** M/S Han Dong Co. Ltd., Address: Han Dong BLDG, 535,Ogeum-ro, songpa-gu, Seoul, Korea. **Importer**: M/S AVON POULTRY House 123 (Ground floor) Road 7 Block-B Section 12 Pallabi Mirpur Dhaka-1223.	**Ciprocin 200 Sol.** Ciprofloxacin 200g/Liter Antibiotic	For treatment of diseases caused by pathogenic bacteria - Mycoplasma - E. coil - Salmonella - Staphylococcos for chicken - Chronic respiratory disease (CRD) - Complicated Chronic respiratory disease (CCRD) - Salmonellosis (Pullorum disease & Fowl typhoid) - Bacterial diarrhea - Intecfious coryza - Staphylococcosis - Colibacillosis	**Contraindication:** It can remain in livestock products when overused or abused. Do not use this product mixed with tetrcycline, chloramphenicol or erythromycin **Side-effect:** None to our present knowledge. **Withdraw Period:** Broiler – Do not use this product for 10 days before a market day. Do not use in laying hens.	Korea	New	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
26.	**Manufacturer:** M/S Shinil Bio-gen Co. Ltd., **Old Address:** 254-10,Dugok-rl, Shinam-myeon,Yeasan-gum, Choongchungnam-do, Korea. New Address: 235-18,Chusa-ro, Choongchungnam-do, Korea. **Importer**: M/S Pharma & Firm	**Phenbrozole sol** **Each L Contains,** Phenylbutazone …….....50g Bromhexine Hydrochloride.....10g Anti-Inflammatory	For the adjuvant therapy of respiratory and infectious disease like Rhinitis, Laryngotracheitis, Bronchitis, Bronchopneumonia, Pheumonia in swine and chicken.	**Contraindication** -None **Side-effect** -None **Withdrawal Period**- Swine 7 days, Chicken-5 days	Korea	New	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

	3/2, City Heart Building, 67, Naya Paltan, Dhaka-1000.							
27.	Laboratorios Calier, S.A., Spain **Local Agent:** NEXUS Distriutor Mymensingh	Indigest Solution for injection. 30 ml Vial Menbutone 100 mg/ml Choleretic agent	Normaliser of the gastric, duodenal and biliary function. It is indicated in those cases when a stimulation of the digestive secretions is required: **Bovine**: Indigestion, toxaemia, ketosis, anorexia, hepatic and pancreatic insufficiencies. **Ovine and caprine**: Indigestion, toxaemia (including the gestation one), poisoning, hepatic and pancreatic insufficiencies. **Porcine**: Indigestion, anorexia, poisoning, hepatic and pancreatic insufficiencies. **Equine**: Toxaemia, anorexia, colic hepatic and pancreatic insufficiencies. **Dogs**: Indigestion, toxaemia, anorexia, constipation, hepatic and pancreatic insufficiencies.	Do not use in case of hypersensitivity to the active substance and/or any of the excipients. Do not administer to animals presenting cardiac disorders, hyperthermia or obstruction of the bile duct. Do not ever administer to cats.	Spain	New	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
28.	MEVET S.A.U, Spain **Local Agent:** Pharmachem Services, 48 Dilkhusha C/A, Motijheel, Dhaka	SANIVALL Disinfectant Solution Didecyl Dimethyl Ammonium Chloride 10% + Glutaraldehyde 4% + Formaldehyde 3.15% + Glyoxal 3.2% Disinfect	SANIVALL is highly effective against bacteria, viruses and fungi. It is broad spectrum against the organisms. It is highly effective against swine fever virus.	**Contraindication:** SANIVALL must not be ingested by animals or human being. Avoid contact with skin. **Side effects**: As other disinfectants, SANIVALL may cause irritation to skin, eyes and mucosa if not used with the recommended protection (mask, gloves, goggles and proper clothing).	Spain	New	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
29.	Green Cross Veterinary Products Co. Ltd., Korea (ACI Ltd.)	Green Cop Water Soluble Powder 100gm, 250gm, 500gm, 1kg & 5kg Potassium monopersulfate (Triple salt) 500gm+Malic acid	Disinfection for pathogen which is sensitive to this product like salmonella typhimurium, Brucella ovis, Avian influenza, Newcastle disease virus, Hog cholera virus,	**Contraindication**: Mix with any other disinfectant or insecticide. **Side effect**: Unknown.	korea	New	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

		100gm+Sodium chloride 15gm+Sulphamic acid 50gm+Sodium hexametaphosphate 181gm+Sodium dodecyl benzene sulphonate 150gm/1kg Disinfectant	Foot and Mouth disease virus. Also disinfection of animal shed and instruments.					
30.	Green Cross Veterinary Products Co. Ltd., Korea (ACI Ltd.)	Cidekill Liquid 1L, 5L, 10L, 18L & 50L Gluteraldehyde 150gm+Dimethylcocobenzyl ammonium chloride 100gm/1L Disinfectant	Disinfection for various virus, bacteria and fungi. Routine disinfection of animal shed, instruments, site, kennel, veterinary hospital etc.	**Contraindication**: Harmful if swallowed. Avoid inhalation of spray mist and contact with eyes and skin. **Side effect**: Skin reactions due to hypersensitivity or direct irritant effect.	KOREA	New	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
31.	Kilco (International) Limited. UK **Importer**: ACE Pharmaceuticals	Kleanline Clear, Colourless liquid. 25KG Paracetic Acid 5%+ Acetic Acid 10% + Hydrozen Peroxide 20% in w/w Disinfectant	Surface and Equipment Disinfectant Concentrate.	**Contraindication**: Do not breathe dust/fumes / gas/mist/vapours spray. **Side Effect:** Harmful by inhalation, in contact with skin and if swallowed. Cause severs burn. Contact with combustible material may cause fire.	United Kingdom	New	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
32.	Kilco (International) Limited. UK **Importer**: ACE Pharmaceuticals	Viroshield Liquid 1L, 5L & 25L Glutaraldehyde 15% + Benzalkonium Chloride 10% in w/w Disinfectant	Surface / Equipment Disinfection, Foot dip use, Wheel dip use.	**Contraindication**: Do not breathe fumes. **Side Effect**: Non Corrosive	United Kingdom	New	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
33.	**Manufacturer:** M/S Shinil Bio-gen Co. Ltd., **Old Address:** 254-10,Dugok-rl, Shinam-myeon,Yeasan-gum, Choongchungnam-do, Korea. New Address: 235-18,Chusa-ro, Choongchungnam-do, Korea.	Super Glu Glutaraldehyde 100g/1000ml Disinfectant	(1) For the sterilization and disinfection of pathogens susceptible to glutaraldehyde. - Bacteria : Salmonella typhimurium. - Virus : Avain influenza virus, Newcastle disease virus, Classical swine fever virus. (2) For the disinfection of medical instruments that can't be treated by heat. (3) For the disinfection of	**Contraindication:** None **Side-effect:** Nausea,Headache -airway obstruction ,eye irritation and dermatitis **Withdraw Period: N/A**	Korea	New	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

	Importer: M/S Pharma & Firm 3/2, City Heart Building, 67, Naya Paltan, Dhaka-1000.		livestock pens, hatchery, equipment, instrument and environment of livestock.					
34.	**Manufacturer:** M/S Komipharma International CO. LTD. 1236-6, Chongwang- Dong, Shihung-SI, Kyonggi-DO, The Republic of Korea. Address: 593, Seonggok-ri, Gyeolseong-Myeon, Hongseong-Gun, Chungcheongnam-Do, Korea **Importer**: M/S. Rafique Medicine 55/B, Hazi zahir Bhaban, Sobahanbag, Saver, Dhaka.	**Multiside Spray** Alkyl-dimethyl Benzyl Ammonium Chloride 50%---170.6 gm + Didecyldimethyl Ammonium Chloride 78.0gm + Glutaraldehyde-107.25gm/1 Litre Disinfectant	Biocidal and Sterillization effect of etiologic agents, which are sensitives to this product. **Bacteria**: General bacteria (salmonella typhimurium),) Brucella ovis ,Haemophilus parasol, Pasturella multocida, Actinobacillus pleuropneumonlae, streptococcus suls, Bordetella bronchiseptical, mycoplasma hyopneumoniae Virus: Avian influenza virus (AIV), Porcine reproductive and respiratory Syndrom virus (PRRSV), porcine Rota Virus,(PRV), foot and mouth disease virus (FMDV), classical swine fever virus (CSFV),Aujeshzkys disease virus (ADV) porcine epidemic diarrhea virus (PEDV), porcine circovirus type 2 (PCV2),Newcastle disease virus (NDV).Transmissible gastroenteritis virus (TGEV), Porcine parvo virus (PPV), Bovine rota virus (BRV),Bovine corona virus (BCV) Infectious bursal disease virus(IBDV), swine Influenza virus (SIV). **Fungus:** Tricophyton mentagrophytes.Strelization effect on livestock building and equipment for livestock	Contraindication & Side effect: None	Korea	New	*Abyjgv` b Kiv †h‡Z cv‡i |*	*Abyjgv` b Kiv nj |*

35.	**Manufacturer:** M/S Komipharma International CO. LTD. 1236-6, Chongwang- Dong, Shihung-SI, Kyonggi-DO, The Republic of Korea. Address: 593, Seonggok-ri, Gyeolseong-Myeon, Hongseong-Gun, Chungcheongnam-Do, Korea **Importer**: M/S. Rafique Medicine 55/B, Hazi zahir Bhaban, Sobahanbag, Saver, Dhaka.	**Farm Guard Solution** \N-Alkyl Dimethyl Benzyl Ammonium Chloride, KVP 100 gm + Citric acid, KVP 200 gm + Phosphoric acid, KVP 100 gm/1 Litre Disinfectant	Eradication of pathogens sensitive to this product:- **1. Bacteria:- i)** Salmonella typhimurium.**(ii)** Brucella ovis, **iii)** Mycobacterium fortutum. **Iv)** Clostridium Perfringens. **2. Virus:-i)** Avian influenza virus (AIV) **ii)** Avian Newcastle disease virus (NDV) **iii)** Classical Swine fever Virus (CSFV).**iv)** Porcine Reproductive and Respiratory Syndrome (PRRSV).**v)** Foot-and-mouth Disease Virus (FMDV), Sterilization effect on livestock building and equipment for livestock.	Contraindication & Side effect: None	Korea	New	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
36.	Green Cross Veterinary Products Co. Ltd., Korea (ACI Ltd.)	Nurelle-20 Liquid 100ml, 500ml, 1L, 5L & 10L Cypermethrin 200gm/1L Ectoparasiticide	For control of animal ectoparasites.	**Contraindication**: Do not contaminate ponds, waterways or ditches with the product or used container. **Side Effect**: Unknown.	KOREA	New	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
37.	Merial, France **Local Agent:** Advance Animal Science Co. Ltd. 2/10 Block-B, Lalmatia, Dhaka-1207)	Frontline Plus Dogs spot on solution FIPRONIL In House 134mg + (S)-methoprene In House 120.60mg/1.34mL Insecticide	*FRONTLINE® PLUS FOR DOGS* provides fast, effective and convenient treatment and control of fleas, ticks and chewing lice for dogs and puppies. - Stops and prevents infestations - Kills adult fleas, flea eggs, and flea larvae - Prevents all flea stages (eggs, larvae, pupae) from developing - Kills fleas which may cause flea allergy dermatitis - Kills all stages of deer ticks (which may carry Lyme disease), brown dog ticks, American dog ticks, and lone star ticks - Prevents and controls	**Contraindication:** None **Side effects**: No remarkable toxicity or side effects were observed during clinical trial.		New	c‡qvRbxq †id v‡iÝmn cieZx mfvq Dc¯vc‡bi Rb¨ ejv †h‡Z cv‡i	c‡qvRbxq †id v‡iÝmn cieZx mfvq Dc¯vc‡bi Rb¨ ejv হল

			reinfestations - Rapidly eliminates infestations with chewing lice - Aids in control of sarcoptic mange infestations - Can also be used on breeding, pregnant and lactating bitches					
38.	**Manufacturer:** M/S Shinil Bio-gen Co. Ltd., **Old Address:** 254-10,Dugok-rl, Shinam-myeon,Yeasan-gum, Choongchungnam-do, Korea. New Address: 235-18,Chusa-ro, Choongchungnam-do, Korea. **Importer**: M/S Pharma & Firm 3/2, City Heart Building, 67, Naya Paltan, Dhaka-1000.	**Super CMP Injection** Calcium Gluconate 230 mg + Magnesium Chloride Hexahydrate 45mg + Sodium Hypophosphite 34.3 mg + Potassium Chloride 2 mg + Dextrose 250mg/ml Mineral	For the prevention and treatment of Hypocalcemia (Parturient paresis, Milk fever), Hypomagnesemia (Grass tetany), and other conditions associated with calcium, magnesium, phosphorous and glucose deficiencies in cattle, sheep and swine.	**Contraindication: None** **Side-effect: None** **Withdraw Period: N/A**	Korea	New	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
39.	Agri Laboratories Ltd., USA (ACI Ltd.)	Cal-Dex CMPK Injection Sterile Aqueous Solution 250ml & 500ml Calcium Gluconate monohydrate (23.3%) 10.8g + Potassium (as potassium chloride) 8.0g + Phosphorus (as sodium hypophosphite H20) 2.5g + Magnesium (as magnesium borogluconate) 1.6g/500ml Minerals	Recommended for use in pre and post-parturitional hypocalcemia, milk fever, Tetany of pregnancy and lactating, Paresis, calcium deficiency syndrome in early pregnancy of cows, sheep, goat, osteomalasia in cow, sheep and goat, disease and weakness of new born animal.	**Contraindication:** Do not administer this product to animals showing signs of cardiac distress. **Side Effect:** Perivascular or subcutaneous deposition may result in severe inflammation at the injection site.	USA	New	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
40.	Ceva Phylaxia Veterinary Biologicals Co., Ltd. Hungary (ACI Ltd.)	Cevac Mass L Vaccine 1000, 2500 & 5000 doses Avian Infectious bronchitis virus	Live, Freeze-dried vaccine for the active immunization of chicken against avian infectious bronchitis virus.	**Contraindication:** No contraindications are known. **Side Effect:** Mild respiratory signs may rarely occur which disappears within a	Hungary	New	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

		(IBV), Massachusetts type B-48……min. 2.8 log10 EID50/dose Vaccine		few days.				
41.	Ceva Salud Animal, S.A De C.V (ACI Ltd.)	Cevac Flukem Vaccine 500ml/1000 doses Avian Influenza virus type A, subtype H5N2 A/Chicken/ Mexico/232/94/CPA strain Vaccine	Cevac Flukem is indicated for immunization against Avian Influenza in susceptible chickens.	**Contraindication**: No contraindications are known. **Side Effect:** It may cause some lesions at the site of injection.	Mexico	New	*c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i|*	*c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv* হল|
42.	Ceva Salud Animal, S.A De C.V (ACI Ltd.)	Cevac New Flukem Vaccine 500ml/1000 doses Avian Influenza virus type A, subtype H5N2 A/Chicken/ Mexico/232/94/CPA strain+Newcastle Disease virus, La Sota Strain Vaccine	Cevac New Flukem is indicated for immunization against Newcastle disease and Avian Influenza in susceptible chickens.	**Contraindication**: No contraindications are known. **Side Effect:** It may cause some lesions at the site of injection.	Mexico	New	*c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i|*	*c‡qvRb ‡bB weavq Av‡e`b bv gÄyi Kiv* হল|
43.	Green Cross Veterinary Products Co. Ltd., Korea (ACI Ltd.)	BBNE Oil Vaccine Vaccine 250, 500 & 1000 doses Inactivated Newcastle disease Virus (B1 strain)….. over $10^{9.5}EID_{50}$+ Infectious Bronchitis virus (M41 strain)…..over $10^{7.0}EID_{50}$+ Infectious bronchitis virus (KM91 strain) ..over $10^{7.0}EID_{50}$/Egg drop syndrome virus (GCVP 813 strain)..over $10^{8.0}EID_{50}$+Oil adjuvent, ISA70-70%//dose Vaccine	The vaccine is a poly-valent oil vaccine for simultaneous immunization against Newcastle disease, Infectious bronchitis (prevailing and variant type) and Egg Drop Syndrome with one vaccine.	**Contraindication**: A satisfactory immune effect may not be obtained from diseased chickens. Only the healthy chickens should be inoculated. **Side Effect:** For forced molting hens, they may suffer stress from inoculation with an oil based adjuvant.	korea	New	*Aby‡gv`b Kiv †h‡Z cv‡i|*	*Aby‡gv`b Kiv nj|*
44.	Green Cross Veterinary Products Co. Ltd., Korea (ACI Ltd.)	BBN Oil Vaccine Vaccine 250, 500 & 1000 doses Inactivated Newcastle disease Virus (LaSota strain) …over $10^{9.5}EID_{50}$+Infectious Bronchitis virus	The vaccine is a poly-valent oil vaccine for simultaneous immunization against Newcastle disease and Infectious bronchitis (Mass and variant type) with one vaccine.	**Contraindication**: A satisfactory immune effect may not be obtained from diseased chickens. Only the healthy chickens should be inoculated. **Side Effect:** For forced molting hens, they may suffer stress from inoculation with an oil based adjuvant.	korea	New	*Aby‡gv`b Kiv †h‡Z cv‡i|*	*Aby‡gv`b Kiv nj|*

		(M41 strain)..over $10^{7.0}EID_{50}$+ Infectious bronchitis virus (KM91 strain)..over $10^{7.0}EID_{50}$ +Oil adjuvent, ISA70-70%//dose Vaccine						
45.	Laboratorios Hipra S.A., Spain (**Importer:** Nasco Agro Product, 307 Sk Mujib Road, Agrabad, Chittagong)	Avisan Multi Vaccine 1000 dose/bottle Inactivated Avian Infectious bronchitis virus, Strain H52……. HAI^1 :2^6-2^8 Inactivated Newcastle disease virus, La Sota strain…… HAI^2 :2^4-2^6 Inactivated Egg drop syndrome virus, Adenovirus 127 strain... HAI^1 :2^7-2^9 /dose(0.5ml)	To prevent Avian infectious bronchitis, Newcastle disease and Egg drop syndrome	**Contraindications**: No contraindications are known. **Side effects:** No palpable reactions were observed following the injection of one dose of vaccine.	Spain	New	*Abyjgv`b Kiv †h‡Z cv¢i |*	*Abyjgv`b Kiv nj |*
46.	Laboratorios Hipra S.A., Spain (**Importer:** Nasco Agro Product, 307 Sk Mujib Road, Agrabad, Chittagong)	STARTVAC 1 dose/bottle 5 dose/bottle 25 dose/bottle 125 dose/bottle *Escherichia coli J5 inactivated > 50 RED_{60} (Rabbit effective dose in 60% of the animals (serology))* *Staphylococcus aureus (CP8) SP 140 strain inactivated, expressing Slime Associated Antigenic Complex (SAAC) > 50 RED_{80} (Rabbit effective dose in 80% of the animals (serology)*	To prevent Mastitis in cows and heifers	**Contraindications**: None **Side effects:** None	Spain	New	*Abyjgv`b Kiv †h‡Z cv¢i |*	*Abyjgv`b Kiv nj |*
47.	Lohmann Animal Health International, USA **(Importer:** C. P. Bangladesh Co., Ltd.; House-28, Alaol Avenue, Sector-6, Uttara, Dhaka-1230**)**	**AviPro MG F Vaccine** A vaccine containing live *Mycoplasma gallisepticum*, F strain, presented in lyophilized (freeze-dried) form in a 30 ml glass vial	This vaccine is recommended as an aid in the prevention of clinical signs of Mycoplasma gallisepticum (MG) infection in healthy chickens. This vaccine is recommended for drinking water use at 12 weeks of age or older but not later than 4 weeks prior to the onset of egg production.	**Contra-indications:** It should not be administered within 1 week before or after vaccination with live Newcastle, bronchitis, or laryngotracheitis vaccines, or within 3 days before to 7 days after treatment with oxytetracycline or chlortetracycline. **Side-effect:** Vaccination may induce mild respiratory signs which can persist for a few days. Overdose = Mild respiratory	USDA	New	*Abyjgv`b Kiv †h‡Z cv¢i |*	*Abyjgv`b Kiv nj |*

				reaction can be observed for few days after vaccination.				
48.	**Manufacturer:** M/s Choong Ang Vaccine Laboratories Co.Ltd., 1476-37 Yuseong-daero, Yuseong-gu, Daejeon, 305-348, Republic of Korea. **Importer :** M/S Pharma & Firm 3/2 Cityheart Bulding, 67 Nayapaltan, Dhaka.	PoulShot® IBD Win+ Infectious bursal disease virus (IBDV, Winterfield 2512 strain) $\geq 10^{2.0}EID_{50}$ Stabilizer……..……25%	As an aid in the control and prevention of infectious bursal disease (Gumboro disease) caused by IBDV in chickens.	**Contraindication:** -None **Side-effect:** Vaccination reaction such as loss of appetite, coughing . **Withdraw Period: Zero(0) day** **Warning specific for target animal**:-Do not vaccinated previously vaccinated animals that showed bad reaction or shock to this vaccine. --Immunity formation in chickens may differ depending on the level of material antibody ,age of animal, health condition or number of vaccination . **Warning specific for user :** **-**Vaccine should avoid direct skin contract or inhalation.	Korea	New	*Abjgv`b Kiv †h‡Z cv‡i |*	*Abjgv`b Kiv nj |*
49.	**Manufacturer:** M/s Choong Ang Vaccine Laboratories Co. Ltd., 1476-37 Yuseong-daero, Yuseong-gu, Daejeon, 305-348, Republic of Korea. **Importer**: M/S Pharma & Firm 3/2, City Heart Building, 67, Naya Paltan, Dhaka-1000.	**CaniShot® RV-K** Each dose: Canine Rabies virus (Pasteur Strain)----------$\geq 10^{7.0}$ $FAID_{50}$ Adjuvant…….....5% Inactivator.....≤1.0mM	For active immunization of dogs, cats and cattle as an aid in the control and prevention of rabies caused by rabies virus.	**Contraindication:** Vaccination in pregnant and lactating animals is not recommended.	Korea	New	*Abjgv`b Kiv †h‡Z cv‡i |*	*Abjgv`b Kiv nj |*
50.	**Manufacturer:** BoehringerIngelheimVetmedica, S.A. de C.V. (Mexico) Calle 30 No. 2614, Zona Industrial Guadalajara, Jal., 44940, Mecxico **Local Agent:** M/s. Square Pharmaceuticals Ltd. (Agrovet Division), 50, C/A Mohakhali Dhaka	Volvac® IBD MLV (1000 doses in a 15ml vial with 30ml diluent) Gumboro disease virus (strain: D78, Lukerts) Each dose contains not less than 10^3 $TCID_{50}$ of vaccinal virus	For the immunization of healthy broilers and replacement breeder and layer pullets against infectious Bursal Disease (IBD)	**Warnings and Contra-indication:** Do not use unhealthy Birds. Sick or weak birds will nt develop adequet immuunit following vaccination. **Side effect:** None	Mexico	New	*Abjgv`b Kiv †h‡Z cv‡i |*	*Abjgv`b Kiv nj |*
51.	**Manufacturer:**	Volvac® IBD MLV	For the immunization of healthy	**Warnings and Contra-indication:**	Mexico	New	*Abjgv`b Kiv †h‡Z cv‡i |*	*Abjgv`b Kiv nj |*

	BoehringerIngelheimVetmedica, S.A. de C.V. (Mexico) Calle 30 No. 2614, Zona Industrial Guadalajara, Jal., 44940, Mecxico **Local Agent:** M/s. Square Pharmaceuticals Ltd. (Agrovet Division), 50, C/A Mohakhali Dhaka	(5000 doses in a 15ml vial with 30ml diluent) Gumboro disease virus (strain: D78, Lukerts) Each dose contains not less than 10^3 $TCID_{50}$ of vaccinal virus	broilers and replacement breeder and layer pullets against infectious Bursal Disease (IBD)	Should not be administered in sick or weak birds, because sick birds will not develop adequate immunity. **Side effect:** None				
52.	**Manufacturer:** BoehringerIngelheimVetmedica, S.A. de C.V. (Mexico) Calle 30 No. 2614, Zona Industrial Guadalajara, Jal., 44940, Mecxico **Local Agent:** M/s. Square Pharmaceuticals Ltd. (Agrovet Division), 50, C/A Mohakhali Dhaka	Volvac® ND conc. KV (2500 doses in a 500ml bottle) Newcastle disease Virus (ND, Strain la sota) Each dose containsnot less than $10^{8.6}EID_{50}$ of vaccinal virus.	The vaccine is recommended for the vaccination of healthy birds since one-day-old, against Newcastle Disease.	**Warnings and Contra-indication:** Should not be administered in sick or weak birds, because sick birds will not develop adequate immunity. **Side effect:** None	Mexico	New	*Abyjgv` b Kiv †h‡Z cv‡i |*	*Abyjgv` b Kiv nj |*
53.	**Manufacturer:** BoehringerIngelheimVetmedica, S.A. de C.V. (Mexico)	Volvac® ND+IB+EDS KV (1000 doses in a 500ml bottle)	The vaccine is recommended as an aid in the prevention of Newcastle Disease, Infectious Bronchitis Disease, and to prevent egg drop	**Warnings and Contra-indication:** Should not be administered in sick or weak birds, because sick birds will not develop adequate immunity.	Mexico	New	*Abyjgv` b Kiv †h‡Z cv‡i |*	*Abyjgv` b Kiv nj |*

	Calle 30 No. 2614, Zona Industrial Guadalajara, Jal., 44940, Mecxico **Local Agent:** M/s. Square Pharmaceuticals Ltd. (Agrovet Division), 50, C/A Mohakhali Dhaka	Newcastle disease Virus (ND, Strain la sota) , avain infectious bronchitis (IB, Strain mass 41), egg drop syndrome (EDS-76, Strain 127) viruses Each dose containsnot less than $10^{8.2}EID_{50}$ Newcastle Disease Virus + $10^{6.7}EID_{50}$Infectious Bronchitis Virus + 1000 HAU Egg Drop Syndrome virus	caused by of EGG Drop Syndrome virus.	**Side effect:** None				
54.	**Manufacturer:** M/S Komipharma International CO. LTD. 1236-6, Chongwang- Dong, Shihung-SI, Kyonggi-DO, The Republic of Korea. Address: 593, Seonggok-ri, Gyeolseong-Myeon, Hongseong-Gun, Chungcheongnam-Do, Korea **Importer**: M/S. Rafique Medicine 55/B, Hazi zahir Bhaban, Sobahanbag, Saver, Dhaka.	**Pro-vac(R) AB Vaccine** **(2 Ml Dose)** ANTHRAX (STERAIN)SULUTION SUSPENDEND IN 50% GLYCERIN---47.5%, + ANTHRAX (STERAIN) SPORE COUNT----≥0.8 x 10^7CFU, + ATTENUATED BLACK LEG CULTURE SOLUTION—47.5%,+ BLACK LEG SPORE COUNT----- ≥2.0x 10^6 CFU + TOLUENE----------4.995%,+ SAPONIN------0.005%,	For the prevention Of Anthrax Disease and black leg at the same time	Contraindication & Side effect: None	Korea	New	*Abjgv` b Kiv †h‡Z cv‡i |*	*Abjgv` b Kiv nj |*
55.	**Manufacturer:** M/S Komipharma International CO. LTD. 1236-6, Chongwang- Dong, Shihung-SI, Kyonggi-DO,	**Pro-vac(R) AE-FP Vaccine** Avian Encephalomyelitis virus (Calnek 1143 strain)---- Min $10^{2.8}$ EID_{50} +	For the prevention againaist Avian Encephalomyelitis (AE) and Fowl Pox (FP) at the same time	Contra-indiction : Do not administer this product to the following once. i) Those with fever or serious nutritional disorder. ii) Those with infectious disease, parasite	Korea	New	*Abjgv` b Kiv †h‡Z cv‡i |*	*Abjgv` b Kiv nj |*

	The Republic of Korea. Address: 593, Seonggok-ri, Gyeolseong-Myeon, Hongseong-Gun, Chungcheongnam-Do, Korea **Importer**: M/S. Rafique Medicine 55/B, Hazi zahir Bhaban, Sobahanbag, Saver, Dhaka.	Attenuated Fowl Pox virus, TCH---- Min $10^{2.8}$ EID_{50} + LPGG---50%		infection, or stress. iii) Those with weakened immunity due to mold or bacterial toxin (aflatoxin or toxin produced by Ecoli or salmonella spp.) iv) those with shock or hypersensitiveness to this vaccine				
56.	MERIAL Italia S.p.A, Italy (Advance Animal Science Co. Ltd. 2/10 Block-B, Lalmatia, Dhaka-1207)	GALLIMUNE 503 ND+IB+EDS+IC2 Vaccine Inactivated Newcastle disease virus, inactivated infectious bronchitis virus, egg drop syndrome virus (EDS '76), inactivated infectious coryza in chicken/0.3ml	Active immunization of laying stock against Newcastle disease, infectious bronchitis, egg drop syndrome (EDS '76) and infectious coryza.	**Contraindication:** None. **Side effects**: Moderate swelling may observed at the site of injection at 7days interval.	Italy	New	*Abyjgv` b Kiv †h‡Z cv¢i |*	*Abyjgv` b Kiv nj |*
57.	MERIAL NANJINJ Animal Health Co. Ltd., China (Advance Animal Science Co. Ltd. 2/10 Block-B, Lalmatia, Dhaka-1207)	Reassortant Avian Influenza Virus Vaccine Inactivated (H5N1 Subtype, Re-6 Strain Each dose contains Inactivated Avian Influenza Virus H5N1 subtype…….≥64HI.U	To prevent Avian Influenza in chicken, duck, goose induced by H5 subtype Avian Influenza Virus. Potency will be obtained 14days post vaccination and lasts for six months in chickens. A booster vaccination after 3 weeks of vaccination in duck and goose, potency last for 4 months.	**Contraindication:** None. **Side effects**: None	China	New	*Abyjgv` b Kiv †h‡Z cv¢i |*	*Abyjgv` b Kiv nj |*
58.	Merial, France **Local Agent:** Advance Animal Science Co. Ltd. 2/10 Block-B, Lalmatia, Dhaka-1207)	AVIENEW NEO Vaccine Virus Vivant De La Maladie De NEWCASTLE Souche VG/GA……≥5.5log $_{10}$DIO $_{50}$ + NEWCASTLE Disease Virus VG/GA Strain...≥5.5log $_{10}$DIO $_{50}$	Active immunization of chickens against Newcastle Disease	**Contraindication:** None. **Side effects**: No reactions could be detected so far.	France	New	*Abyjgv` b Kiv †h‡Z cv¢i |*	*Abyjgv` b Kiv nj |*

59.	Merial, France **Local Agent:** Advance Animal Science Co. Ltd. 2/10 Block-B, Lalmatia, Dhaka-1207)	Gallimune 208 ND+Flu H9 M.E. Vaccine Virus Inactive de la GRIPPE AVIAIRE Souche H9N2, Titer minimum Avant Inactivation……10^8DIO50 Virus Inactive de la NEWCASTLE Souche ULSTER 2C, Titer minimum Avant Inactivation……10^8DIO50 Inactivated Avian Influenza Virus H9N2 Strain, Minimum Titer before Inactivation……10^8EID50 Inactivated NEWCASTLE Virus ULSTER 2C Strain, Minimum Titer before Inactivation……10^8EID50	Preventive Vaccination of poultry against Avian Influenza (H9N2) and Newcastle Disease	**Contraindication:** None. **Side effects**: Moderate swelling may observed at the site of injection at 7days interval.	France	New	*c‡qvRb ‡bB wea‡q Av‡e`b bvgÄyi Kiv †h‡Z cv‡i |*	*c‡qvRb ‡bB wea‡q Av‡e`b bvgÄyi Kiv nj |*
60.	**Manufacturer:** BoehringerIngelheimVetmedica, S.A. de C.V. (Mexico) Calle 30 No. 2614, Zona Industrial Guadalajara, Jal., 44940, Mecxico **Local Agent:** M/s. Square Pharmaceuticals Ltd. (Agrovet Division), 50, C/A Mohakhali Dhaka	Avian Coryza, Newcastle, Infectious Bronchitis and Egg Drop Syndrome Killed virus vaccine. *Avibacterium paragallinarum* Srotpe A, Harvest fluid 10^8 EID_{50} + *Avibacterium paragallinarum* Srotpe B, Harvest fluid 10^8 EID_{50} + *Avibacterium paragallinarum* Srotpe C, Harvest fluid 10^8 EID_{50} + Newcastle Disease, harvest fluid $10^{8.2}$ EID_{50} + Infectious Bronchitis, harvest fluid $10^{6.7}$ EID_{50} + Egg Drop Syndrome, harvest fluid 1000 HAU + Mineral Oil USP 234.5mg + Span 80 USP 22.720mg +Tween 80 USP 8.262mg + Propylene Glycol USP 4.144mg + Formaldehyde Solution USP 0.432mg **Killed virus vaccine**	Recommended for the immunization of healthy birds as an aid in the prevention and control of Infectious Coryza, Newcastle Disease, Infectious Bronchitis and EGG Drop Syndrome. It is generally recommended for use in layers, with the vaccine usually being administered at between 14 an 17 weeks of age. It is also recommended that, where Infectious Coryza is a threat to production.	**Warnings and Contra-indication:** Should not be administered in sick or weak birds, because sick birds will not develop adequate immunity. **Side effect:** None	New	Volvac AC PLUS + ND + IB + EDS KV Boehringer Ingelheim	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
61.	**Ceva Phylaxia Veterinary Biologicals Co., Ltd. Hungary** **(ACI Ltd.)**	Vectormune ND Vaccine 1000, 2000 & 4000 doses *Live recombinant turkey herpesvirus with inserted NDV….at least 2500 PFU/dose*	Live, Frozen vector vaccine for active immunization of chickens against Newcastle and Marek's disease.	**Contraindication:** Do not use in birds in lay. **Side Effect:** No undesirable effects are known.	Hungary	New	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

62.	**Eagle Vet. Tech Co., Ltd. Korea (ACI Ltd.)**	Buscom Injectable solution 10ml, 20ml, 50ml, 100ml & 200ml *Scopolamine butylbromide 4.0mg+Dipyrone 500.0mg/ml*	Supplementary treatment of functional bloat, colic, esophageal obstruction, diarrhea, pain related to enteritis and the malfunction of digestive system and so on for cattle and calves.	**Contraindication**: Inject slowly and do not mix with other medicines. **Side Effect**: Rarely, hypersensitivity can occur.	korea	New	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
63.	**M/S. Quat-chem Ltd. United kingdom** **Gentry Pharmaceuticals Ltd., House: 08, Road: 28, Sector: 7, Uttara, Dhaka**	Viruquat 300 Solution *Benzalkonium Chloride 10% W/W + Glutaraldehyde 15-16%*	• Infection Control In Livestock Agriculture • Disinfection Of Livestock Housing & Farm Buildings • Ideal for Terminal Disinfection • Foot & Wheel Dips • Disinfection Of All Farmyard Equipment	Contraindication & Side effect: None	UK	Benzalkonium Chloride BP 5 ml + Glutaraldehyde 50% BP 8 ml / 100 ml	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
64.	**Merial, France** **Local Agent:** **Advance Animal Science Co. Ltd.** **2/10 Block-B, Lalmatia, Dhaka-1207)**	NexGard Chewable Tablet *AfoxolanNer 68mg*	For the treatment and control of flea infestations in dogs (Ctenocephalides felis and C. canis) for at least 5 weeks. The product can be used as part of a treatment strategy for the control of flea allergy dermatitis (FAD). Treatment of tick infestations in dogs (Dermacentor reticulates, Ixodes ricinus, Rhipicephalus sanguineus) One treatment kills ticks for up to one month. Fleas and Ticks must attach to the host and commence feeding in order to be exposed to the active substances.	**Contraindication:** Do not use in case of hypersensitivity to the active substances or to any of the excipients. **Side effects**: None	France	New	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

Annex-E

Jla wbqšY KwgwU Gi 245Zg mfvq †iwR‡÷kb ewZ‡ji Rb¨ mycvwikKZ.Jl‡ai ZvwjKv|

bs	c¨Zôv‡bi bvg		Jl‡ai eªvYR¨K bvg	‡RbwiK bvg	‡iwR‡÷kb bs (wWGAvi bs)	c‡UÝx	e¨vP bs	MÖnZ e¨e¯'v`	‡iwR‡÷kb mvgwqK ewZ‡ji ZwiL
01	‡gmvm Avi Gb dvgvmDwUK¨vjm wjt, C`MuU K¤úvDÛ, gqgbwmsn|	1.	wm‡fU-‡fU †evjvm	c¨vivwmUvgj weic 2 Mvg	172-21(V)-83	54.61% mxgv:(90%-110%)	8	gvbwnfy© nIqvq c`wUi ‡iwR‡÷kb mvgwqK ewZj Kiv nBqv‡Q|	wWG/GgGj-172/77/515; ZwiL: 17/01/2016
02	‡gmvm wdwb· ‡KwgK¨vj j¨ve‡iUixR wjt, biwms`x|	2.	wdwb· ‡iwbwUwWb U¨ve‡jU	‡iwbwUwWb BDGmwc-150wgtMvt	20-01-29	41.29% (95%-105%)	wcGBP-008	c`wUi ‡iwR‡÷kb mvgwqK ewZj Kiv nBqv‡Q|	wWwRwWG/GmGm -032/2011/659; ZwiL: 20/01/2016
03	‡gmvm ‡gwWKb dvgvmDwUK¨vjm wjt, wgicyi, XvKv	3.	wi‡evdwfb U¨ve‡jU	wi‡evdwfb weic 5 wgt Mvt	238-04-29	78.60% (95%-115%)	061114	gvbwnfy© nIqvq c`wUi ‡iwR‡÷kb mvgwqK ewZj Kiv nBqv‡Q|	wWwRwWG/GmGm -238/16/1040; ZwiL: 25/01/2016
04	‡gmvm we÷j dvgvwjt Gm-13,G-32 Ask, †Kvbvevox, MvRxcyi|	4.	‡K‡Uv‡ivjvK U¨ve‡jU	‡K‡Uv‡ivjvK U‡g_vgvBb BDGmwc	054-91-006	c‡hvR¨ bq	c‡hvR¨ bq	‡iwR‡÷k‡bi kZ f½ nIqvi Kvi‡b c`wUi ‡iwR‡÷kb mvgwqK ewZj Kiv nBqv‡Q|	wWG/GgGj-054/87/3724; ZwiL: 10/03/2016
		5.	‡jv‡mK 20 K¨vcmyj	IwgcvRj weic 20 wgt Mv	054-25-29	84.45% (95%-105%)	14 ‡R001	gvbwnfy© nIqvq c`wUi ‡iwR‡÷kb mvgwqK ewZj Kiv nBqv‡Q|	wWG/GmGm -054/03/3053; ZwiL: 2/02/2016
05	মেসার্স ক্রিস্টাল ফার্মাসিউটিক্যালস লিঃ, চর্থা, Kzgjv	6.	wbwWb U¨ve‡jU	†iwbwUwWb	182-021-055	59.73%	03 (m¨v¤új)	gvbwnfy© nIqvq c`wUi ‡iwR‡÷kb mvgwqK ewZj Kiv nBqv‡Q|	wWG/GgGj-182/03/3147; ZwiL: 19/02/2016
06	‡gmvm we÷j dvgvwjt Gm-13,G-32 Ask, †Kvbvevox MvRxcyi|	7.	‡jv‡mK 20 K¨vcmyj	IwgcvRj weic 20 wgt Mvt	054-25-29	84.45% (95%-105%)	14 ‡R 001	gvbwnfy© nIqvq c`wUi ‡iwR‡÷kb mvgwqK ewZj Kiv nBqv‡Q|	wWG/GmGm-054/03/3053; ZwiL: 28/02/2016
07	‡gmvm ÷¨vÛvW j¨ve‡iUixR wjt PÆMvg|	8.	‡fvë-50 U¨ve‡jU	WvB‡Kv‡dbvK †mvwWqvg BDGmwc 50wgtMvt	021-153-65	83.56% (90%- 110%)	010614	gvbwnfy© nIqvq c`wUi ‡iwR‡÷kb mvgwqK ewZj Kiv nBqv‡Q|	wWwRwWG/GmGm-180/15/1143; ZwiL: 26/01/2016

bs	cÖwZôv‡bi bvg		Jl‡ai eªvÛ/R¨K bvg	‡RbwiK bvg	‡iwR‡÷ªkb bs (wWGAvi bs)	c‡UÝx	e¨vP bs	Mߧ© e¨e¯'v`	‡iwR‡÷ªkb mvgwqK evwZ‡ji ZvwiL
08	‡gmvm ‡Mv‡e· dvgvmDwUK¨vjm wjt, Kvgvjcyi, bvivqbMÄ|	**9.**	wgD‡Kv‡UK wmivc	‡evg‡nw· wWb nvB‡Wv‡KvivBW wewc 0.080Mvt/100wgwj	180-71-031	52.25% (90%-110%)	01	gvbwbnfÝ nIqvq c`wUi ‡iwR‡÷ªkb mvgwqK evwZj Kiv nBqv‡Q|	wWwRwWG/GmGm-180/15/1029; ZvwiL: 24/01/2016
09	‡gmvm ¯úvK dvgvmDwUK¨vjm 175, Picvovev‡jb, Picvov gqgbwmsn|	**10.**	¯úvivg K¨vcmyj	IwgcvRj weGc 20wgtMvt	329-08-29	13.55% (mxgv 95%-105%)	wm-140403	gvbwbnfÝ nIqvq c`wUi ‡iwR‡÷ªkb mvgwqK evwZj Kiv nBqv‡Q|	wWwRwWG/GmGm-329/06/4429 ZvwiL: 23/03/2016
		11.	¯úvivfU K¨vcmyj	WvBW †divm mvj‡dU weGc 150.03 wgt Mvt + dwjK GwmW weGc 0.5 wgtMvt + wRsKmvj‡dU g‡bvnvB‡WU weGc 61.8 wgtMvt	329-11-39	The sample does not contain any Ferrous Sulphate, Folic Acid & Zinc Sulphatre at all.	Gj140606	gvbwbnfÝ nIqvq c`wUi ‡iwR‡÷ªkb mvgwqK evwZj Kiv nBqv‡Q|	wWwRwWG/GmGm-329/06/4426 ZvwiL: 23/03/2016
		12.	¯úviW·-100 K¨vcmyj	Wv·mvBwKm weGc 100wgtMvt	329-10-60	17.87% (mxgvt 95%-105%)	‡K-1110	gvbwbnfÝ nIqvq c`wUi ‡iwR‡÷ªkb mvgwqK evwZj Kiv nBqv‡Q|	wWG/29-2/1099; ZvwiL: 20/01/2015
		13.	¯úvig· 250 K¨vcmyj	G‡gv·wmwjb weGc 250 wgtMvt	329-06-60	G‡gv·wmwjb mbv³ nq bvB|	G-120701	gvbwbnfÝ nIqvq c`wUi ‡iwR‡÷ªkb mvgwqK evwZj Kiv nBqv‡Q|	wWG/29-2/1099; ZvwiL: 20/01/2015
		14.	¯úvigj G·Uv	c¨vivwmUvgj weGc 500 wgtMvt + K¨v‡dBb weGc 65 wgtMvt	329-07-06	c¨vivwmUvgj 56.27% I K¨v‡dBb 10.78% (mxgvt 95%-105%)	we2 Gg 0515	gvbwbnfÝ nIqvq c`wUi ‡iwR‡÷ªkb mvgwqK evwZj Kiv nBqv‡Q|	wWG/29-2/1099; ZvwiL: 20/01/2015
10	‡gmvm c¨vivWvBm ‡KwgK¨vj BÛvóR, ewikvj|	**15.**	c¨vi‡dbvK 50	WvB‡Kv‡dbvK †mvwWqvg 50 wgt Mvt	130-59-65	67.18% (95%-105%)	0113	gvbwbnfÝ nIqvq c`wUi ‡iwR‡÷ªkb mvgwqK evwZj Kiv nBqv‡Q|	wWwRwWG/GmGm-130/2002/7264; ZvwiL: 01/06/2015
11	‡gmvm ‡ej‡mb dvgvmDwUK¨vjm wjt, Kgjvcyi, dwi`cyi|	**16.**	Acv U¨ve‡jU	c¨vivwmUvgj weGc 500 wgtMvt	069-34-06	97.28wg:Mv: (19.46%) mxgv: (95%-	0212	gvbwbnfÝ nIqvq c`wUi ‡iwR‡÷ªkb mvgwqK evwZj Kiv nBqv‡Q|	wWG/GmGm-200/2014/827; ZvwiL: 15/01/2015

<table>
<tr><th>bs</th><th>cwZôv‡bi bvg</th><th></th><th>Jl‡ai evwYwR¨K bvg</th><th>‡RbwiK bvg</th><th>‡iwR‡÷kb bs (wWGAvi bs)</th><th>c‡UÝx</th><th>e¨vP bs</th><th>MÖnZ e¨e¯'v</th><th>‡iwR‡÷kb mvgwqK evwZ‡ji ZvwiL</th></tr>
<tr><td></td><td></td><td></td><td></td><td></td><td></td><td>105%)</td><td></td><td></td><td></td></tr>
<tr><td>12</td><td>‡gmvm I‡q÷vi dvgwmDwUK¨vjm wjt, †evinvbDwÏb †ivW, KvkxNvU, wm‡jU|</td><td>17.</td><td>AvjmwWb U¨ve‡jU</td><td>‡iwbwUwWb BDGmwc 150wgtMvt</td><td>101-24-29</td><td>80.27%
mxgv: 90%-110%</td><td>010214</td><td>gvbewnfy©Z nIqvq c`wUi ‡iwR‡÷kb mvgwqK evwZj Kiv nBqv‡Q|</td><td>wWwRwWG/GmGm-101/015/17929; ZvwiL: 22/12/2015</td></tr>
<tr><td>13</td><td>‡gmvm wdwb· ‡KwgK¨vj j¨ve‡iUixR wjt, biwms`x|</td><td><b>18.</b></td><td>wdwb· IwgcvRj K¨vcmyj</td><td>IwgcvRj wewc 20 wgtMvt</td><td>20-02-34</td><td>86.76%<br>(95%-105%)</td><td>wewU 002</td><td>gvbewnfy©Z nIqvq c`wUi ‡iwR‡÷kb mvgwqK evwZj Kiv nBqv‡Q|</td><td>wWwRwWG/GmGm-032/2011/7650; ZvwiL: 09/06/2015</td></tr>
<tr><td rowspan="2">14</td><td rowspan="2">‡gmvm ‡Rwb_ dvgwmDwUK¨vjm wjt, we-67, wewmK wkíbMix, Pvixcyi, †dbx</td><td>19.</td><td>WvB‡Kwbj GmAvi U¨ve‡jU</td><td>WvB‡Kv‡dbvK †mwWqvg wewc 100 wgtMvt</td><td>080-162-64</td><td>42.17%
(95%-105%)</td><td>001</td><td>gvbewnfy©Z nIqvq c`wUi ‡iwR‡÷kb mvgwqK evwZj Kiv nBqv‡Q|</td><td>wWwRwWG/GmGm - 080/15/11456; ZvwiL: 10/08/2015</td></tr>
<tr><td>20.</td><td>UvUv‡Rb K¨vcmj</td><td>‡UUvmvBwKb nvB‡Wv‡KvivBW wewc 250wgtMvt</td><td>080-80-60</td><td>75.28%
(90%-125%)</td><td>001</td><td>gvbewnfy©Z nIqvq c`wUi ‡iwR‡÷kb mvgwqK evwZj Kiv nBqv‡Q|</td><td>wWwRwWG/GmGm - 080/15/18142; ZvwiL: 28/12/2015</td></tr>
<tr><td>15</td><td>‡gmvm ‡UK‡bv WvMm wjt MwUicvov, biwms`x|</td><td><b>21.</b></td><td>cvB‡iv‡fU †evjvm</td><td>c¨vivwmUvgj wewc 2 Mvt</td><td>304-52(V)-83</td><td>53.88%<br>(95%-105%)</td><td>001</td><td>gvbewnfy©Z nIqvq c`wUi ‡iwR‡÷kb mvgwqK evwZj Kiv nBqv‡Q|</td><td>wWwRwWG/GmGm - 304/04/18247; ZvwiL: 30/12/2015</td></tr>
<tr><td>16</td><td>‡gmvm ev‡qvm dvgwmDwUK¨vjm wjt, fvIqvj, wgRvcyi m`i, MvRxcyi|</td><td><b>22.</b></td><td>ógwWb U¨ve‡jU</td><td>‡iwbwUwWb BDGmwc</td><td>57-32-29</td><td>85.33%<br>(90.0%-110.00%)</td><td>0101</td><td>gvbewnfy©Z nIqvq c`wUi ‡iwR‡÷kb mvgwqK evwZj Kiv nBqv‡Q|</td><td>wWwRwWG/GmGm - 057/15/17907, ZvwiL: 22/12/2015</td></tr>
<tr><td>17</td><td>‡gmvm Uz¨W dvgv wjt wewmK wkí bMix, Kzgjv|</td><td>23.</td><td>‡cBbAvDU 400 U¨ve‡jU</td><td>AvBeyc‡db 400 wgtMvt</td><td>066-19-65</td><td>AvBeyc‡db mbv³ nq bvB|</td><td>001</td><td>gvbewnfy©Z nIqvq c`wUi ‡iwR‡÷kb mvgwqK evwZj Kiv nBqv‡Q|</td><td>wWwRwWG/GgGj-66/80(Ask-1)/15613; ZvwiL: 09/11/2015</td></tr>
<tr><td>18</td><td>‡gmvm ggZvR dvgwmDwUK¨vjm wjt, cU bs-551, wc, eo †`Iov, U½x, MvRxcyi</td><td><b>24.</b></td><td>fv‡qv‡Rb 2% †jvmb</td><td>‡Rbwmqvb fv‡qv‡jU wewc 2%</td><td>344-19-76</td><td>27.40%<br>(95%-105%)</td><td>1585</td><td>gvbewnfy©Z nIqvq c`wUi ‡iwR‡÷kb mvgwqK evwZj Kiv nBqv‡Q|</td><td>wWwRwWG/GmGm-344/2013/6730; ZvwiL: 21/05/2015</td></tr>
<tr><td>19</td><td>‡gmvm wgw÷K dvgwmDwUK¨vjm wjt, 16, evM`x, biwms`x|</td><td>25.</td><td>cvg-20 K¨vcmyj</td><td>IwgcvRj wewc 20 wgtMvt</td><td>121-31-29</td><td>82.90%
(95%-105%)</td><td>227</td><td>gvbewnfy©Z nIqvq c`wUi ‡iwR‡÷kb mvgwqK evwZj Kiv nBqv‡Q|</td><td>wWG/GmGm-121/15/17816; ZvwiL: 20/12/2015</td></tr>
<tr><td>20</td><td>‡gmvm dwgK j¨ve‡iUixR wjt Lyjkx, PÆMvg|</td><td>26.</td><td>‡UUvmvBwKb K¨vcmyj</td><td>‡UUvmvBwKb nvB‡Wv‡KvivBW 250 wgtMvt</td><td>179-43-60</td><td>‡UUvmvBwKb nvB‡Wv‡KvivBW mbv³ nq bvB|</td><td>130934</td><td>c`wUi ‡iwR‡÷kb mvgwqK evwZj Kiv nBqv‡Q|</td><td>wWwRwWG/GmGm - 179/06/14975; ZvwiL: 25/10/2015</td></tr>
</table>

bs	cÖZôv‡bi bvg		Jl‡ai eªvÛ bvg	‡RbvwiK bvg	‡iwR‡÷ªkb bs (wWGAvi bs)	c‡UÝx	e¨vP bs	MÖnY e¨e¯’v	‡iwR‡÷ªkb mvgwqK evwZ‡ji ZvwiL
	‡gmvm dvgvK j¨ve‡iUixR wjt Lyjbv, PÆMÖvg|	27.	wg‡KvRj-400wgtMvt U¨ve‡jU	‡g‡UªvbWvRj wewc 400wgtMvt	179-49-57	97.91% (95%-105%) c`wU wWRBb‡U‡Mkb cixÿvq DËxb nqwbGes c¨v‡K (wewa÷vi) G Drcv`b I †gqv‡`vËxY ¯úó bq|	150402	gvbwnfyZ nIqvq c`wUi ‡iwR‡÷ªkb mvgwqK evwZj Kiv nBqv‡Q|	wWwRwWG/GmGm - 179/06/14975; ZvwiL: 25/10/2015
		28.	wce¨vK U¨ve‡jU	wm‡cvd·wmb nvB‡Wª‡K¬vivBW BDGmwc 500 wgtMvt	179-42-60	wm‡cvd·wmb nvB‡Wª‡K¬vivBW mbv³ nq bvB	141101	gvbwnfyZ nIqvq c`wUi ‡iwR‡÷ªkb mvgwqK evwZj Kiv nBqv‡Q|	wWwRwWG/GmGm - 179/06/14975; ZvwiL: 25/10/2015
21	‡gmvm B‡›`v-evsjv dvgvwmDwUK¨vjm IqvKm wjt, K‡jR †ivW, ewikvj|	29.	B‡›`v‡dbvK 100 GmAvi K¨vcmyj	WvB‡K¬v‡dbvK †mvwWqvg wewc 100 wgtMvt	032-49-06	59.38% (95%-105%)	010115	gvbwnfyZ nIqvq c`wUi ‡iwR‡÷ªkb mvgwqK evwZj Kiv nBqv‡Q|	wWwRwWG/GmGm - 032/13/17012 ZvwiL: 03/12/2015
		30.	‡KwUg· wWGm U¨ve‡jU	mvjdv‡gš_v·vRj wewc 800wgtMvt + UvBwg‡_wcg wewc 160wgtMvt	032-28-59	32.88% (92.5%-107.5%)	020714	gvbwnfyZ nIqvq c`wUi ‡iwR‡÷ªkb mvgwqK evwZj Kiv nBqv‡Q|	wWwRwWG/GmGm - 032/13/9776; ZvwiL: 13/07/2015
		31.	‡Ubmwwij U¨ve‡jU	WvqwRcvg wewc 5 wgtMvt	032-63-02	66.6% (92.5%-107.5%)	010114	gvbwnfyZ nIqvq c`wUi ‡iwR‡÷ªkb mvgwqK evwZj Kiv nBqv‡Q|	wWwRwWG/GmGm- 032/13/13154; ZvwiL: 17/09/2015
		32.	B‡›`wi÷b Avi 150 U¨ve‡jU	‡ivwbwUwWb BDGmwc 150wgtMvt	032-93-29	50.93% (90%-110%)	230912	gvbwnfyZ nIqvq c`wUi ‡iwR‡÷ªkb mvgwqK evwZj Kiv nBqv‡Q|	wWwRwWG/GmGm - 032/13/2081; ZvwiL: 17/01/2014
	‡gmvm B‡›`v-evsjv dvgvwmDwUK¨vjm IqvKm wjt, K‡jR †ivW, ewikvj|	33.	‡KwUg· mvm‡cbmb	mvjdv‡gš_v·vRj wewc 4.0Mvt + UvBwg‡_wcg wewc 0.80	032-16-59	সক্রিয় উপাদান সনাক্ত nq bvB	171211	cixÿvZ bgybvwU evwn¨Kfv‡e `yÜhy³ I Kvj‡P ev`vgx is aviY	wWwRwWG/GmGm - 032/13/2081; ZvwiL: 17/01/2014

bs	cwZôv‡bi bvg		JI‡ai ewYwR¨K bvg	‡Rbwi‡K bvg	‡iwR‡÷kb bs (wWGAvi bs)	c‡UÝx	e¨vP bs	MÖnZ e¨e¯¯w`	‡iwR‡÷kb mvgwqK ewZ‡ji ZwiL
				Mvt/100wgwj				Kivq gvbewnfÿZ nIqvq c`wUi ‡iwR‡÷kb mvgwqK ewZj Kiv nBqv‡Q|	
		34.	‡gUj mvm‡cbmb	‡g‡UwbWvRj we‡c 4.0Mvg/100wgtwjt	032-26-56	40.09% (95%-105%)	150812	gvbewnfÿZ nIqvq c`wUi ‡iwR‡÷kb mvgwqK ewZj Kiv nBqv‡Q|	wWRwWG/GmGm - 032/13/2081; ZwiL: 17/01/2014
		35.	‡gUj U¨ve‡jU	‡g‡UwbWvRj 400 wgtMvt	032-25-56	39.65% (95%-105%)	050412	gvbewnfÿZ nIqvq c`wUi ‡iwR‡÷kb mvgwqK ewZj Kiv nBqv‡Q|	wWRwWG/GmGm - 032/13/2081; ZwiL: 17/01/2014
		36.	B‡›`v‡c·-we U¨ve‡jU	wfUwgb we-Kg‡c·	032-30-39	সক্রিয় উপাদান সনাক্ত nq bvB	071011	gvbewnfÿZ nIqvq c`wUi ‡iwR‡÷kb mvgwqK ewZj Kiv nBqv‡Q|	wWRwWG/GmGm - 032/13/2081; ZwiL: 17/01/2014
		37.	c¨vivwmUvgj U¨ve‡jU	c¨vivwmUvgj we‡c 500wgtMvt	032-18-06	38.85% (95%-105%)	020212	gvbewnfÿZ nIqvq c`wUi ‡iwR‡÷kb mvgwqK ewZj Kiv nBqv‡Q|	wWRwWG/GmGm - 032/13/2081; ZwiL: 17/01/2014
		38.	wi‡evdwfb U¨ve‡jU	wi‡evdwfb we‡c 5wgtMvt	032-08-39	38.32% (95%-115%)	141011	gvbewnfÿZ nIqvq c`wUi ‡iwR‡÷kb mvgwqK ewZj Kiv nBqv‡Q|	wWRwWG/GmGm- 032/13/2722; ZwiL: 20/02/2013
		39.	B‡›`vd·K¨vcmÿ	dK·wmwjb we‡c 500wgtMvt	032-36-60	31.22% (92.5%-110%)	020811	gvbewnfÿZ nIqvq c`wUi ‡iwR‡÷kb mvgwqK ewZj Kiv nBqv‡Q|	wWRwWG/GmGm- 032/13/2722; ZwiL: 20/02/2013
		40.	B‡›`vg·bK¨vcmÿ	G‡gw·wmwjb we‡c 500wgtMvt	032-34-60	27.01% (92.5%-110%)	010112	gvbewnfÿZ nIqvq c`wUi ‡iwR‡÷kb mvgwqK ewZj Kiv nBqv‡Q|	wWRwWG/GmGm- 032/13/2722; ZwiL: 20/02/2013
		41.	BbW·K¨vcmÿ	Ww·mvBwKb we‡c 100 wgtMvt	032-50-60	44.27% (95%-105%)	031211	gvbewnfÿZ nIqvq c`wUi ‡iwR‡÷kb mvgwqK ewZj Kiv nBqv‡Q|	wWRwWG/GmGm- 032/13/2722; ZwiL: 20/02/2013
22	‡gmvm BDwbqb dvgv wjt mvfvi, XvKv|	**42.**	G‡Wvj mvm‡cbkb	c¨vivwmUvgj we‡c 120wgtMvt/5wgtwjt	207-19-06	79.4% (95%-105%)	010709	gvbewnfÿZ nIqvq c`wUi ‡iwR‡÷kb mvgwqK ewZj Kiv nBqv‡Q|	wWRwWG/GmGm- 207/05/2769; ZwiL: 03/03/2013

bs	cwZôv‡bi bvg		JI‡ai ewYwR¨K bvg	‡RvbwiK bvg	‡iwR‡÷kb bs (wWGAvi bs)	c‡UÝx	e¨vP bs	MyjnZ e¨e¯wn`	‡iwR‡÷kb mvgwqK ewZ‡ji ZwiL
23	‡gmvm I‡qwmm j¨ve‡iUixR wjt wmivRMÄ|	43.	I-dwfb U¨ve‡jU	wi‡fvdwfb 5 wgtMvt	161-30-39	সক্রিয় উপাদান সনাক্ত nq bvB	Aviwe-06	gvbewnfyZ nIqvq c`wUi ‡iwR‡÷kb mvgwqK ewZj Kiv nBqv‡Q|	wWwRwWG/GmGm-161/13/6190; ZwiL: 21/05/2013
			44.	‡iwbwmm U¨ve‡jU	‡iwbwUwWb nvB‡Ww‡KvivBW 150wgtMvt	161-47-29	7.75% (90%-110%)	AviGm-02	gvbewnfyZ nIqvq c`wUi ‡iwR‡÷kb mvgwqK ewZj Kiv nBqv‡Q|	wWwRwWG/GmGm-161/13/6190; ZwiL: 21/05/2013
24	‡gmvm ÷vim dvgwmDwUK¨vjm wjt, ewikvj|	45.	‡iwbwUwWb 150 U¨ve‡jU	‡iwbwUwWb wewc 150 wg:Mv:	263-15-29	‡iwbwUwWb mbv³ nq bvB|	0110	gvbewnfyZ nIqvq c`wUi ‡iwR‡÷kb mvgwqK ewZj Kiv nBqv‡Q|	wWwRwWG/GmGm-263/11/11241; ZwiL: 02/10/2011

Annex-F

Proposed Product for locally manufacture (Medical Devices)

নং	প্রস্তুতকারকের নাম	‡gvW‡Kj wWfvB‡mi bvg		জেনেরিক নাম	wb‡`kbv/e¨envi	Contra-indication & Side-effect	আবেদনকারী প্রদত্ত USFDA/BNF/ MHRA Ref.	‡UKwbK¨vj mve-KwgwUi wm×vš	mfvi wm×vš
1.	OMC Healthcare (Pvt.) Ltd., Rupnagar Industrial Area, Plot#44, Block-K, Road#4, Mirpur-2, Dhaka-1216, Bangladesh	a)-1	One-Step Pregnancy Test **Class: B**	Human Chorionic Gonadotropin (HCG) Pregnancy Test	Self-performing in-vitro immunoassay in urine for early detection of pregnancy	**Contraindication & side-effect:** None	-	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
2.		a)-2	One-Step Dengue Test **Class:B**	Dengue Test	Dengue Antigen and Antibody in-vitro immunoassay for early detection of Dengue	**Contraindication & side-effect:** None	-	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

Proposed Product for Import (Medical Devices):

নং	প্রস্তুতকারকের নাম	ewYRK bvg	‡gvW‡Kj vWfvB‡mi bvg	wb‡`kbv/e¨envi	Contra-indication & Side-effect	FSC/CPP	‡UKwbK¨vj mve-KgvUi wm×vš	mfvi wm×vš
1.	Romsons International, India **Importer:** Barisal Surgical, 34/1, Mitford Road, Dhaka-1100	Guedel Airways (Sterile) **Class: B**	Oro Pharyngeal Airways	Used to maintain an unobstructed airway during general anaesthesia and in patients who are unconscious for other reasons.	Contraindication: None Side-effect : None	FSC-India EC Certificate-Norway	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj	
2.	**Name & Address of Manufacturer** C.R. Bard, INC. 8195 Industrial BLVD, Covington, GA, USA-30014 **Manufacturing Site** Bard Peripheral Vascular, Inc. USA (**Local Agent;** Lilac pvt. Ltd 72, New Elephant Road, Dhaka-1205)	**Bard Max-Core** Disposable Core Biopsy Instrument **Class: B**	Disposable Core Biopsy Instrument (Needle)	It is intended for use in obtaining biopsies from soft tissue such as Liver, Kidney, prostate spleen lymph nodes, and various soft tissues tumors.	**Contraindication**: Not for use in Bone. **Warning**: *Good medical judgement should be exercised in considering biopsy on patients who are receiving anticoagulant therapy or who have a bleeding disorder. *The collection of multiple needle cores may help to ensure the detection of any cancer tissue. *A "negative" biopsy in the presence of suspicious radiographic findings does not preclude the presence of carcinoma. *Post-biopsy patient care may vary with the biopsy technique utilized and the individual patients physiological condition. Observation of vital signs and other precautions should be taken to avoid and/or treat potential complications that may be associated with biopsy procedures. *Core biopsy needles are for single patient use only. Do not reuse. Do not desterilize. *After use, this product may be potential biohazard. Handle and dispose of in accordance with accepted medical practice and with applicable laws and regulations. *This device has been designed for clean once body fluids or tissues with potential	Certificate to Foreign Gov.-USFDA CE Certificate-bsi, UK	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj	

3.	**Name & Address of Manufacturer** C.R. Bard, INC. 8195 Industrial BLVD, Covington, GA, USA-30014 **Manufacturing Site** Bard Peripheral Vascular, Inc. USA (**Local Agent;** Lilac pvt. Ltd 72, New Elephant Road, Dhaka-1205)	**Bard Magnum** Disposable Core Tissue Biopsy Needle **Class: B**	Disposable Core Tissue Biopsy Needle	It is intended for use in obtaining biopsies from soft tissue such as Liver, Kidney, prostate spleen lymph nodes, and various soft tissues tumors.	**Contraindication:** Not for use in bone. **Warning**: *Good medical judgement should be exercised in considering biopsy on patients who are receiving anticoagulant therapy or who have a bleeding disorder. *The collection of multiple needle cores may help to ensure the detection of any cancer tissue. *A "negative" biopsy in the presence of suspicious radiographic findings does not preclude the presence of carcinoma. *Post-biopsy patient care may vary with the biopsy technique utilized and the individual patients' physiological condition. Observation of vital signs and other precautions should be taken to avoid and/or treat potential complications that may be associated with biopsy procedures. *Core biopsy needles are for single patient use only. Do not reuse. Do not desterilize. *If collecting multiple samples, inspect the needle for damaged point, bent shaft or other imperfections after each sample is collected. Do not use needle if any imperfection is noted. *After use, this product may be potential biohazard. Handle and dispose of in accordance with accepted medical practice and with applicable laws and regulations. *This device has been designed for single use only. *Do not resterilize. Afterresterilization the sterility of the product is not guaranteed because of an inderterminable degree of potential pyrogenic or microbial contamination which may lead to infectious complications, cleaning, reprocessing.	Certificate to Foreign Gov.-USFDA CE Certificate-bsi, UK	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj
4.	Microtrack Surgicals A38, Adarsh2 Industrial Estate, B/h.Ashish Cinema, Odhav Ahmedabad -382415 India. **<u>Local agent:</u>** Zas Corporation 80/22, Mymensingh Road, Banglamotor , Dhaka-1000	**Microtrack** Ophthalmic Micro Surgical Full Handle Knife **Class: B**	Ophthalmic Blades	Surgical intervention in various types of ophthalmic surgeries.	**Contraindication:** None **Side Effect:** None	FSC-India EC Certificate (Czech Republic)	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj

5.	**Manufacturer:** Onbo Electronic (Shenzhen), Co. Ltd, China **Supplier:** **Microlife Corporation** 9F, 431, RuiGang Road, NehHu, Taippei 114, Taiwan, R.O.C **Local agent:** **Transcom Distribution Co.** Ltd 52, Motijheel C/A, Dhaka-1000	Digital Blood Pressure Monitor (Mannual) **Class: B**	Blood Pressure Machine	It is used to mesure Blood Pressure	Contraindication: None Side effect: None	Certificate for Exportation of Medical Products-China	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj	
6.	**Manufacturer:** Onbo Electronic (Shenzhen), Co. Ltd, China **Supplier:** **Microlife Corporation** 9F, 431, RuiGang Road, NehHu, Taippei 114, Taiwan, R.O.C **Local agent:** **Transcom Distribution Co.** Ltd 52, Motijheel C/A, Dhaka-1000	Digital Blood Pressure Monitor (Automatic) **Class: B**	Blood Pressure Monitor	It is used to mesure Blood Pressure	Contraindication: None Side effect: None	Certificate for Exportation of Medical Products-China	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj	

7.	Terumo Clinical Supply Co. Ltd., 3 Kawashima-Takehayamachi, Kakamigahara, Japan **Local Agent:** UniMed Ltd. (Medical Device) 34/1 Sonagaon Road, Paribag, Dhaka	Eliminate (EG 1602 & EG 1652) **Class: B**	Aspiration Catheter	It is intended for the removal of fresh, soft emboli and thrombi from vessels in the coronary and peripheral vasculature.	**Contra-indication :** It is contraindicated in: * Vassels <1.8mm in diameter as for ELT6FGC, <2.05mm in diameter as for ELT7FGC and<2.2mm in diameter as for ELT8FGC • The removal of fibrous, adherent or calcified material • To be use in the venous system **Complications:** Complications associated with the use of the eliminate are similar to the ones associated with standard percutaneous interventional procedures. Possible complications may include, but are not limited to : Local or systemic infection, Local haematomas, Intimal disruption, Arterial dissection, perforation, rupture or injury, Arterial thrombosis, Distal embolization of blood clots and plaque, Arterial spasm, Arteriovenous fistula formation, Pseudoaneurysm or Bleeding Complication at Access Site, Acute myocardial infarction, Arryhythmias, including life-threatening ventricular fibrillation, Stroke/ CVA, Death, Emergent or Non- emergent Bypass Graft Surgery, Haemorrhage, Myocardial Ischemia & Hypotension.	FSC-Japan CE Marking of Conformity	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj

8.	Sewoon Medical Co. Ltd, South Korea. **Local Agent:** **M. Enterprise,** H-9, R-20, Block-C, Sec-10, Mirpur, Dhaka-1216	ALL-Silicone Foley Balloon Catheter Size 2 Way: 6,8,10,12,14,16,18, 20Fr Size 3Way 16, 18, 20, 22Fr **Class: B**	Balloon Catheter, Urological	**Diagnostic indications :** Collection of uncontaminated urine specimen, Monitoring of urine output, Imaging of the urinary tract **Therapeutic Indication:** Acute urinary retention, Chronic obstruction that causes hydronephrosis, Initiation of continuous bladder irrigation, Intermittent decompression for neurogenic bladder, Hygienic care of bedridden patients.	**Contraindications:** Urethral catheterization is contraindicated in the presence of traumatic injury to the lower urinary tract (e.g. urethral tear). This condition may be suspected in male patients with a pelvic or straddle-type injury. Signs that increase suspicion for injury are a high-riding or boggy prostate, perineal hematoma, or blood at the meatus. When any of these findings are present in the setting of possible trauma, a retrograde urethrogram should be performed to rule out a urethral tear prior to placing a catheter into the bladder. **Side Effects:** Infections, including urethritis, cystitis, pyelonephritis and transient bacteremia, Paraphimosis, caused by failure to reduce foreskin after catherization, Creation of false passages, Urethral strictures, Urethral perforation, Bleeding.	FSC-South Korea & EC Certificate (UK)	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
9.	Ashitaka Factory of Terumo Corporation , Japan **Local Agent**: UniMed Ltd. (Medical Device) 34/1 Sonagaon Road, Paribag, Dhaka	**Radifocus Introducer II** [It is consits of Introducer (a sheath and a dialation), a mini guide wire, an entry needle and a syringe] **Class: B**	Catheter Introducer	It is intended to be inserted percutaneously into a vessel to facilitate the insertion of angiographic, electrode, balloon or similar catheters.	**Contra-indication :** None **Side effect :** None	FSC-Japan EC Certificate	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj

10.	Sewoon Medical Co. Ltd, South Korea. **Local Agent:** **M. Enterprise,** H-9, R-20, Block-C, Sec-10, Mirpur, Dhaka-1216	ALL- Silicone Gastric Duodenal LEVIN Tube Size : 8,10,12,14,16,18,20,22 Fr **Class: B**	Catheter, Gastrointestinal	**Diagnostic Indication:** Evaluation of upper gastrointestinal (GI) bleed, Aspiration of gastric fluid content, Identification of the esophagus and stomach on a chest radiograph, Administration of radiographic contrast to the GI tract. **Therapeutic Indication:** Gastric decompression, including maintenance of a decompressed state after endotracheal intubation, often via the oropharynx, Relief of symptoms and bowel rest in the setting of small-bowel obstruction, Aspiration of gastric content from recent ingestion of toxic material, Administration of medication, Feeding, Bowel irrigation.	**Contraindications:** Severe midface trauma, Recent nasal surgery relative contraindications, coagulation abnormality, Esophageal varices or stricture, Recent banding or cautery of esophageal varices, Alkaline ingestion. **Side Effects:** Patient discomfort, Epistaxis may be prevented by generously lubricating the tube tip and using a gentle technique, Respiratory tree intubation, Esophageal perforation.	FSC-South Korea & EC Certificate (UK)	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

11.	Sewoon Medical Co. Ltd, South Korea. **Local Agent:** **M. Enterprise,** H-9, R-20, Block-C, Sec-10, Mirpur, Dhaka-1216	Urinary Self Catheter **Size:** 10, 12,14,16 Fr **Class: B**	Catheter, Urological	**Diagnostic indications :** Collection of uncontaminated urine specimen, Monitoring of urine output, Imaging of the urinary tract **Therapeutic Indication:** Acute urinary retention, Chronic obstruction that causes hydronephrosis, Initiation of continuous bladder irrigation, Intermittent decompression for neurogenic bladder, Hygienic care of bedridden patients.	**Contraindications:** Urethral catheterization is contraindicated in the presence of traumatic injury to the lower urinary tract (e.g. urethral tear). This condition may be suspected in male patients with a pelvic or straddle-type injury. Signs that increase suspicion for injury are a high-riding or boggy prostate, perineal hematoma, or blood at the meatus. When any of these findings are present in the setting of possible trauma, a retrograde urethrogram should be performed to rule out a urethral tear prior to placing a catheter into the bladder. **Side Effects:** Infections, including urethritis, cystitis, pyelonephritis and transient bacteremia, Paraphimosis, caused by failure to reduce foreskin after catherization, Creation of false passages, Urethral strictures, Urethral perforation, Bleeding.	FSC-South Korea & EC Certificate (UK)	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
12.	**Name & Address of Manufacturer** C.R. BARD, INC. 8195 Industrial BLVD, Covington, GA, USA-30014 **Manufacturing Site** Bard SDN BHD LOT- 57-C,Kulim Industrial Estate Kulim Kedah , Malaysia 09000, (**Local Agent;** Lilac pvt. Ltd 72, New Elephant Road, Dhaka-1205)	Bard Biocath Suprapubic Cathterisation Set Size: 12 & 16 Fr **Class: B**	Cathterisation Set	It is indicated for use in urethral strictures or fistula and for outflow obstruction, chronic retention, urethral trauma, pre-, intra- and post-operative bladder drainage requested by physician.	**Contraindications:** None **Side Effects:** None	Certificate to Foreign Gov.-USFDA	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

13.	**Manufacturer :** Boston Scientific Corporation, 300 Boston Scientific Way, USA **Local Agent :** Vastech Limited, Nurjehan Tower (6th Floor) 80/22 Mymensingh Road Dhaka-1000. Bangladesh.	Extractor Pro **Class: B**	Retrieval Balloon Catheter	The Extractor Pro XL Retrieval Balloon Catheter is used endoscopically to 1) remove stones from the biliary system; or 2) to facilitate injection of contrast medium while occluding the duct with the balloon.	**Contraindications:** Contraindications for this device are those specific to endoscopic retrograde cholangiopancreatography (ERCP) and endoscopic sphincterotomy (ES). **Adverse events:** Possible complications include, but may not be limited to: pancreatitis, perforation, hemorrhage, hematoma, cholangitis, stone impaction, septicemia/infection, allergic reaction to contrast medium	Certificate to foreign Government-USA CE Marking of Conformity (DEKRA)	Abjgv`b Kiv †h‡Z cv‡i |	Abjgv`b Kiv nj |
14.	Romsons International, India **Importer:** Barisal Surgical, 34/1, Mitford Road, Dhaka-1100	Suction catheter thumb control **Class: B**	Suction catheter (Suction Catheter-TC is a suction catheter equipped with feature -Thumb Control Vacuum Valve for optional intermittent)	Suctioning of tracheo-bronchial secretions.	Contraindication: None Side-effect : None	FSC-India EC Certificate-Norway	Abjgv`b Kiv †h‡Z cv‡i |	Abjgv`b Kiv nj |
15.	Romsons International, India **Importer:** Barisal Surgical, 34/1, Mitford Road, Dhaka-1100	Chest drainage catheter **Class: B**	Thoracic drainage catheter	Chest Drainage Catheter is sterile, single use device intended for post operative drainage after cardiothoracic surgery.	Contraindication: None Side-effect : None	FSC-India EC Certificate-Norway	Abjgv`b Kiv †h‡Z cv‡i |	Abjgv`b Kiv nj |
16.	Romsons Scientific & Surgical Industries Pvt. Ltd, India **Importer:** Barisal Surgical, Dhaka	Catheter Mount **Class: B**	Catheter Mount 'T' Piece with intermittent suction port collapsible tubing 22mm x15mm swivel connector	Catheter Mount ('T' Piece) is generally used for connection with breathing/ventilation Circuit for convenience of intermittent suction and gas sampling	Contraindication: None Side-effect : None	FSC-India EC Certificate-Norway	Abjgv`b Kiv †h‡Z cv‡i |	Abjgv`b Kiv nj |

17.	SIMEKS TIBBI URUNLER SANAYI Ve TICARET Limited Sirketi, Turkey (**Importer:** Biva International)	**SIMPASS PLUS** PTCA Balloon Dilatation Catheter **Class: D**	PTCA Balloon Dilation Catheter	It is indicated for balloon dilatation of the stenotic portion of a coronary artery for the purpose of improving myocardial perfusion.	**Contraindication:** Unprotected left main artery. **Adverse Events:** Possible adverse effects include, but are not limited to the following: Death, Acute myocardial infarction, Total occlusion of the coronary artery or bypass graft, Restenosis of dilated vessel, Coronary vessel dissection, perforation, rupture or injury, Hemorrhage or hematoma, Unstable angina, Arrhythmias , including ventricular fibrillation, Drug reactions, allergic reaction to contrast media, Hypo/ hypertension, Infection, Coronary artery spasm, Arteriovenous fistula, Embolism.	FSC- Turkey EC Certificate	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

18.	GELITA MDICAL GmbH: Uferstrasse 7 69412 Eberbach Germany [**SURGITA LTD**. : 58/3, Lake Circus, Kalabagan, Dhanmondi, Dhaka – 1205, Bangladesh]	**GELITA-CEL® CA POWDER** **Class: D**	Absorbable Oxidized Cellulose Powder Hemostat	Topical hemostat for use as an adjunct to hemostasis in particular where control of capillary, venous, and arteriolar bleeding, by pressure, ligature, andother conventional procedures, is either ineffective or impractical.The GELITA-CEL® CA POWDER is especially suitable for vulnerable applications on the brain parenchyma, cranial nerves, the brainstem (where surgicalmanipulation may lead to neuronal damage) or on the dura mater. Because of the possibility of applying a very fine thin layer, it will not clog up openings in the nervous system such as the mesencephalic duct, obex and ventricles. For spinal surgery the GELITA-CEL® CA POWDER applicator facilitates application in narrow or difficult to reach areas between bones as in discectomiesand laminectomies.	**Contraindication:** It should not be used in bleeding from large arteries. Washing away of the preparation by blood can be considered as anindication of inappropriate use. It should not be used for implantation in bone defects, such as fractures, since there is a possibility of interference with callusformation and a theoretical chance of cyst formation. It should not be used in conjunction with methyl methacrylate adhesives, for example in orthopedic surgery, because theirpresence may reduce the adhesive strength of the bonding agent to bone. It is a bio-degradable hemostat, and should not be used as an adhesion prevention product. It should not be used on non-hemorrhagic serous oozing surfaces, since body fluids other than whole blood, such as serum, do not react with the product to produce satisfactory hemostatic effect. **Side Effects:** Occasional reports of “burning” and “stinging” sensations and sneezing when oxidized cellulose has been used as packing in epistaxis, are believed to be due to the low pH of the product. Burning has been reported when oxidized cellulose was applied after nasal polyp removal and after hemorrhoidectomy.	FSC- Germany CE Marking of Conformity EC Certificate (Germany)	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj

19.	M/S Comed B.V De Marne 118 8701 MC Bolsward The Netherlands (**Importer:** Advanced Meditech Suite: 115 (1st Floor), Krishnachura Commercial Complex, 24/B.C Shahid Minar Road, Kallanpur, Dhaka-1207, Bangladesh)	Hyperion PDA Occuluder **Class: D**	Occuluder with Delivery System	The Hyperion PDA Occuluder can be used in treating patent ducts arteriousus. It is applicable for various simple patent ducts arteriousus with the patent arteriousus diameter of between 2.15mm.	**Contraindications:** ➢ Patient ducts arteriousus complicated with severe pulmonary hypertension at the presence of left to right shunt. ➢ Hemorrhagic disease such as active ulcer. ➢ Other abnormalities are present in need of surgical treatment. ➢ Recent infection is found patient. ➢ Patient s weighing less than 6kg or patient less than 7 months of age. ➢ Presence of thombus at the intended site of implant or documented evidence of venous thrombus in the vessel through which access to the defected is gain ➢ Patients whose vasculature through which access to the defects is gained, is inadequate to accommodate the appropriate sheath size. **Adverse events:** The implanted Occuluder should completely occlude the patent ductus arteriousus which has same therapeutic effects with surgical treatment. However, the following complications may emerge during and after the implanting process; Hemorrhage and homeostasis around the puncture site, formation of Haematoma, impairment of femoral artery, femoral vein and femoral nurve, cardiac arrythmia, endocarditis, incomplete occlusion accompanied by mechanical hemolysis.	The Netherlands EC Certificate (Germany)	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj

20.	Fresenius Medical Care Homburg, Germany **Local agent :** Janata Traders	**FXClassix** **Class: C**	Capillary High-Flux Dialyser	It is designed for single use in chronic haemodialysis.	**Contraindication:** Special contraindications are unknown.Generally contraindication for haemodialysis are applicable. Patients with known hypersensitivity to any of the dialyser's material must not treated with the dialyzer. **Side Effects:** Certain side effects may occur during dialysis and may result from factors specific to the patient, operating parameters, equipment, priming procedure, dialysis solution anticoagulation, medication etc. Hypersensitivity or hypersensitivity like reaction has been observed. Symptomatology can vary and may include dyspnoea, Chest congestion, bronchospasm, respiratory arrest, hypotension, tachycardia, urticaria, abdominal pain, nausea, convaltion and unconsciousness.	FSC-Germany	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj
21.	Fresenius Medical Care Homburg, Germany **Local agent :** Janata Traders	**FXClass** **Class: C**	Capillary Low-Flux Dialyzer	It is designed for single use in chronic haemodialysis or haemo filtration respectively.	**Contraindication:** Special contraindication is unknown.Generally contraindication for haemodialysis are applicable. Patients with known hypersensitivity to any of the dialyser's material must not treated with the dialyzer. **Side effect:** During dialysis and may result from factors specific to the patient, operating parameters, equipment, priming procedure, dialysis solution anticoagulation, medication etc. Hypersensitivity or hypersensitivity like reaction has been observed. Symptomatology can vary and may include dyspnoea, Chest congestion, bronchospasm, respiratory arrest, hypotension, trachycardia, urticaria, abdominal pain, nausea, convaltion and unconsciousness.	FSC-Germany	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj

22.	Fresenius Medical Care Homburg, Germany **Local agent :** Janata Traders	**Hemoflow Dialyzer** **Class: C**	Dialyzer	Used for Dialysis & Apheresis Therapies	**Contraindications:** Specific contraindications for the dialyzer are unknown. Generally, the contraindications for hemodialysis are applicable. The dialyzer should only be used as directed by a physician. **Side effects**: In rare cases hypersensitivity reactions may occur during hemodialysis treatment. A history of allergies is an indication for careful monitoring of hypersensitivity reactions. Dialyzers of this type should not be used again on any patient exhibiting a hypersensitivity reaction. With severe reactions, dialysis must be discontinued and aggressive, first line therapy for anaphylactic reactions must be initiate.	FSC-Germany	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
23.	Fresenius Medical Care Homburg, Germany **Local agent :** Janata Traders	Citrosteril **Class: B**	Heat Disinfection of Haemodialysis Machines with recirculation	The synergistic effect of its components makes Citrosteril a potent disinfectant solution. Citrosteril at 84○C has a broad Spectrum of microbiocidal activity and work bactericidal, virus inactivating (HBV, HCV, HIV) and fungicidal.	**Contraindications:** Do NOT use Benzalkonium Chloride Solution if: you are allergic to any ingredient in Benzalkonium Chloride Solution. you have a deep puncture wound, animal bite, or serious burn. **Side effects:** Severe allergic reactions (rash; hives; itching; difficulty breathing; tightness in the chest; swelling of the mouth, face, lips, or tongue); infection; severe or persistent irritation,	FSC-Germany	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
24.	**Manufacturer :** Smith & Nephew Inc., 101 Hessle Road, Hull, HU3 2BN, UK **Local Agent :** Vastech Limited, Nurjehan Tower (6th Floor) 80/22 Mymensingh Road Dhaka-1000. Bangladesh	Opsite Post-Op **Class: B**	Absorbent, waterproof and bacteria proof film dressing	It is indicated to dress wounds such as lacerations, cuts, abrasions, post-operative wounds, minor burns, where levels of exudate are low/moderate.	**Contra-Indication:** None **Side Effects:** None	FSC-UK EC Certificate (bsi)	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

25.	**Manufacturer :** Smith & Nephew Inc., 101 Hessle Road, Hull, HU3 2BN, UK **Local Agent :** Vastech Limited, Nurjehan Tower (6th Floor) 80/22 Mymensingh Road Dhaka-1000. Bangladesh.	Allevyn Ag Adhesive/Non-adhesive/Sacrum **Class: B**	Adhesive Hydrocellular Antimicrobial Dressing	It is indicated as an antimicrobial absorbent dressing for the management, by secondary intention of chronic and acute full thickness, partial thickness or shallow granulating, exuding wounds such as: pressure ulcers, venous ulcers, diabetic ulcers, burns, donor sites, fungating/malignant wounds and surgically dehisced wounds	**Contra-indications:** None **Side Effects:** None	FSC-UK EC Certificate (bsi)	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |
26.	BSN Private Ltd.India **Contract manufacturer** Adeshwar meditex Pvt.Ltd.India **Local agent :** **Janata Traders**	**Cutisorb** **Class: B**	Combine dressing sterile	Heavily exuding wounds, which are chronic or healing by secondary intent, such as leg ulcers, pressure ulcers, diabetic foot ulcers and similar types of wounds.	**Contraindication:** None **Side effect:** There are no known side effects. The dressing should not be used on dry wounds, low exuding wounds, eyes, mucous membranes or in wound cavities (the dressing swells considerably after fluid absorption!)	FSC-India	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |
27.	ConvaTec Limited USA. **Local agent** : Janata Traders, TCB Bhabon , 1 kawran Bazar,Dhaka	**DuoDerm-** Extra Thin, CGF **Class: B**	Hydrocolloid Dressing	Management of superficial, dry to lightly exudating dermal ulcers, post operative wound, protective dressing.	**Contraindication:** Should not be used on individual who are sensetive to or who have had an allergic reaction to the dressing or its components. **Side effect:** None	CFG-USFDA	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |
28.	**ConvaTec Inc, USA** **Local agent :** Janata Traders TCB Bhabon , 1 kawran Bazar,Dhaka	**Aquacel-** Ag , Extra, Burn, Glove, Foam ADH, Foam N/ADH ,SCD **Class: B**	Hydrofiber Dressing	It is used for abrasions, lacerations, minor cut, burns, leg ulcers , pressure ulcer, diabetic ulcer, traumatic wounds.	**Contraindication:** It should not be used on individuals who are sensitive to or who have had an allergic reaction to the dressing or its components. **Side effect:** None.	CFG-USFDA	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |

29.	**Manufacturer :** Smith & Nephew Inc., 101 Hessle Road, Hull, HU3 2BN, UK **Local Agent :** Vastech Limited, Nurjehan Tower (6th Floor) 80/22 Mymensingh Road Dhaka-1000. Bangladesh	Intrasite Gel **Class: B**	Hydrogel wound dressing	It is is indicated for the management of shallow and deep open wounds healing by secondary intent, e.g. Venous leg ulcers, Diabetic foot ulcers, Surgical wounds, Pressure sores, Extravasation injuries, Radiation damage, Burns, Fistulae, Amputations, Fungating ulcers.	**Contra-Indication:** None **Side Effects:** None	FSC-UK EC Certificate (bsi)	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
30.	**Manufacturer :** Smith & Nephew Inc., 101 Hessle Road, Hull, HU3 2BN, UK **Local Agent :** Vastech Limited, Nurjehan Tower (6th Floor) 80/22 Mymensingh Road Dhaka-1000. Bangladesh	Melolin **Class: B**	Low adherent absorbent dressing	It is indicated for the management of a wide variety of light to moderately exuding wounds including clean sutured wounds, abrasions, lacerations and minor burns.	**Contra-Indication:** None **Side Effects:** None	FSC-UK EC Certificate (bsi)	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
31.	**ICIM INTERNATIONAL S.R.L.,** located in Via Peloritana, 28, 20024 Garbagnate Milanese (Milano), Italy. **Local agent:** Radiant Export Import Enterprise. Uttara, Dhaka.	Bionike PROCUTASE Ionic Hydrogel SPRAY for Skin Lesion Dressing Spray, 100 ml/bottle **Class: B**	Natural hydrophilic polymers; a patented solution of ionized trace metals; Metalloproteinases inhibiting plant peptides (TIMPS)	It is used for dressing skin lesions like abrasions, scalds and burns, graze and wounds, vascular and decubitus ulcers. It is highly effective in controlling the lession humid microenvironment while encouraging fibroblast proliferation as well as the healing process.	**Contraindications:** Not applicable **Side effects:** No side effects are known that relate specifically to the use of the product. The product is usually well tolerated. In the event of individual hypersensitivity to any of the specific components, avoid using the product. In case of intolerance to the product, stop the treatment and consult your doctor. Consult your doctor if the lesions do not heal normally.	FSC-Italy	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

32.	**ICIM INTERNATIONAL S.R.L.,** located in Via Peloritana, 28, 20024 Garbagnate Milanese (Milano), Italy. **Local agent:** Radiant Export Import Enterprise. Uttara, Dhaka.	Bionike PROCUTASE Ionic Hydrogel for Skin Lesion Dressing Gel 50 ml tube with applicator **Class: B**	Natural hydrophilic polymers; a patented solution of ionized trace metals; Metalloproteinases inhibiting plant peptides (TIMPS)	It is used for dressing skin lesions like abrasions, scalds and burns, graze and wounds, vascular and decubitus ulcers. It is highly effective in controlling the lession humid microenvironment while encouraging fibroblast proliferation as well as the healing process.	**Contraindications:** Not applicable **Side effects:** No side effects are known that relate specifically to the use of the product. The product is usually well tolerated. In the event of individual hypersensitivity to any of the specific components, avoid using the product. In case of intolerance to the product, stop the treatment and consult your doctor. Consult your doctor if the lesions do not heal normally.	FSC-Italy	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj
33.	**Manufacturer :** Smith & Nephew Inc., 101 Hessle Road, Hull, HU3 2BN, UK **Local Agent :** Vastech Limited, Nurjehan Tower (6th Floor) 80/22 Mymensingh Road Dhaka-1000. Bangladesh	Primapore **Class: B**	Non-woven adhesive wound dressing	It is indicated for post-operative wounds, and cuts, lacerations and sutured wounds.	**Contra-Indication:** None **Side Effects:** None	FSC-UK EC Certificate (bsi)	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj

34.	**Manufacturer :** Smith & Nephew Inc., 101 Hessle Road, Hull, HU3 2BN, UK **Local Agent :** Vastech Limited, Nurjehan Tower (6th Floor) 80/22 Mymensingh Road Dhaka-1000. Bangladesh	Allevyn Ag Gentle Border **Class: B**	Silicone Gel Adhesive Antimicrobial Hydrocellular foam dressing	ALLEVYN Ag Gentle Border is an Absorbent antimicrobial dressing for the management, by secondary intention of chronic and acute full thickness, partial thickness or shallow granulating, exuding wounds such as: pressure ulcers, venous ulcers, diabetic ulcers, burns, donor sites, fungating / malignant wounds and surgically dehisced wounds. ALLEVYN Ag Gentle Border may be used on infected wounds. Where the product is used on infected wounds the wounds should be treated as per local protocol. ALLEVYN Ag Gentle Border can be used in conjunction with INTRASITE™ Gel for necrotic or sloughy wounds. ALLEVYN Ag Gentle Border is suitable for use on fragile skin.	**Contra-indications** Do not use on patients known to be hypersensitive to silver sulfadiazine or sulfonamides. • As sulfonamides are known to cause kernicterus, ALLEVYN Ag Gentle Border should not be used on females who are at, or near term pregnancy or lactating, on premature infants or on newborn infants during the first months of life. **Side Effects:** None	FSC-UK EC Certificate (bsi)	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

35.	**Manufacturer :** Smith & Nephew Inc., 101 Hessle Road, Hull, HU3 2BN, UK **Local Agent :** Vastech Limited, Nurjehan Tower (6th Floor) 80/22 Mymensingh Road Dhaka-1000. Bangladesh	Cica-Care **Class: B**	Silicone Gel sheet	It is designed for temporary use: 1. in the management of both existing and new hypertrophic scars and keloids and 2. as a prophylactic therapy on closed wounds to help to prevent hypertrophic scarring and keloids	**Contra-indications:** Do not use on patients with complicating medical factors which would make them unable to use the gel sheet properly. Do not use on patients with dermatological conditions (e.g. Psoriasis & Acne). DO NOT USE ON OPEN WOUNDS or where the skin integrity is compromised. **Side Effects:** None	FSC-UK EC Certificate (bsi)	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj
36.	**Manufacturer :** Smith & Nephew Inc., 101 Hessle Road, Hull, HU3 2BN, UK **Local Agent :** Vastech Limited, Nurjehan Tower (6th Floor) 80/22 Mymensingh Road Dhaka-1000. Bangladesh	Opsite Incise **Class: B**	Transparent adhesive film	It is indicated to dress minor burns, scalds, skin grafts and donor sites, pressure sores, abrasions and Lacerations; for the prophylaxis of pressure sores; for stoma and fistula therapy; for use in all types of surgery; IV fixation (central peripheral or venous catheters).	**Contra-Indication:** None **Side Effects:** None	FSC-UK EC Certificate (bsi)	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj

37.	**Manufacturer :** Boston Scientific Corporation, 300 Boston Scientific Way, USA **Local Agent :** Vastech Limited, Nurjehan Tower (6th Floor) 80/22 Mymensingh Road Dhaka-1000. Bangladesh.	Speed Band Superview Super 7 **Class: B**	Endoscopic Ligator	The Speedband Superview Super 7™ Multiple Band Ligator is used for endoscopic ligation of esophageal varices and anorectal hemorrhoids.	**Contraindications:** The Speed band Superview Super 7 Multiple Band Ligator is contraindicated for patients with bleeding disorders, unless the bleeding disorder is first identified and treated appropriately, and is contraindicated for any other condition which would otherwise contraindicate gastrointestinal endoscopy. The Speedband Superview Super 7 Multiple Band Ligator is not intended for ligation of esophageal varices below the gastroesophageal junction. **Adverse Events:** Of Esophageal Variceal Ligation • Esophageal ulceration. • Retrosternal chest pain secondary to initial banding or ulceration at banding sites. • Treatment-related bleeding secondary to ulceration at banding sites. • Esophageal stricture formation. • Esophageal perforation. • Esophageal obstruction. Of Anorectal Hemorrhoidal Ligation Severe pain, which may result from treatment of hemorrhoids below the dentate line	Certificate to foreign Government-USA EC Certificate (bsi)	Abygv`b Kiv †h‡Z cv‡i |	Abygv`b Kiv nj |
38.	Romsons Scientific & Surgical Industries Pvt. Ltd, India **Importer:** Barisal Surgical, Dhaka	Endotracheal Tube Cuffed **Class: B**	Oral/Nasal Tracheal Tube with cuff & with Murphy eye & Radio opaque Line, pilot Balloon	Endotracheal Tube- Cuffed intended to provide a direct & uninterrupted to & fro airway Passage to lungs. (A special soft, high volume, low pressure cuff is provided for air tight seal between the tube and the tracheal wall.	Contraindication: None Side-effect : None	FSC-India EC Certificate-Norway	Abygv`b Kiv †h‡Z cv‡i |	Abygv`b Kiv nj |

39.	Romsons Scientific & Surgical Industries Pvt. Ltd, India **Importer:** Barisal Surgical, Dhaka	Endotracheal Tube Plain **Class: B**	Oral/Nasal Tracheal Tube without cuff & with Murphy eye & Radio opaque Line	Endotracheal Tube is intended to provide a direct & uninterrupted to & fro airway to lung.	Contraindication: None Side-effect : None	FSC-India EC Certificate-Norway	Abjgv`b Kiv †h‡Z cv‡i |	Abjgv`b Kiv nj |
40.	Romsons International, India **Importer:** Barisal Surgical, 34/1, Mitford Road, Dhaka-1100	VEIN-O-LINE **Class: B**	Large Bore, Low Pressure, Extension Line with Male luer Lock & 3-Way stop Cock at ends	Device intended for administration of intravenous infusions & nutrition etc. 3 Way Stop Cock facilitates selective running of multiple infusion lines.	Contraindication: None Side-effect : None	FSC-India EC Certificate-Norway	Abjgv`b Kiv †h‡Z cv‡i |	Abjgv`b Kiv nj |
41.	Romsons International, India **Importer:** Barisal Surgical, 34/1, Mitford Road, Dhaka-1100	PM-O-Line **Class: B**	Pressure monitoring line	It is single use device, used for administration of intravenous fluids in critical care for low prime volume high pressure applications.	Contraindication: None Side-effect : None	FSC-India EC Certificate-Norway	Abjgv`b Kiv †h‡Z cv‡i |	Abjgv`b Kiv nj |
42.	Romsons Scientific & Surgical Industries Pvt. Ltd, India **Importer:** Barisal Surgical, Dhaka	IRRIGATTO **Class: B**	Syringe with catheter mount suitable for irrigation and feeding	IRRIGATTO Syringe with catheter mount nozzle is used for irrigation and feeding.	Contraindication: None Side-effect : None	FSC-India EC Certificate-Norway	Abjgv`b Kiv †h‡Z cv‡i |	Abjgv`b Kiv nj |
43.	Romsons Scientific & Surgical Industries Pvt. Ltd, India **Importer:** Barisal Surgical, Dhaka	Romolene **Class: B**	Ryle's Tube with funnel & Luer connector	Intended for naso-gastric introduction, for Enteral Feeding as well as for aspiration of gastro-Intestinal secretions.	Contraindication: None Side-effect : None	FSC-India EC Certificate-Norway	Abjgv`b Kiv †h‡Z cv‡i |	Abjgv`b Kiv nj |

44.	Ashitaka Factory of Terumo Corporation , Japan **Local Agent**: UniMed Ltd. (Medical Device) 34/1 Sonagaon Road, Paribag, Dhaka	Capiox Hemoconcentrator **Class: B**	Hemoconcentration Filter	It is intended to be used during and after surgical procedures requiring cardiopulmonary bypass (up to 6 hours) when the removal of excess fluid from blood is required. It should not be used as dialyzer, hemofilter or other device.	**Contraindication:** There are no known absolute Contraindication for ultrafiltration therapy; ultrafiltration can affect the effective concentration and clearance of concoomitant drugs and medication. Use of drugs and medications during ultrafiltration therapy must beclosely monitored by the prescribing physician. **Side effect :** None	FSC-Japan EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
45.	**Manufacturer :** Smith & Nephew Inc., 101 Hessle Road, Hull, HU3 2BN, UK **Local Agent :** Vastech Limited, Nurjehan Tower (6th Floor) 80/22 Mymensingh Road Dhaka-1000. Bangladesh	Endobutton **Class: C**	Fixation Button	The fixation device is used for fixation of tendons and ligaments during orthopedic reconstruction procedures such as anterior cruciate ligament (ACL) reconstruction.	**Contraindication:** Known hypersensitivity to the implant material. Where material sensitivity is suspected, appropriate tests should be made and sensitivity ruled out prior to implantation Insufficient quantity or quality of bone. Blood supply and previous infections which may tend to retard healing. Active infection. Conditions which tend to limit the patient's ability or willingness to restrict activities or follow directions during the heating period. **Adverse Reactions:** Complications are those seen with any method of internal fixation. Adverse reactions associated with suture include: wound dehiscence, calculi formation in urinary or biliary tract such as bile or urine occurs, infected wounds, minimal acute inflammatory tissue reaction and transitory local irritation.	Certificate to Foreing Gov.-USA EC Certificate (bsi)	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

46.	**Manufacturer :** Smith & Nephew Inc., 101 Hessle Road, Hull, HU3 2BN, UK **Local Agent :** Vastech Limited, Nurjehan Tower (6th Floor) 80/22 Mymensingh Road Dhaka-1000. Bangladesh	BIORCI Scrue **Class: C**	Fixation Screws	The Smith & Nephew BIORCI Screw is indicated for interference fixation of bone-tendon-bone or soft tissue grafts in anterior/posterior cruciate ligament ACL/PCLI reconstruction procedures.	**Contra-Indication:** Known hypersensitivity to the implant material. Where material sensitivity is suspected, appropriate tests should be made and sensitivity ruled out prior to implantation. . Conditions that would reduce the support of the screw threads, e.g., insufficient quantity or quality of bone including tumors and severe osteoporosis. The presence of infection. Conditions which would limit the patient's ability or willingness to restrict activities or follow directions during the healing period. o Contraindications may be relative or absolute and must be carefully weighed against the patient's entire evaluation. **Adverse Reactions:** Complications which are seen with any method of internal fixation include failure to regain full extension or flexion, patela femoral complications, fixation complications, hardware irritation, impingement to the graft, and arthrofibrosis. Additional complications may include fixation failure, and migration of the screw. As with any bio-absorbable implant, there is a chance of an inflammatory response during the degradation period of the device.	Certificate to Foreing Gov.- USA EC Certificate (bsi)	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

47.	**Manufacturer :** Smith & Nephew Inc., 101 Hessle Road, Hull, HU3 2BN, UK **Local Agent :** Vastech Limited, Nurjehan Tower (6th Floor) 80/22 Mymensingh Road Dhaka-1000. Bangladesh	RCI Fixation Screw **Class: C**	Fixation Screws	Smith & Nephew RCI Fixation Screws are used for interference fixation of bone-tendon-bone or soft tissue grafts in anterior or posterior cruciate [ligament reconstruction'	**Contra-Indication:** Known hypersensitivity to the implant material. Where material sensitivity is suspected, appropriate tests should be made and sensitivity ruled out prior to implantation. Conditions that would reduce the support of the screw threads; e.g. insufficient quantity or quality of bone including tumors and severe Osteoporosis. The presence d infection Mental or neurological conditions which would reduce compliance with the restrictions and demands of the rehabilitation program, especially during the first two weeks; e.g. drug use or mental illness. **Adverse Reactions:**. Complications are those seen with any method of internal fixation. Laceration of the graft material This may occur as a result of poor technique in the placement of the screws. It is more likely to result from operative instruments than from the screws. . Graft material may fail to incorporate into the tunnel following anterior cruciate reconstruction and, white this is a rare occurrence, this may lead to the dissolution of the graft material and recurrent instability of the knee'	Certificate to Foreing Gov.- USA EC Certificate (bsi)	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
48.	Romsons International, India **Importer:** Barisal Surgical, 34/1, Mitford Road, Dhaka-1100	Romo Vac Set **Class: B**	Closed wound suction set	Romo Vac Set is intended for close wound drainage under negative pressure with the option to use one perforated to catheter or two catheters simultaneously to facilitate desired drainage.	Contraindication: None Side-effect : None	FSC-India EC Certificate-Norway	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
49.	Romsons International, India **Importer:** Barisal Surgical, 34/1, Mitford Road, Dhaka-1100	ROMO SEAL (ADULT, MIDI, & KID) **Class: B**	Under water seal drainage system	Romo Seal is sterile, single, use device intended for under water seal chest drainage application after cardiac and cardio-thoracic procedures.	Contraindication: None Side-effect : None	FSC-India EC Certificate-Norway	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

50.	Romsons Scientific & Surgical Industries Pvt. Ltd, India **Importer:** Barisal Surgical, Dhaka	Romo Drain 1000ml **Class: B**	Under water seal chest drainage system	It is an under water seal drainage bag intended for use in pleural cavity/thoracic drainage	Contraindication: None Side-effect : None	FSC-India EC Certificate-Norway	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |
51.	**Manufacturer :** Boston Scientific Corporation, 300 Boston Scientific Way, USA **Local Agent :** Vastech Limited, Nurjehan Tower (6th Floor) 80/22 Mymensingh Road Dhaka-1000. Bangladesh.	**EndoVive** Percutaneous Endoscopic Gastrostomy Kit **Class: C**	Percutaneous Endoscopic Gastrostomy Kit	The Initial Placement PEG is indicated for enteral nutrition directly into the stomach in both pediatric and adult patients who are unable to consume nutrition by conventional means.	**Contraindications:** • Patient on anticoagulant drugs. • Obstruction of the esophagus which may prevent the introduction or removal of the feeding tube. • Inability to achieve transabdominal illumination or needle/cannula placement. • Multiple surgical procedures near the gastrostomy site. • High medical risk patients. **ADVERSE EVENTS:** Possible complications include, but may not be limited to: fever, gastric distention, infection, blockage/occlusion, tissue necrosis, migration, peritonitis, sepsis, erosion/embedding into the gastric wall ("Buried Bumper Syndrome"), aspiration, bleeding, fistula, gastroparesis, GE reflux, pain, perforation, ulceration, tube clogging, malposition, leakage, kinking, inadvertent removal, small bowel obstruction, granulation tissue, and pneumoperitoneum.	Certificate to foreign Government-USA EC Certificate (bsi)	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |

52.	GELITA MDICAL GmbH: Uferstrasse 7 69412 Eberbach Germany [**SURGITA LTD**. : 58/3, Lake Circus, Kalabagan, Dhanmondi, Dhaka – 1205, Bangladesh]	**GELITA-CEL® STANDARD** **Class: D**	Absorbable Oxidized Cellulose Gauze Hemostat	Topical hemostat for use as an adjunct to hemostasis by tamponade effect, in particular where control of capillary, venous, and arteriolar bleeding, by pressure, ligature, and other conventional procedures, is either ineffective or impractical.	**Contraindication:** It should not be used for implantation in bone defects, such as fractures, since there is a possibility of interference with callus formation and a theoretical chance of cyst formation. It should not be used in conjunction with methyl methacrylate adhesives, for example in orthopedic surgery, because their presence may reduce the adhesive strength of the bonding agent to bone. It should not be used in bleeding from large arteries. It should not be used in chemically cauterized areas; its use should not be preceded by application of escharotic chemicals (silver nitrate or any other). It should not be used as a surface dressing except for immediate control of bleeding as it inhibits new skin growth. It should not be used in closure of skin incisions as it may interfere with healing of the skin edges. This is due to mechanical interposition of oxidized cellulose and not to intrinsic interference with wound healing. **Side Effects:** Occasional reports of "burning" and "stinging" sensations and sneezing when oxidized cellulose has been used as packing in epistaxis, are believed to be due to the low pH of the product. Burning has been reported when oxidized cellulose was applied after nasal polyp removal and after hemorrhoidectomy. Stinging has also been reported where oxidized cellulose was applied on surface wounds (varicose ulcerations, dermabrasions, and donor sites).	FSC- Germany CE Marking of Conformity EC Certificate (Germany)	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

53.	GELITA MDICAL GmbH: Uferstrasse 7 69412 Eberbach Germany [**SURGITA LTD**. : 58/3, Lake Circus, Kalabagan, Dhanmondi, Dhaka – 1205, Bangladesh]	**GELITA-CEL® X-SORB** **Class: D**	Absorbable Oxidized Cellulose Gauze Hemostat for Heavier Bleeding	Topical hemostat for use as an adjunct to hemostasis by tamponade effect, in particular where control of capillary, venous, and arteriolar bleeding, by pressure, ligature, and other conventional procedures, is either ineffective or impractical.	**Contraindication:** It should not be used for implantation in bone defects, such as fractures, since there is a possibility of interference with callus formationand a theoretical chance of cyst formation. It should not be used in conjunction with methyl methacrylate adhesives, for example in orthopedic surgery, because their presencemay reduce the adhesive strength of the bonding agent to bone. It should not be used in bleeding from large arteries. It should not be used in chemically cauterized areas; its use should not be preceded by application of escharotic chemicals (silver nitrate or any other). It should not be used as a surface dressing except for immediate control of bleeding as it inhibits new skin growth. It should not be used in closure of skin incisions as it may interfere with healing of the skin edges. This is due to mechanical interposition of oxidized cellulose and not to intrinsic interference with wound healing. **Side Effects:** Occasional reports of "burning" and "stinging" sensations and sneezing when oxidized cellulose has been used as packing in epistaxis, are believed to be due to the low pH of the product. Burning has been reported when oxidized cellulose was applied after nasal polyp removal and after hemorrhoidectomy. Stinging has also been reported where oxidized cellulose was applied on surface wounds (varicose ulcerations, dermabrasions, and donor sites).	FSC- Germany CE Marking of Conformity EC Certificate (Germany)	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

54.	GELITA MDICAL GmbH: Uferstrasse 7 69412 Eberbach Germany [**SURGITA LTD**. : 58/3, Lake Circus, Kalabagan, Dhanmondi, Dhaka – 1205, Bangladesh]	**GELITA-CEL® FIBRILLAR** **Class: D**	Absorbable Oxidized Cellulose Non-Woven Hemostat	Topical hemostat for use as an adjunct to hemostasis by tamponade effect, in particular where control of capillary, venous, and arteriolar bleeding, by pressure, ligature, and other conventional procedures, is either ineffective or impractical.	**Contraindication:** It should not be used for implantation in bone defects, such as fractures, since there is a possibility of interference with callus formation and a theoretical chance of cyst formation. It should not be used in conjunction with methyl methacrylate adhesives, for example in orthopedic surgery, because their presencemay reduce the adhesive strength of the bonding agent to bone. It should not be used in bleeding from large arteries. It should not be used in chemically cauterized areas; its use should not be preceded by application of escharotic chemicals (silver nitrate or any other). It should not be used as a surface dressing except for immediate control of bleeding as it inhibits new skin growth. It should not be used in closure of skin incisions as it may interfere with healing of the skin edges. This is due to mechanicalinterposition of oxidized cellulose and not to intrinsic interference with wound healing. **Side Effects:** Occasional reports of “burning” and “stinging” sensations and sneezing when oxidized cellulose has been used as packing in epistaxis, are believed to be due to the low pH of the product. Burning has been reported when oxidized cellulose was applied after nasal polyp removal and after hemorrhoidectomy.Stinging has also been reported where oxidized cellulose was applied on surface wounds (varicose ulcerations, dermabrasions, and donor sites).	FSC- Germany CE Marking of Conformity EC Certificate (Germany)	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj

55.	GELITA MDICAL GmbH: Uferstrasse 7 69412 Eberbach Germany [**SURGITA LTD**. : 58/3, Lake Circus, Kalabagan, Dhanmondi, Dhaka – 1205, Bangladesh]	**GELITA-SPON ® POWDER** **Class: D**	Absorbable Gelatin Powder Hemostat	Topical hemostat for use as an adjunct to hemostasis by tamponade effect, in particular where control of capillary, venous, and arteriolar bleeding, by pressure, ligature, and other conventional procedures, is either ineffective or impractical. It is particularly useful in procedures involving cancellous bone bleeding and can be used in the same way and for the same indications as would a “bone wax”. The paste having been mixed to the required consistency may be smeared or pressed against the bleeding osseous surface to control bleeding. Due to its complete bio degradation, it is not known to cause sternal dehiscence unlike conventional bone waxes. It can also be mixed to a “slurry” consistency and administered via a syringe in difficult to reach areas, e.g. in spinal surgeries, during laminectomy and discectomy, or spinal fusion to stop capillary bleeding. It used dry, as a paste or more viscous fluid, has clinical utility in broad surface bleeding, cavity bleeding and uneven surface bleeding.	**Contraindication:** Hypersensitivity to porcine products. It should not be used in closure of skin incisions as it may interfere with healing of the skin edges. This is due to mechanicalinterposition of gelatin and not to intrinsic interference with wound healing.The product should not be used without antibiotics in infected wounds. **Side Effects:** There have been no reported adverse reactions for product used correctly according to these Instructions for Use and when the product is not over-packed.Formation of tissue granulation during middle ear procedures has been reported in at least one animal study.	FSC- Germany CE Marking of Conformity	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj

56.	GELITA MDICAL GmbH: Uferstrasse 7 69412 Eberbach Germany [**SURGITA LTD**. : 58/3, Lake Circus, Kalabagan, Dhanmondi, Dhaka – 1205, Bangladesh]	**GELITA-SPON ® STANDARD** **Class: D**	Absorbable Gelatin Sponge Hemostat	Topical hemostat for use as an adjunct to hemostasis by tamponade effect, in particular where control of capillary, venous, and arteriolar bleeding, by pressure, ligature, and other conventional procedures, is either ineffective or impractical. 2.1 In Dental surgery, the GELITA-SPON® STANDARD Cube is an aid in providing hemostasis and filling dead space created by extraction of the teeth, rootamputations and removal of cysts, tumors and impacted teeth. 2.2 For ENT surgery a "High Density" sponge is available. It has proven to be effective in supporting and keeping the fascia or perichondrium in the middleear (dry application) in place or in the outer ear canal after tympanoplasty. 2.3 For rectal surgery, hemorrhoid operations and gynaecology the GELITA-SPON® STANDARD"Tampon" form is available.	**Contraindication:** Hypersensitivity to porcine products. It should not be used in closure of skin incisions as it may interfere with healing of the skin edges. This is due to mechanicalinterposition of gelatin and not to intrinsic interference with wound healing.The product should not be used without antibiotics in infected wounds. **Side Effects:** There have been no reported adverse reactions for product used correctly according to these Instructions for Use and when the product is not over-packed. Formation of tissue granulation during middle ear procedures has been reported in at least one animal study.	FSC- Germany CE Marking of Conformity	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

57.	GELITA MDICAL GmbH: Uferstrasse 7 69412 Eberbach Germany [**SURGITA LTD**. : 58/3, Lake Circus, Kalabagan, Dhanmondi, Dhaka – 1205, Bangladesh]	**GELITA-SPON ® RAPID³** **Class: D**	Absorbable Gelatin Sponge Hemostat	Topical hemostat for use as an adjunct to hemostasis by tamponade effect, in particular where control of capillary, venous, and arteriolar bleeding, by pressure, ligature, and other conventional procedures, is either ineffective or impractical.	**Contraindication:** Hypersensitivity to porcine products. • It should not be used in closure of skin incisions as it may interfere with healing of the skin edges. This is dueto mechanical interposition of gelatin and not to intrinsic interference with wound healing. The product should not be used without antibiotics in infected wounds. **Side Effects:** There have been no reported adverse reactions for product used correctly according to these Instructions for Use and when overpackingis not applied.Formation of tissue granulation during middle ear procedures has been reported in at least one animal study.	FSC- Germany CE Marking of Conformity	Abjgv`b Kiv †h‡Z cv‡i |	Abjgv`b Kiv nj |
58.	BIOTRINIK AG Ackerstrasse 6 CH-8180 Bulach Switzerland (**Cardiac Solution** Ltd.House #1/1/B,Flat# 1/C,SiliconArcade, Ring Road, Adabor, Dhaka)	Cruiser Hydro **Class: B**	Coronary and Peripheral Artery Guide Wire	Use for Coronary Artery	**Contraindication:** None **Side Effects:** None	FSC-Switzerland EC Design examination Certificate(bsi)	Abjgv`b Kiv †h‡Z cv‡i |	Abjgv`b Kiv nj |
59.	**Manufacturer :** Boston Scientific Corporation, 300 Boston Scientific Way, USA **Local Agent :** Vastech Limited, Nurjehan Tower (6th Floor) 80/22 Mymensingh Road Dhaka-1000. Bangladesh.	Hydra Jagwire High Performance Guide wire **Class: B**	Endoscopy Guide wires	It is indicated for use in selective cannulation of the biliary ducts including the common bile, pancreatic, cystic, right and left hepatic ducts and to aid in the placement of diagnostic and therapeutic devices during bronchoscopic procedures.	**Contraindications:** None known. **Adverse events:** Use of guide wires in the gastrointestinal tract and the tracheobronchial tree may be associated with the following complications: Infection, Bleeding, Breakage with retention of fragment, Perforation, Peritonitis, Inflammation, Edema, Hemorrhage, Pancreatitis, Tissue Trauma, Wire Fracture, Failure to Pass, Hemothorax, Pneumothorax, Foreign Object in Body, Septicemia	Certificate to foreign Government-USA EC Certificate (bsi)	Abjgv`b Kiv †h‡Z cv‡i |	Abjgv`b Kiv nj |

60.	Ashitaka Factory of Terumo Corporation , Japan **Local Agent**: UniMed Ltd. (Medical Device) **34/1 Sonagaon Road, Paribag, Dhaka**	TR Brand (Sterile) **Class: B**	Radial Artery Hemostasis Band	It is a compression device to assist hemostasis of the radial artery after a transradial procedure.	**Contra-indication :** None **Side effect :** None	FSC-Japan	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj	
61.	B L Lifesciences Pvt Ltd., India [**Inogen Systems**, Cha 73/2, Pragati Sarani, North Badda, Badda, Dhaka-1212]	**Perfx-** Heart Lung Pack **Class: D**	Disposable Perfusion Set	This set is intended for use during surgery requiring cardiopulmonary or other surgical techniques. It can be used together with other devices such as pumps, oxygenators, reservoirs, filters, heat exchangers and cannulae. It is suggested to consult the specific instructions for use of the above listed devices. The blood to be treated must contain anticoagulant. The device must not be used for longer than 6 hours. Contact with blood for longer periods is not advisable.	**Contraindication:** None **SideEffects:** None	FSC-India CE Mark-Norway	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj	
62.	Terumo Medical Products(Hangzhou)Co. Ltd., China **Local Agent:** UniMed Ltd. (Medical Device), 34/1 Sonagaon Road, Paribag, Dhaka	**Terufuion** Solution Administration Set for Infusion Pump **Class: B**	Solution Administration Set	Solution Administration Set for Infusion Pump	**Contraindication:** None **Side Effects:** None	FSC-China EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj	

63.	B L Lifesciences Pvt Ltd., India [**Inogen Systems**, Cha 73/2, Pragati Sarani, North Badda, Badda, Dhaka-1212]	**Perfx-** Perfusion Accessories **Class: D** **(Component:** Arterial Filter- Adult/ Pediatric/ Semi Adult, Hemoconcentrator Kit- Adult/ Pediatric, Suction Tubing, HLP Connector- Straight/Y With/ Without Luer Lock, Tourniquet Set- Adult/ Pediatric, Mister Blower- With/ Without Handle, Purge Line, Gas Filter, BT Filter Set, Heat Exchanger- With/ Without aluminium spiral)	Disposable Perfusion Set	This set is intended for use during surgery requiring cardiopulmonary or other surgical techniques. It can be used together with other devices such as pumps, oxygenators, reservoirs, filters, heat exchangers and cannulae. It is suggested to consult the specific instructions for use of the above listed devices. The blood to be treated must contain anticoagulant. The device must not be used for longer than 6 hours. Contact with blood for longer periods is not advisable.	**Contraindication:** None **SideEffects:** None	FSC-India CE Mark-Norway	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj
64.	**Manufacturer :** Boston Scientific Corporation, 300 Boston Scientific Way, USA **Local Agent :** Vastech Limited, Nurjehan Tower (6th Floor) 80/22 Mymensingh Road Dhaka-1000. Bangladesh.	Ultratome **Class: B**	Sphincterotomes	It is indicated for use in the selective cannulation of the Common Bile Ducts (CBD) and the transendoscopic sphincterotomy of the Papilla of Vater and/or the Sphincter of Oddi. The Sphincterotome can also be used to inject contrast medium.	**Contraindications:** Contraindications for this device are those specific to endoscopic retrograde cholangiopancreatography (ERCP) and endoscopic sphincterotomy (ES). **Adverse Effect:** Potential Complications include, but are not limited to: Pancreatitis; Perforation; Hemorrhage; Hematoma; Cholangitis; Stone Impaction; Septicemia/Infection; Allergic Reaction to Contrast Medium. Possible electrosurgical adverse effects include: Fulguration; Burns; Stimulation; Cardiac Arrhythmias.	Certificate to foreign Government-USA EC Certificate (bsi)	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj

65.	Romsons Scientific & Surgical Industries Pvt. Ltd, India **Importer:** Barisal Surgical, Dhaka	T.U.R. Set **Class: B**	Irrigation set for endoscopic Trans urethral Resection of Prostrate (T.U.R.P.)	Sterile, Single Use device intended for continuous irrigation during Trans urethral Resection of Prostrate (T.U.R.P.)	Contraindication: None Side-effect : None	FSC-India EC Certificate-Norway	Abygv`b Kiv †h‡Z cv‡i |	Abygv`b Kiv nj |
66.	Romsons Juniors India Unit-II, India **Importer:** Barisal Surgical, Dhaka	**Microguard** Vented infusion set with 0.2µ filter, DEHP free **Class: B**	Sterile disposable perfusion set	It is intended for use in intervenous infusion of I.V. Fluids and parenteral drugs, to Neonates, Paediatrics, Adults in critical care units and for chemo therapy.	Contraindication: None Side-effect : None	FSC-India EC Certificate-Norway	Abygv`b Kiv †h‡Z cv‡i |	Abygv`b Kiv nj |
67.	Bausch & Lomb,INC 1400,North Goodman Street,Rochester,NY, USA. <u>Local agent :</u> Janata Traders.	Akreos **Class: C**	Advanced Optics Aspheric Lens (Intraocular Lens)	Intraocular Lens	Contraindication: None Side effect: None	CFG-USFDA	Abygv`b Kiv †h‡Z cv‡i |	Abygv`b Kiv nj |
68.	Eyekon Medical Inc. 2451 Enterprise Rd. Clearwater , Florida, USA 33763. **<u>Local agent:</u>** Zas Corporation 80/22, Mymensingh Road, Banglamotor, Dhaka-1000	DGR Hydrophilic Acrylic **Class: C**	Hydrophilic Sterile Intra Ocular Lenses	Optical implant for the replacement of human lens in surgical treatment of cataract	**Contraindication:** Significant complications during the cataract extraction procedure. **Side Effect:** Glare or halos around lights, or decreased sharpness of vision (contrast sensitivity) may occur, especially at night or in dim light	Certificate to Foreign Gov. – USA EC Certificate (UK)	Abygv`b Kiv †h‡Z cv‡i |	Abygv`b Kiv nj |
69.	Bausch & Lomb,INC 1400,North Goodman Street,Rochester,NY, USA. <u>Local agent :</u> Janata Traders.	EnVista **Class: C**	Hydrophobic Acrylic Intraocular Lens	Indicated for primary implantation for the visual correction of aphakia in adult patients in whom the cataractous lens has been removed. The lens is intended for placement in the capsular bag.	Contraindication: None Side effect: None	CFG-USFDA	Abygv`b Kiv †h‡Z cv‡i |	Abygv`b Kiv nj |

70.	Eyekon Medical Inc. 2451 Enterprise Rd. Clearwater , Florida, USA 33763. **Local agent:** Zas Corporation 80/22, Mymensingh Road, Banglamotor, Dhaka-1000	DGR Hydrophobic Acrylic **Class: C**	Hydrophobic Sterile Intra Ocular Lenses	Optical implant for the replacement of human lens in surgical treatment of cataract	**Contraindication:** Significant complications during the cataract extraction procedure. **Side Effect:** Glare or halos around lights, or decreased sharpness of vision (contrast sensitivity) may occur, especially at night or in dim light	Certificate to Foreign Gov. – USA EC Certificate (UK)	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj
71.	Bausch & Lomb,INC 1400,North Goodman Street,Rochester,NY, USA. Local agent : Janata Traders.	**Crystalens** Accommodating Intraocular Lens **Class: C**	Intraocular Lens	It is intended for primary **implantation in the capsular bag of the eye** for the visual correction of aphakia secondary to the removal of cataractous lens in patients with and without presbyopia.The crystalens provides approximately one diaper of monocular accommodation which allows for near, intermediate and distance vision without spectacles.	**Warning:** Careful preoperative evaluation and sound clinical judgment should used by the surgeon to decide the risk/benefit ratio before implanting a lens in a patient. **Side effect:** Hypopyon, intraocular infection ,acute corneal decomposition and secondary surgical intervention.	CFG-USFDA	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj
72.	Bausch & Lomb,INC 1400,North Goodman Street,Rochester,NY, USA Local agent : Janata Traders.	**INCISE** Microincision IOL **Class: C**.	Microincision IOL.	It is designed for microincision cataract Surgery. The Incise Microincision posterior chamber lens is indicated for **primary implantation** for visual correction of aphakia in the adult patients where a cataractous lens has been removed.the lens is design to be a folded prior to insertion in the eye and implantation in the capsular bag.	**Warning:** Physicians considering lens implantation must weigh the risk /benefit ratio i9nh any of the following circumstances. Recurrent severe anterior or posterior segment Inflammation or uveitis .Iris damage, Persistent bleeding, vitreous prolapse or loss .Patient under two years are not suitable candidate for lol. **Side effect:** wound leak , corneal edema,Hyphema, Virtritis,Pupillary membrane ,Virtrious aspiration ,Glucoma.	CFG-USFDA	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj

73.	Bausch & Lomb,INC 1400,North Goodman Street,Rochester,NY, USA. Local agent : Janata Traders.	**Akreos AO** Micro-incision Lens **Class: C**	Microincision Lens.	The Incise Micro incision posterior chamber lens is indicated for primary implantation for visual correction.	**Warning :** Physicians considering lens implantation must weigh the risk /benefit ratio i9nh any of the following circumstances. **Side effect:** wound leak , corneal edema,Hyphema, Virtritis, Pupillary membrane ,Virtrious aspiration ,Glucoma.	CFG-USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
74.	Eyekon Medical Inc. 2451 Enterprise Rd. Clearwater , Florida, USA 33763. **Local agent:** Zas Corporation 80/22, Mymensingh Road, Banglamotor, Dhaka-1000	DGR PMMA **Class: C**	PMMA Sterile Intra Ocular Lenses	Optical implant for the replacement of human lens in surgical treatment of cataract	**Contraindication:** Significant complications during the cataract extraction procedure. **Side Effect:** Glare or halos around lights, or decreased sharpness of vision (contrast sensitivity) may occur, especially at night or in dim light	Certificate to Foreign Gov. – USA EC Certificate (UK)	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
75.	Romsons International, India **Importer:** Barisal Surgical, 34/1, Mitford Road, Dhaka-1100	CVP Manometer **Class: B**	Central Venous Pressure Monitoring Manometer with extension line	Central Venous Pressure Manometer is used for continuous and intermittent monitoring of central venous Pressure during short term infusion administration.	Contraindication: None Side-effect : None	FSC-India EC Certificate-Norway	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
76.	**Legal Manufacturer:** Davol Inc., Sub. C.R. Bard, Inc. 100 Crossings Boule vard warwick, RI, USA **Manufacturing Facility:** BARD SHANNON Lt., San Geronimo Ind. Park, Lot No-1, R-1, KM 79.7, Humacao, USA **Local Agent:** **Unicorn Healthcare Solution** Ltd., Rupayun Karim Tower 14/D, 80 Kakrile, Dhaka-1000	BARD Composix L/P Mesh **Class: C**	Low profile, large pore polypropylene Mesh	It is indicated for use in the reconstruction of soft tissue deficiencies such as for the repair of hernias and chest wall defects.	**Contraindications:** 1. Literature reports there may be a possibility for adhesion formation when the polypropylene is placed in direct contact with the bowel or viscera. 2. Do not use the Bard® Composix™ L/P Mesh in infants or children whereby future growth will be compromised by use of such material. 3. Do not use Bard® Composix™ L/P Mesh for the reconstruction of cardiovascular defects. **Adverse Reactions:** Possible complications include seroma, adhesions, hematomas, inflammation, extrusion, fistula formation and recurrence of the hernia or soft tissue defect.	Certificate to Foreign Government-USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

77.	**Legal Manufacturer:** Davol Inc., Sub. C.R. Bard, Inc. 100 Crossings Boule vard warwick, RI, USA **Manufacturing Facility:** BARD SHANNON Lt., San Geronimo Ind. Park, Lot No-1, R-1, KM 79.7, Humacao, USA **Local Agent:** **Unicorn Healthcare Solution** Ltd., Rupayun Karim Tower 14/D, 80 Kakrile, Dhaka-1000	BARD 3DMax Light Mesh **Class: C**	Monofilament Polypropylene Mesh	It is indicated to reinforce soft tissue where weakness exists, i.e., repair of hernias and chest wall defects.	**Contraindications:** 1. Do not use BARD® mesh in infants or children, whereby future growth will be compromised by use of such mesh material. 2. Literature reports there may be a possibility for adhesion formation when BARD® mesh is placed in direct contact with the bowel or viscera. **Adverse Reactions:** Possible complications include seroma, adhesions, hematoma, infl ammation, extrusion, fi stula formation and recurrence of the hernia or soft tissue defect. Erosion and migration of the mesh have been reported in gastric banding procedures.	Certificate to Foreign Government-USFDA	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj
78.	**Legal Manufacturer:** Davol Inc., Sub. C.R. Bard, Inc. 100 Crossings Boule vard warwick, RI, USA **Manufacturing Facility:** BARD SHANNON Lt., San Geronimo Ind. Park, Lot No-1, R-1, KM 79.7, Humacao, USA **Local Agent:** **Unicorn Healthcare Solution** Ltd., Rupayun Karim Tower 14/D, 80 Kakrile, Dhaka-1000	BARD Mesh **Class: C**	Monofilament Polypropylene Mesh	It is indicated to reinforce soft tissue where weakness exists, i.e., repair of hernias and chest wall defects	**Contraindications:** 1. Do not use the Bard® Mesh in infants or children whereby future growth will be compromised by use of such material. 2. Literature reports there may be a possibility for adhesion formation when the polypropylene is placed in direct contact with the bowel or viscera. **Adverse Reactions:** Possible complications include seroma, adhesions, hematoma, inflammation, extrusion, fistula formation and recurrence of the hernia or soft tissue defect. Erosion and migration of the mesh have been reported in gastric banding procedures.	Certificate to Foreign Government-USFDA	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj

79.	**Legal Manufacturer:** Davol Inc., Sub. C.R. Bard, Inc. 100 Crossings Boule vard warwick, RI, USA **Manufacturing Facility:** BARD SHANNON Lt., San Geronimo Ind. Park, Lot No-1, R-1, KM 79.7, Humacao, USA **Local Agent:** **Unicorn Healthcare Solution** Ltd., Rupayun Karim Tower 14/D, 80 Kakrile, Dhaka-1000	Permafix Fixation System **Class: C**	Delivary system and sample absorbable fasteners It is disposable, single-use system designed to deliver a permanent fastener into tissue or prosthesis during general surgery procedures such as hernia repair. The fistener delivery system consists of'an ergonomic handle with trigger', shaft and penetrating tip. The shaft is available in either a 36cm length for laparoscopic use or a 20 cm length for' open surgical procedures. The device is preloaded with 15 or' 30 permanent fasteners.. Each permanent fastener contains threads for mesh and tissue delivery.	It is indicated for the approximation of soft tissue and fixation of surgical mesh to tissues during open or laparoscopic surgical procedures, such as hernia repair.	**Contraindications:** (1). This device is not intended for use except as Indicated. (2). Do not use this device where hemostasis cannot beverified visually after application. (3). Contraindications associated with laparoscopic and open surgical procedures relative to mesh fixation apply, including but not limited to: • Fixation of vascular or neural structures • Fixation of bone and cartilage .(4) this device should not be used in patients with a known allergy or hypersensitivity to acetal polymers.(5) Carefully inspect the area in the vicinity of the tissue being fastened to avoid inadvertent penetration of underlying structures such as nerves, vessels, viscera or bone. Use of it in the close vicinity of such underlying structures is contraindicated. For reference, the length of the fastener is 6.0 mm, the fastener head is another 0.7 mm (total 6.7 mm). **Adverse Reactions:** Adverse reactions and potential complications associated with fixation devices such as the PermaFix™ Fixation System may include, but are not limited to the following: hemorrhage, pain, edema, and erythema at wound site; septicemia/infection; allergic reaction to acetal; hernia recurrence/wound dehiscence.	Certificate to Foreign Government-USFDA	*Abyţgv`b Kiv †h‡Z cv‡i* |	*Abyţgv`b Kiv nj* |

80.	Romsons International, India **Importer:** Barisal Surgical, 34/1, Mitford Road, Dhaka-1100	Mucus extractor **Class: B**	Mucus Extractor	Mucus Extractor is used for aspiration of secretion from Oropharynx in newly born babies to ensure free airway for un-interrupted respiration	Contraindication: None Side-effect : None	FSC-India EC Certificate-Norway	Abjygv`b Kiv †h‡Z cv‡i	Abjygv`b Kiv nj
81.	Romsons International, India **Importer:** Barisal Surgical, 34/1, Mitford Road, Dhaka-1100	Aeromist **Class: B**	Aerosol mask with Nebulization Chamber	Used for convienient Nebulizer therapy.	Contraindication: None Side-effect : None	FSC-India EC Certificate-Norway	Abjygv`b Kiv †h‡Z cv‡i	Abjygv`b Kiv nj
82.	HONSUN (Nantong) Co. ltd., China **Local Agent:** **Padma Inter Reade,** 52 New Eskaton Road, TMC Bulding, Dhaka	Nebulizer (Ultrasonic and Compressor) **Class: B**	Nebulizer	Use for remove or reduce Cough and cold	Contraindication: None Side-effect : None	FSC-China CE, Germany	Abjygv`b Kiv †h‡Z cv‡i	Abjygv`b Kiv nj
83.	**Manufacturer:** Onbo Electronic (Shenzhen), Co. Ltd, China **Supplier:** **Microlife Corporation** 9F, 431, RuiGang Road, NehHu, Taippei 114, Taiwan, R.O.C **Local agent:** **Transcom Distribution Co.** Ltd 52, Motijheel C/A, Dhaka-1000	Nebulizer **Class: B**	Nebulizer	It is used to deliver medications along the respiratory tract and is indicated to various respiratory problems and diseases such as: Broncho-spasms Chest tightness Excessive and thick mucus secretions Respiratory congestions Pneumonia	**Contra Indication** :In some cases, nebulization is restricted or avoided due to possible untoward results or rather decreased effectiveness such as: • Patients with unstable and increased blood pressure • Individuals with cardiac irritability (may result to dysrhythmias) • Persons with increased pulses • Unconscious patients (inhalation may be done via mask but the therapeutic effect may be significantly low) **Side effect:** Possible effects and reactions after nebulization therapy are as follows: Palpitations, Tremors, Tachycardia, Headache, Nausea, Broncho-spasms (too much ventilation may result or exacerbate Broncho-spasms)	Certificate for Exportation of Medical Products-China	Abjygv`b Kiv †h‡Z cv‡i	Abjygv`b Kiv nj

84.	Pu Yuan Biotech Co., Ltd., Taiwan (Benvue International, 14/1 Joy Tower, Chattesory Road, Joy nagor R/A, Chittagonj	Azmaler Vapour User Portable Nebulizer **Class: B**	Portable Nebulizer	The function of this nebulizer is basically with compressed air to break up medical suspensions, solutions, COPD, other respiratory disorders. After inhalation with the device the drug content (diluents)/liquid breaks up in the small droplets that can easily inhaled and reached to onset of action (Lungs & alveoli). This device is indicated for respiratory complications like asthma, COPD or for those patients where physician recommended nebulization.	**Contraindications:** None **Side Effects:** None	FSC:Taiwan CE: SGS, UK Ltd.	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

85.	GEOTEK MEDIKAL VE SAGLIK HIZMETLERI TIC.SAN.LTD.STI. Lvedik OSB Agac Metal Sitesi 1436. Sokak No: 12 Yenimahalle, Ankara, Turkey (Local Agent: **Visilex Corporation** Section 2, Avenue 3/12 Haziroad, Mirpur, Dhaka)	**Estacore** Automatic Biopsy Needle **Class: B**	Automatic Biopsy Needle	It is a disposable full automatic biopsy device used for histological core samples from soft tissue and organs.	**Contraindications:** This needle should be used by a physician trained in interventional techniques and familiar with the possible side effects, typical findings, limitations, indications and contraindications of biopsy procedures, pyrogenic judgment is required when considering biopsy on patient's with bleeding disorder, or receiving anti-coagulant medications. **Caution:** Ethylen oxide sterilized. Non-pyogenic, Check the package. If package damaged or opened do not use the products. Check the expiry date and the gauge. Store in dy place, between 15-25^{0}C temperatures, product from sun light Possible allergic reactions should be considered. It must be used only in hospitals dry physicals who familiar with the device and procedures **for single patient use** only. Do not attempt to clean or resterilize this protect. Do not use the device under MRI. After use, this product may be potential biohazard. Handle in a manner which will prevent accidental puncture. Dispose in Accordance with applicable laws and regulations.	FSC-Turkey, EC Certificate (Czech Republic)	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
86.	B.Braun Mesulgen AG, Germany. **Local Agent:** Asia Pacific Medicals Ltd.	Intradyne Puncture Needle **Class: B**	Needle for Angiogram	Needle for Angiography which smoothly tapered design for easy introduction of the guide wire up to max. 0.089 mm	Contraindication: None Side-effect : None	FSC- Germany EC Certificate	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
87.	ConvaTec INC 211, Greensboro. NC.USA <u>Local agent :</u> Janata Traders. TCB Bhabon, 1 kawranbazar,Dhaka 1215	Stomahesive Powder **Class: B**	Ostomy Pouch And accessories	Use for skin requiring absorption of moisture and protection from further damage.	Do not use if a known sensitivity to the powder , do not apply if irritation	CFG-USFDA	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj

88.	ConvaTec INC 211, Greensboro. NC.USA <u>Local agent :</u> Janata Traders.	Somahesive paste **Class: B**	Ostomy Pouch And accessories	Protective skin barrier & filler, convenient, highly effective fillerb, sealant and specially design for arond clostomy.ileostomy or urinary stomas.The paste can also be used to protect exposed skin around fistula sites.between the base of stoma and the opening in the skin barrier, and filler for skin fold, uneven skin surfaceand scars.	Do not use if a known sensitivity to the paste, do not apply if irritation.	CFG-USFDA	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
89.	ConvaTec INC 211, Greensboro. NC.USA <u>Local agent :</u> Janata Traders. TCB Bhabon, 1 kawranbazar,Dhaka 1215	Surfit Plus WFR Flex Tan 38 mm, 45mm, 57mm,70mm , Surfit plus WFR Stoma 38mm ,45mm, 57mm, 70mm , With pouch **Class: B**	Ostomy Pouch And accessories	For Management of Stomal output	Care should be excercised when using adhesive products around flush urostomy .and absence of urinary reserver.	CFG-USFDA	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
90.	ConvaTec INC 211, Greensboro. NC.USA <u>Local agent :</u> Janata Traders. TCB Bhabon, 1 kawranbazar,Dhaka 1215	Surfit Natura CVX INS 19x38mm, 22x38mm, 25x38mm, 29x45mm , 32x45mm, 35x45mm, 38x 57mm **Class: B**	Ostomy Pouch And accessories	For Management of Stomal output	Care should be excercised when using adhesive products around flush urostomy .and absence of urinary reserver.	CFG-USFDA	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

91.	ConvaTec INC 211, Greensboro. NC.USA Local agent : Janata Traders. TCB Bhabon, 1 kawranbazar,Dhaka 1215	Surfit Natura Somahesive WFR Flex 38mm,45mm, 57mm, 70mm **Class: B**	Ostomy Pouch And accessories	For Management of Stomal output	Care should be excercised when using adhesive products around flush urostomy .and absence of urinary reserver.	CFG-USFDA	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |
92.	ConvaTec INC 211, Greensboro. NC.USA Local agent : Janata Traders. TCB Bhabon, 1 kawranbazar,Dhaka 1215	Natura + DRN PCH TANWF **Class: B**	Ostomy Pouch And accessories	For Management of Stomal output	Care should be excercised when using adhesive products around flush urostomy .and absence of urinary reserver.	CFG-USFDA	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |
93.	ConvaTec INC 211, Greensboro. NC.USA Local agent : Janata Traders. TCB Bhabon, 1 kawranbazar,Dhaka 1215	Natura WFR S/Hesive 38mm, 45mm,57mm .70mm and D/Hesive **Class: B**	Ostomy Pouch And accessories	For Management of Stomal output	Care should be excercised when using adhesive products around flush urostomy .and absence of urinary reserver.	CFG-USFDA	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |
94.	ConvaTec INC 211, Greensboro. NC.USA Local agent : Janata Traders. TCB Bhabon, 1 kawranbazar,Dhaka 1215	Natura Stoma FLATAMOLD SM 45MM, 57MM, 70 MM **Class: B**	Ostomy Pouch And accessories	For Management of Stomal output	Care should be excercised when using adhesive products around flush urostomy .and absence of urinary reserver.	CFG-USFDA	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |

95.	ConvaTec INC 211, Greensboro. NC.USA <u>Local agent :</u> Janata Traders. TCB Bhabon, 1 kawranbazar,Dhaka 1215	C/Hesive Nat WFR MLD CVX 13/22mm, 22/33mm, 33/57mm **Class: B**	Ostomy Pouch And accessories.	For Management of Stomal output	Care should be excercised when using adhesive products around flush urostomy .and absence of urinary reserver.	CFG-USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
96.	Romsons Scientific & Surgical Industries Pvt. Ltd, India **Importer:** Barisal Surgical, Dhaka	Hi-Mask **Class: B**	High concentration Face mask	Hi-Mask is a single use device equipped with Mask, NRV Reservoir intended for high concentration of oxygen therapy.	Contraindication: None Side-effect : None	FSC-India EC Certificate-Norway	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
97.	Romsons International, India **Importer:** Barisal Surgical, 34/1, Mitford Road, Dhaka-1100	POWER Drool **Class: B**	Nebulizer Chamber with "T" Shaped connector and mouth piece	Used for convenient Nebulizer therapy through Mouth	Contraindication: None Side-effect : None	FSC-India EC Certificate-Norway	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
98.	Romsons International, India **Importer:** Barisal Surgical, 34/1, Mitford Road, Dhaka-1100	Flexi Mask **Class: B**	Oxygen face mask with swivel connector & multi channel tubing	Used for administration of Oxygen.	Contraindication: None Side-effect : None	FSC-India EC Certificate-Norway	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

99.	Ashitaka Factory of Terumo Corporation , Japan **Local Agent**: UniMed Ltd. (Medical Device) **34/1 Sonagaon Road, Paribag, Dhaka**	Capiox SX **Class: C**	Extra-corporeal Membrane Oxygenator [An optional level alarm system, using ultrasonic waves is available for use with the CAPIOX SX]	It is Intended to be used during open heart surgical procedure requiring cardiopulmonary bypass for period up to 6 hours. It is also intended for use in vacuum assisted venous drainage procedure, post-operative chest drainage and autotransfusion procedures to aseptically return the blood to the patient for blood volume replacement.	**Contra-indication :** None **Side effect :** None	FSC-Japan EC Certificate	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
100.	Ashitaka Factory of Terumo Corporation , Japan **Local Agent**: UniMed Ltd. (Medical Device) **34/1 Sonagaon Road, Paribag, Dhaka**	Capiox RX **Class: C**	Extra-corporeal Membrane Oxygenator [Differ in the positioning of the arterial blood port of the oxygenator, the blood contacting surfaces are coated with Xcoating]	It is Intended to be used during open heart surgical procedures to transfer oxygen and remove carbon dioxide from blood and to control the bold temperature during cardiopulmonary bypass for periods up to 6 hours. The patient weight and BSA should be considered upon use. It is also intended for use in vacuum assisted venous drainage procedures.	**Contra-indication :** None **Side effect :** None	Japan EC Certificate	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

101.	**Novo Nordisk A/S** **Device Manufacturing Development** **Brennum Park, 3400 Hilleroed** **Denmark** Local Office: [Novo Nordisk Pharma (Private) Limited Nina Kabbo, Level-9, 227/A, GulshanTejgaon Link Road, Tejgaon, Dhaka-1208, Bangladesh]	NordiPen® 5 **Class: C**	Growth Hormone Pen	It is intended for use with Norditropin®SimpleXx® (Somatropin) 5mg cartridges & NovoFine® Needle.	**Contra Indication & Side-effect:** Not applicable for this product because it is a non-invasive & category C medical device and not directly contact with the body.	FSC-Denmark	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj	
102.	**Novo Nordisk (China) Pharmaceuticals Co. Ltd** **No. 99 Nanhai Road, TEDA, 300457 Tianjin** **P.R. China** Local Office: [Novo Nordisk Pharma (Private) Limited Nina Kabbo, Level-9, 227/A, GulshanTejgaon Link Road, Tejgaon, Dhaka-1208, Bangladesh]	NovoPen® 4 **Class: C**	Insulin Pen	It is a durable handheld injection device intended for the subcutaneous injection of insulin. It will be used to treat diabetes requiring subcutaneously insulin therapy. The key specific claims in relation to the safety and performance of the device are the 1 Unit dosing increments up to 60 Units.	**Contra Indication & Side-effect:** Not applicable for this product because it is a non-invasive & category C medical device and not directly contact with the body.	FSC-Denmark	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj	

103.	**Manufacturer :** Boston Scientific Corporation, 300 Boston Scientific Way, USA **Local Agent :** Vastech Limited, Nurjehan Tower (6th Floor) 80/22 Mymensingh Road Dhaka-1000. Bangladesh.	Wallstent Biliary Metal Stent **Class: C**	Biliary Metal Stent	It is indicated for use in the treatment of biliary strictures produced by malignant neoplasms.	Contraindication: Contraindications associated with the use of the WALLSTENT BILIARY Endoscopic Biliary Endoprosthesis include: • Use of the device in very small intrahepatic ducts. • Stenting of a perforated duct, where leakage from the duct could be exacerbated by the prosthesis and leakage could occur across the uncovered mesh of the stent. • All of the customary contraindications associated with the endoscopic manipulation of 8F (2.7 mm) caliber catheters within the biliary system. **Adverse Effect:** Complications associated with the use of the WALLSTENT BILIARY Endoscopic Biliary Endoprosthesis Partially Covered may include the usual complications reported for conventional biliary stents and endoscopic procedures such as: • Infection or sepsis • Stent misplacement • Stent migration • Stent obstruction secondary to tumor in growth through the stent • Tumor overgrowth at the stent ends • Sludge occlusion • Bile duct perforation or ulceration • Bleeding • Cholangitis • Pancreatitis	Certificate to foreign Government-USA CE Marking of Conformity (DEKRA)	*Abyjgv` b Kiv †h‡Z cv‡i |*	*Abyjgv` b Kiv nj |*

104.	**Manufacturer :** Boston Scientific Corporation, 300 Boston Scientific Way, USA **Local Agent :** Vastech Limited, Nurjehan Tower (6th Floor) 80/22 Mymensingh Road Dhaka-1000. Bangladesh.	Percuflex biliary stent **Class: C**	Biliary Stents (Plastic)	The Biliary Stents is indicated for use in the treatment of biliary strictures.	**Contraindications:** Contraindications associated with the use of the Biliary Stents include**.** **Adverse Events:** Potential complications that may result from a biliary stent placement procedure include, but may not be limited to: • perforation of bile ducts, liver and/or duodenum • hemorrhage • hematoma • septicemia/infection • bile peritonitis • allergic reaction to contrast medium • stent migration Check for proper position of the stent and delivery system using endoscopy and fluoroscopy. Insertion and placement in an improper location may lead to patient injury.	Certificate to foreign Government-USA EC Certificate (bsi)	Abjgv`b Kiv †h‡Z cv‡i |	Abjgv`b Kiv nj |
105.	M/s. Abbott Vascular, Cashel Road, Clonmel, CO. Tipperary, Ireland **Local Agent:** **The Spondon Ltd** 102, Kazi Nazrul Islam Avenue, BSEC Bhabon Lebel-9, Kawran Bazar, Dhaka	**Absorb GT1** Bioresorbable Vascular Scaffold (BVS) System **Class: D**	Bioresorbable Vascular Scaffold (BVS) System	The Absorb GT1 Bioresorbable Vascular Scaffold (BVS) is a temporary scaffold that will fully resorb over time and is indicated for improving coronary luminal diameter in patients with ischemic heart disease due to de novo native coronary artery lesions (length ≤24 mm) with a reference vessel diameter of ≥2.5 mm and ≤3.75 mm.	**Contraindication:** It is contraindication for use in: • Patients in whom antiplatelet and/ or anticoagulant therapy is contraindicated • Patients with a know hypersensitivity or contraindication to aspirin. both heparin and bivalirudin, clopidogrel, ticlopidine, prasugrel, and ticagrelor, everolimus, poly (L-lactide), poly (D, L-lactide), or platinum, or with contrast sensitivity, who cannot be adequately premedicated **Side-effect:** The clinical data exhibits one side effect that may cause late thrombosis formation for less than 1%.	FSC-Ireland EC Design Examination Certificate	Abjgv`b Kiv †h‡Z cv‡i |	Abjgv`b Kiv nj |

106.	SIMEKS TIBBI URUNLER SANAYI Ve TICARET Limited Sirketi, Turkey (**Importer:** Biva International)	**SIMFLEX** Coronary Stent System **Class: D**	Cobalt Chromium Coronary Stent System	It is indicated for improving coronary luminal diameter in patients with symptomatic ischemic heart disease in de novo coronary artery lesion in native coronary arteries with reference vessel diameter of 2.00 mm to 4.50 mm and lesion length 40mm. The SimFlex is intended for use patients eligible for Percutaneous Transluminal Coronary Angioplasty (PTCA)	Contraindication: Use of SimFlex coronary Stent Delivery System is contraindicated in the following patient types: Patients who are not candidates for coronary bypass surgery. Patients who exhibit angiographic evidence of existing thrombus. Patients who are contraindicated for anti-platelet/anit-coagulant therapy, this includes patients who had major surgery, an obstetrical delivery organ biopsy, or puncture of a non-compressible vessel within 4 days of procedure. Also excluded are those patients with a history of gastrointestinal bleeding, recent C.V.A, diabetic hemorrhage retinopathy, or any condition compromised by prolonged anti-coagulant. Pregnant woman or woman of childbearing potential. Patients who have experienced a recent (less then 1 weak) actual myocardial infarction. Patients with diffuse, disease, defined as long segments of abnormal vessel without interposed normal vessel. Transplant Patients **Adverse effects**: Potential adverse events which may be associated with the use of a coronary stent include but are not limited to: Abrupt stent closure, Acute myocardial infarction, Allergic reactions to anti-coagulant and or anti-thrombotic therapy or contrast medium angina, Arrhythmia, including ventricular fibrillation (VF) and ventricular tachycardia (VT), Arterial perforation, Arterial rupture, Arteriovenous fistula, Bleeding complications, Bradycardia, Cardiac tapenade, Carcinogenic hock, Coronary spasm, Coronary or stent embolism, Coronary or stent thrombosis, Death, Dissection of the coronary artery, Drug reaction to antiplatelet agents / anticoagulant agents/ contrast medium, Emboli distal (fair, tissue or thrombotic emboli), Emergency or non-emergent Coronary Artery Bypass Graft Surgery, Entry site complication & Heart Failure.	FSC- Turkey EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

107.	BIOTRINIK AG Ackerstrasse 6 CH-8180 Bulach Switzerland (**Cardiac Solution** Ltd.House #1/1/B,Flat# 1/C,SiliconArcade, Ring Road, Adabor, Dhaka)	Dynamic Renal **Class: C**	CoCr. Renal Stent	Use for KidneyArtery(Blockage)	**Contraindication:** None **SideEffects:** None	Switzerland	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |
108.	Legal Manufacturer: Biosensors Europe SA Rue de Lausanne 29 1110 Morges Switzerland **Manufacturing Facility:** Biosensors Interventional Technologies Pte Ltd 36 Jalan Tukang Singapore **Local Agent:** Omega Health Care 581, Shewrapara Begum Rokeya Sarani, Mirpur, Dhaka - 1216 Bangladesh	**Chroma** Coronary Stent System **Class: D**	Coronary Stent System	It is indicated for improving coronary luminal diameter for the treatment of de novo lessions in native coronary arteries with a reference diameter ranging between 2.25 mm and 4.00 mm. Stents with length 33 mm and 36 mm are only available for artery diameters ranging between 2.5 mm and 3.5 mm	Contraindication: None Side Effects: None	FSC-Switzerland EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |

109	Legal Manufacturer: Biosensors Europe SA Rue de Lausanne 29 1110 Morges Switzerland **Manufacturing Facility:** Biosensors Interventional Technologies Pte Ltd 36 Jalan Tukang Singapore **Local Agent:** Omega Health Care 581, Shewrapara Begum Rokeya Sarani, Mirpur, Dhaka - 1216 Bangladesh	**BioFreedom** Drug Coated Coronary Stent System (BioFreedom DCS) **Class: D**	Drug Coated Coronary Stent System	It is indicated for improving coronary luminal diameter for the treatment of de novo lessions in native coronary arteries with a reference diameter ranging between 2.25 mm and 4.00 mm. Stents with length 33 mm and 36 mm are only available for artery diameters ranging between 2.5 mm and 3.5 mm	Contraindication: None Side Effects: None	Switzerland EC Certificate CE Marking of Conformity	*Abygv`b Kiv †h‡Z cv‡i |*	*Abygv`b Kiv nj |*

110	Legal Manufacturer: Biosensors Europe SA Rue de Lausanne 29 1110 Morges Switzerland **Manufacturing Facility:** Biosensors Interventional Technologies Pte Ltd 36 Jalan Tukang Singapore **Local Agent:** Omega Health Care 581, Shewrapara Begum Rokeya Sarani, Mirpur, Dhaka - 1216 Bangladesh	**BioMatrix NeoFlex** Drug Eluting Coronary Stent System (BioMatrix NeoFlex DES) **Class: D**	Drug Eluting Coronary Stent System	It is indicated for improving coronary luminal diameter for the treatment of de novo lessions in native coronary arteries with a reference diameter ranging between 2.25 mm and 4.00 mm. Stents with length 33 mm and 36 mm are only available for artery diameters ranging between 2.5 mm and 3.5 mm. The BioMatrix NeoFlex DES with stent length up to 28 mm is also indicated for use in patients with: * ST Elevated Myocardial Infrarction (STEMI), *Acute Coronary Syndromes (ACS) including ACS-STEMI, and unstable Angina. * Diabetes Melitus	Contraindication: None Side Effects: None	Switzerland EC Certificate CE marking of Conformity	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

111.	Legal Manufacturer: Biosensors Europe SA Rue de Lausanne 29 1110 Morges Switzerland **Manufacturing Facility:** Biosensors Interventional Technologies Pte Ltd 36 Jalan Tukang Singapore **Local Agent:** Omega Health Care 581, Shewrapara Begum Rokeya Sarani, Mirpur, Dhaka - 1216 Bangladesh	**BioMatrix Flex** Drug Eluting Coronary Stent System (BioMatrix Flex DES) **Class: D**	Drug Eluting Coronary Stent System (BioMatrix Flex DES)	It is indicated for improving coronary luminal diameter for the treatment of de novo lessions in native coronary arteries with a reference diameter ranging between 2.25 mm and 4.00 mm. Stents with length 33 mm and 36 mm are only available for artery diameters ranging between 2.5 mm and 3.5 mm The BioMatrix Flex DES with stent length up to 28 mm is also indicated for use in patients with: * ST Elevated Myocardial Infrarction (STEMI) * Acute Coronary Syndromes (ACS) including ACS-STEMI, and unstable Angina. * Diabetes Melitus	Contraindication: None Side Effects: None	Switzerland Singapore CE Marking of Conformity	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj

112	**Manufacturer :** Boston Scientific Corporation, 300 Boston Scientific Way, USA **Local Agent :** Vastech Limited, Nurjehan Tower (6th Floor) 80/22 Mymensingh Road Dhaka-1000. Bangladesh.	Ultraflex Esophageal NG Stent System **Class: C**	Esophageal Stent (Metal)	The Ultraflex Esophageal NG Stent System is intended for maintaining esophageal luminal patency in esophageal strictures caused by intrinsic and/or extrinsic malignant tumors only. The Ultraflex Esophageal NG Covered Stent System is also indicated for occlusion of concurrent esophageal fistula.	**Contraindication** • Placement for occlusion of esophageal fistula of any type, unless a covered stent is being used. • Placement in esophageal strictures caused by benign tumors, as the long-term effects of the stent in the esophagus are unknown at this time. • Placement in strictures that cannot be dilated enough to pass the endoscope or the delivery system. • Placement of the stent's proximal end within 2 cm of the cricopharyngeal muscle. • Placement in an esophago-jejunostomy (following gastrectomy), as peristalsis may displace stent. • Placement in necrotic chronically bleeding tumors, if bleeding is active at the time of placement. • Placement in polypoid lesions. • Those patients for whom endoscopic techniques are contraindicated. Any use other than those specifically outlined under indications for use. **Adverse Effect:** The following complications have been reported in the literature for esophageal prosthesis. These include, but are not necessarily limited to: **Procedural Complications:** Bleeding, Perforation, Pain, Aspiration, Oxygen desaturation related to sedation, Infection **Post-stent Placement Complications** Bleeding Perforation, Pain, Stent migration, Tumor in-growth through stent, Tumor overgrowth around ends of stent, Foreign body sensation, Food bolus impaction (lavage and debridement may be necessary on a periodic basis), Reflux, Esophagitis, Edema, Ulceration, Infection and Septic shock, Fever, Septicemia, Recurrent dysphagia, Fistula with trachea, bronchi, or pleural space (other than that due to normal disease progression), Death (other than that due to normal disease progression), Stent Fracture, Tracheal compression/Airway compression	Certificate to foreign Government-USA CE Marking of Conformity (DEKRA)	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj

113.	QualiMed Innovative Medizinprodukte GmbH, Germany **Importer:** Biva International	**MAGMA** Rapamycin-Eluting Coronary Stent System **Class: D**	Rapamycin-Eluting Coronary Stent System	It is indicated for improving coronary luminal diameter in patients with symptomatic ischemic disease due to discrete de novo and in stent restenotic lesions (length S38) in native coronary arteries with vessel diameter of 2.0mm to 4.0mm. It has been shown to significantly reduce binary restenosis target lesion revascularization and angiographic late loss. The stent is also indicated for treatment of abrupt or threatening closure in patients with failed interventional therapy. The treated lesion (>50%) length should be less than the nominal stent length (10 to 38mm) with reference vessel diameters from 2.0 to 4.0.	**Contraindications:** The Rapamycin-Eluting Coronary Stent System is contraindicated for patients with: Known sensitivity to Rapamycin, Known allergy to stainless steel, Known allergy to PLGA polymer, Severe reaction to contrast agents, Patients in whom anti-platelet and /or anticoagulant therapy is contraindicated, In-stent Restenosis, Myocardial infarction <72hours, Stenting of saphenous Vein Grafts, Unprotected left main coronary artery, Total occlusion of target vessel, Heavily calcified lesions, Lesions involving arterial segments with highly tortuous anatomy, Lesions involving a bifurcation, Left ventricular ejection fraction <30% Cardiogenic shock, Presence of definite or probable intraluminal thrombus, Any patients judged to have a lesion which may interfere with proper stent deployment. **Aadverse events**: Adverse events may be associated with the implantation of a coronary stent in coronary arteries, but are not limited to the following: Allergic reaction, Aneurysm, Arrhythmias Death, Dissection, Drug reactions to antiplatetet agents/anticoagulation agents/contrast medium, Emboil, distal (tissue, air or thrombis emboli), Embolization,stent, Failure to deliver the stent to intended site, Hemorrhage, Hypotension/Hypertension, Infection and pain at the insertion site, Myocardial ischemia and /or infarction, Occlusion, Restenosis of stented segment (> 50% obstruction), Stroke, Thrombosis (acute, subacute or late), Ventricular fibrillation, Vessel spasm, Stent migration, Stent Collapse.	FSC-Germany EC design Examination Certificate	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj

114.	C.R. Bard INC. 8195 Industrial BLVD, Covington, GA, USA-30014 (**Local Agent;** Lilac Pvt. Ltd 72, New Elephant Road,)	Bard Fluoro-4 Silicone Ureteral Coil Stent **Class: C**	Silicone Ureteral Coil Stent	It is indicated to relive obstruction in a verity of benign, malignant and post – traumatic condition in the ureter such as stone and stone fragments, carcinoma of abdominal organs, retroperitoneal fibrosis or ureteral trauma or in association with Extracorporeal Shock Wave Lithotropsy	**Contraindication**: There are no known contraindications to use. **Warning**: This is a single use only. Do not desterilize any portion of this device. Reuse or repacking may create risk of patient or user infection.	Certificate to Foreign Gov.- USFDA CE Certificate- bsi, UK	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
115.	COVIDIEN LLC, Athlone, Ireland **Local Agent:** Medi-Q-Medicals	**DAR,** Close suction System	Close suction System	The incorporation of a suction system into a mechanical ventilator that permits airway suctioning without disconnecting patients from the ventilator.	Contraindication: None Side-effect : None	FSC Ireland	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
116.	Romsons International, India **Importer:** Barisal Surgical, 34/1, Mitford Road, Dhaka-1100	Vaccu suck suction set (Sterile) **Class: B**	Yankaur Suction Set	Vaccu Suck Suction Set is sterile, single use device intended for per-operative suction & removal of secretions, blood and surgical debris.	Contraindication: None Side-effect : None	FSC-India EC Certificate- Norway	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
117.	Shandong Yaohua Medical Instrument Corporation, China **Local Agent:** VIP Trader's, 52 New Eskaton Road, TMC Building (3rd Floor), Dhaka	Urine Test Strip **Class: B**	Urine Test Strip	Basic diagnostic tool used to determine pathological changes in a patient's urine in standard urinalysis	Contraindication & Side Effect : None	FSC-China	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

118.	**Manufacturer:** Onbo Electronic (Shenzhen), Co. Ltd, China **Supplier:** **Microlife Corporation** 9F, 431, RuiGang Road, NehHu, Taippei 114, Taiwan, R.O.C **Local agent:** **Transcom Distribution Co.** Ltd 52, Motijheel C/A, Dhaka-1000	Digital Thermometer	Digital Thermometer	It is used to take an oral temperature. It is a small hand-held device with a "window" showing your temperature in numbers.	Contraindication: None Side effect: None	Certificate for Exportation of Medical Products-China	Abjygv`b Kiv †h‡Z cv‡i	Abjygv`b Kiv nj
119.	Romsons Scientific & Surgical Industries Pvt. Ltd, India **Importer:** Barisal Surgical, Dhaka	Trachea Tee Plus **Class: B**	T-Connector with suction port & Tube	Used for connecting with breathing / ventilator circuits. 'T' connector with oxygen nipple, suction port and 2 mtr. long oxygen tube with connectors	Contraindication: None Side-effect : None	FSC-India EC Certificate-Norway	Abjygv`b Kiv †h‡Z cv‡i	Abjygv`b Kiv nj
120.	Romsons Scientific & Surgical Industries Pvt. Ltd, India **Importer:** Barisal Surgical, Dhaka	Tracheostomy Tube Plain **Class: B**	Tracheostomy Tube catheter Plain, without cuffed with stylet, with radio opaque line	Tracheostomy Tube- Plain is intended for respiration & venting after Tracheostomy.	Contraindication: None Side-effect : None	FSC-India EC Certificate-Norway	Abjygv`b Kiv †h‡Z cv‡i	Abjygv`b Kiv nj
121.	Romsons Scientific & Surgical Industries Pvt. Ltd, India **Importer:** Barisal Surgical, Dhaka	Tracheostomy Tube Cuffed **Class: B**	Tracheostomy Tube Cuffed, with pilot balloon with stylet with radio opaque line	Tracheostomy Tube- Cuffed is intended for respiration & venting after Tracheostomy.	Contraindication: None Side-effect : None	FSC-India EC Certificate-Norway	Abjygv`b Kiv †h‡Z cv‡i	Abjygv`b Kiv nj

122.	COVIDIEN LLC, Athlone, Ireland **Local Agent:** Medi-Q-Medicals	**Bronco.** Cath Endo Bronchial Tube **Class:B**	Endo Bronchial Tube	It is used in tracheal intubation during thoracic surgery and other medical conditions, to achieve the selective one sided ventilation of either the right or the left lung.	Contraindication: None Side-effect : None	FSC Ireland	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |
123.	COVIDIEN LLC, Athlone, Ireland **Local Agent:** Medi-Q-Medicals	**Argyle,** FeedingTube/ Suction control Tube **Class:B**	FeedingTube/ Suction control Tube	It is use for who have a deformity of mouth or esophagus, who have difficulty swallowing or keeping food down and ho aren't getting enough nutrition or fluids by mouth.	Contraindication: None Side-effect : None	FSC Ireland	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |
124.	COVIDIEN LLC, Athlone, Ireland **Local Agent:** Medi-Q-Medicals	**Taper Guard**-Evac, Oral Tracheal Tube **Class:B**	Oral Tracheal Tube	It is use for the people who are mechanically ventilated intensive care patients at risk of ventilator-associated pneumonia (VAP).	Contraindication: None Side-effect : None	FSC Ireland	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |

125.	COVIDIEN LLC, Athlone, Ireland **Local Agent:** Medi-Q-Medicals	**Lo-Contour** Oral/Nasal Tracheal Tube **Class: B**	Tracheal Tube	It is indicated for airway management by oral / nasal intubation of the trachea for anesthesia or other short-term procedures. Ordinarily, the cuff pressure should not exceed 25 cm H2O. However, clinical situations may arise where a higher sealing pressure is clinically indicated. Although uncuffed tracheal tubes are ordinarily used in pediatric airway management, in some cases (e.g., pulmonary function testing) use of a small diameter cuffed tracheal tube may be indicated.	**Contraindications:** Lo-Contour Cuffed Tracheal Tubes in procedures which will involve the use of a LASER or an electrosurgical active electrode in the immediate area of the device is contraindicated. Contact of the beam or electrode with the tracheal tubes, especially in the presence of oxygen-enriched or nitrous oxide containing mixtures, could result in the rapid combustion of the tube with harmful thermal effects and with emission of corrosive and toxic combustion products including hydrochloric acid (HCl). It has been reported by Hirshman and Smith that mixtures of nitrous oxide and oxygen support combustion about the same as pure oxygen and that in addition to ignition by direct contact with the beam, the interior of the tube can also be ignited by contact with flaming tissue in close proximity to the tip of the tracheal tube. **Adverse events:** The following adverse reactions have been reported to be associated with the use of cuffed tracheal tubes during the intubation procedure, during the intubation period, or subsequent to extubation. The order of listing is alphabetical and does not indicate frequency or severity. Reported adverse reactions* include: abrasion of the arytenoid cartilage vocal process; cartilage necrosis; cicatrix formation; consequences of failure to ventilate including death; damage to the perichondrium; development of dense or diffuse fibrosis invading the entire glottic area; emphysema; endobronchial aspiration; endobronchial intubation (hypoxemia); endotracheobronchial aspiration; epistaxis; esophageal intubation (stomach distention); excoriated membranes of the pharynx; eye trauma; fibrin deposition; formation of subglottic web; fracture-luxation of cervical column (spinal injury); fragmentation of cartilage; glottic edema (supraglottic, subglottic retroarytenoidal); granuloma of the inner arytenoid area; infections (laryngitis, sinusitis, abscess, respiratory tract infection); inflammation; intermittent aphonia and recurrent sore throat;	FSC Ireland	*Ab‡gv`b Kiv †h‡Z cv‡i |*	*Ab‡gv`b Kiv nj |*

126.	COVIDIEN LLC, Athlone, Ireland **Local Agent:** Medi-Q-Medicals	**Shiley,** Tracheostomy Tube **Class:B**	Tracheostomy Tube	It is use for the people who are on ventilator support, who have conditions that affect coughing or block the airways and who have swallowing problems.	Pain& inflammation	FSC Ireland	Abyôgv`b Kiv †h‡Z cv‡i |	Abyôgv`b Kiv nj |
127.	Romsons International, India **Importer:** Barisal Surgical, 34/1, Mitford Road, Dhaka-1100	Oxy Set **Class: B**	Nasal oxygen Twin bore cannula	Used for oxygen administration.	Contraindication: None Side-effect : None	FSC-India EC Certificate-Norway	Abyôgv`b Kiv †h‡Z cv‡i |	Abyôgv`b Kiv nj |
128.	**Manufacturer :** Smith & Nephew Inc., 101 Hessle Road, Hull, HU3 2BN, UK **Local Agent :** Vastech Limited, Nurjehan Tower (6th Floor) 80/22 Mymensingh Road Dhaka-1000. Bangladesh	Pico **Class: B**	Negative Pressure Wound Therapy System	It is indicated for patients who would benefit from a suction device (negative pressure wound therapy) as it may promote wound healing via removal of low to moderate levels of exudate and infectious materials. Examples of appropriate wound types include: Chronic, Acute, Traumatic, Subacute and dehisced wounds, Partial thickness burns, Ulcers (such as diabetic or pressure), Flaps and grafts, Surgically closed incision sites, PICO Single Use Negative pressure Wound Therapy System is suitable for use both in a hospital and homecare setting.	**Contraindications** The use of PICO is contraindicated in the presence of: • Patients with malignancy in the wound bed or margins of the wound (except in palliative care to enhance quality of life). • Previously confirmed and untreated osteomyelitis. • Non-enteric and unexplored fistulas. • Necrotic tissue with eschar present. • Exposed arteries, veins, nerves or organs. • Anastomotic sites. • Emergency airway aspiration. • Pleural, mediastinal or chest tube drainage. • Surgical suction. **Side Effects:** None	FSC-UK EC Certificate (bsi)	Abyôgv`b Kiv †h‡Z cv‡i |	Abyôgv`b Kiv nj |
129	**Legal Manufacturer:** Becton Dickinson and company 1 Becton Drive, Franklin Lake NJ 07417-1880 USA	BD Luer Adapter **Class: B**	Adapter	This is a specialized category of Adapter for which is used to connect venous access devices such as needle, blood collection set, and infusion sets to blood collection tubes. It is	**Contraindications:** None **Side-effects:** none	FSC-USA	Abyôgv`b Kiv †h‡Z cv‡i |	Abyôgv`b Kiv nj |

	Manufacturing site: Becton Dickinson and company (BD) 1575 Airport Road Sumter , SC, 29153 USA **Local Agent:** M/s Becton Dickinson & Company, 80, Kakrail, Dhaka			also sued in collection with non-needle devices for collection of blood from catheters.				
130	**Manufacturer:** Sichuan Guangyuan Kangkang Medical Instrument Co. Ltd. Zhongzi Industrial Park, Chaotian District, Guangyuan City, Sichuan Province, China. **Local Agent:** National Healthcare Services, 441/1, Senpara Parbata, Mirpur, Dhaka.	Single-use High-pressure Angiographic Extension Line with/without T tubing **Class: D**	Single-use High-pressure Angiographic Extension Line with/without T tubing	It is used as an interface between CT syringes or other syringes and needle catheters for use in delivering contrast media into vascular system.	**Contraindications:** Do not use if package is unsealed or damaged. Do not use the Extensions Line after the expiration date. **Side Effects:** None	FSC China, EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
131.	**Manufacturer:** Sichuan Guangyuan Kangkang Medical Instrument Co. Ltd. Zhongzi Industrial Park, Chaotian District, Guangyuan City, Sichuan Province, China. **Local Agent:** National Healthcare Services, 441/1, Senpara Parbata, Mirpur, Dhaka.	Single-use High-pressure Angiographic Syringes Kit without needle **Class: B**	Single-use High-pressure Angiographic Syringes Kit without needle	It is used with automatic injector machine for angiography.	**Contraindications:** Do not use if package is unsealed or damaged. Do not use the Extensions Line after the expiration date. **Side Effects:** None	FSC China, EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
132.	**Manufacturer:** Poly Medicure Ltd. Regd Office: 1st Floor, 12 Sant Nagar, East of Kailash,	Poly drain under water seal drainage system **Class: B**	Poly drain under water seal drainage system	It is used to drain the fluid from body collect due to various reasons like larger pleural effusions, post-operative	**Contraindications:** None **Side Effects:** None.	FSC-India EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

	New Delhi-110 065 India. Industry: Plot No. 104-105 & 115-116, Sector-59, HSIIDC Industrial Area, Ballabhagra-121004, Faridbad (HARYANA) India. **Local Agent:** Aglow International, 114, 1st Floor, Aziz Co-operative Super Market, Shahbag, Dhaka.			drainage etc. from a patient.				
133.	**Manufacturer:** Meril Healthcare PVT. India. **Local Agent:** Meril Bangladesh PVT. Ltd. Union Heights-1 13th Floor, Level-14, 55/2, Bir Uttam Qazi Nuruzzaman Saraka, West Panthapath, Dhaka.	Armar Bone Plates **Class: C**	Bone Plates	It is used for the surgical treatment of fractures, deformities, and tumor diseases of long bones, such as those of the arms and legs.	**Contraindications**: It is contraindicated in: Active Infection; Conditions which tend to retard healing such as blood supply limitations previous infections; Insufficient quantity or quality of bone to permit stabilization of the osteotomy; Lack of musculo-cutaneous cover; Muscular deficit, neurological deficiency or behavioral disorders which could submit the osteosynthesis to abnormal mechanical strains; Cases with malignant primary or metastatic; tumors which preclude adequate bone support or screw fixations; Conditions that restrict the patient's ability or willingness to follow postoperative instructions during the healing process; Foreign body sensitivity. **Adverse Events:** The following are specific adverse effects, which should be understood by the surgeon and explained to the patient. These do not include all adverse effects, which can occur with surgery in general, but are important considerations specific to	FSC India EC Certificate (Norway)	*Abygv`b Kiv †h‡Z cv‡i |*	*Abygv`b Kiv nj |*

					metallic internal stabilization devices. General surgical risks should be explained to the patient prior to surgery: - Infection or adverse reactions for a foreign body; - Pain, discomfort, or abnormal sensations due to the presence of the implant; - Loosening, bending, cracking, or fracture of the components or loss of fixation of bone attributable to nonunion, osteoporosis, markedly unstable comminuted fractures; loss of anatomic position with nonunion or malunion with rotation or angulation; - Migration of the implant, loosening of the implant; - Delayed correction in alignment; - Decrease in bone density due to stress shielding; - Bursitis.			
134.	**Manufacturer:** Ningbo Boya Medical Equipment Co. Ltd, No. 102, Jingsan Road, Yaobei Industrial Part, YuYao City, Zhejiang Province, China. **Local Agent:** Pride Medical Appliances Ltd. 19, Indira Road, Farmgate, Dhaka.	TornadoCare Ventilation Circuit/Catheter Mount **Class: B**	Anesthesia Breathing Circuit	It is used for bacteria, particle filtration in breathing machine and anesthesia machine and to increase the gas moisture degree	**Contraindication:** Product should not use after validity period. Do not use if the package is damaged. Not use for different user. **Side Effects:** None	FSC China	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj
135.	**Manufacturer:** Changzhou Chuangjia Medical Appliance Co. Ltd, Sanhekou Development Zone, Zhenglu, Town, Zhangzhou 213115, China.	TornadoCare Sterile Nasla Oxygen Cannula **Class: B**	Sterile Nasla Oxygen Cannula	This product is used mainly for clinical gynecological checkup.	**Contraindication:** Do not use if the package is damaged. For single use only. Do not use if the package is opened for long time. **Side Effects:** None	FSC China	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj

	Local Agent: Pride Medical Appliances Ltd. 19, Indira Road, Farmgate, Dhaka.							
136.	Manufacturer: Poly Medicure Ltd. Regd Office: 1st Floor, 12 Sant Nagar, East of Kailash, New Delhi-110 065 India. Industry: Plot No. 104-105 & 115-116, Sector-59, HSIIDC Industrial Area, Ballabhagra-121004, Faridbad (HARYANA) India. **Local Agent:** Aglow International, 114, 1st Floor, Aziz Co-operative Super Market, Shahbag, Dhaka.	Polymed Nasocath **Class: B**	Sterile Nasal Oxygen Cannula	It is used to provide oxygen for respiration with maximum comfort to the patient. System of attachment provides maximum freedom to the patient and leaves the patients mouth free for nutrition & communication.	**Contraindications:** Not to be used in patients with known hypersensitivity to any of the materials used. **Side Effects:** None.	FSC-India & EC Certificate	Abjygv`b Kiv †h‡Z cv‡i	Abjygv`b Kiv nj
137.	Manufacturer: Becton Dickinson Medical (S) Pte Ltd. 30 Tuas Avenue 2 Singapore. **Local Agent:** BD-Becton Dickinson. Flat 68, 6th Floor, Rupayan Karim Tower, 80 Kakrail V.I.P Road. Dhaka.	BD Arterial Cannula **Class: C**	Arterial Cannula	It is used for Arterial pressure monitoring.	Minor acute irritating	FSC Singapore	Abjygv`b Kiv †h‡Z cv‡i	Abjygv`b Kiv nj
138.	**Manufacturer:** Jafron Biomedical Co. Ltd., No.98 Technology Six Road, High-tech Zone, Zhuhai, Guangdong, China. **Local Agent:** Bangladesh Health Products, 2/4 Nawab Habibullah Road, Shahbag, Dhaka.	Jafron Disposable Hemoperfusion Cartridge **Class: C**	Disposable Hemoperfusion Cartridge	It is used for Renal failure, chronic nephropathy, uremia caused by all kinds of reasons; Related complications of maintenance hemodialysis: Renal osteopathy, refractory itch of skin, peripheral neuropathy, cardiovascular disease, refractory hypertension, nephro-encephalopathy, malnutrition	**Contraindication:** None **Side Effect**: None	FSC China, EC Certificate	Abjygv`b Kiv †h‡Z cv‡i	Abjygv`b Kiv nj

139.	**Manufacturer:** Goodman Co. Ltd., Goodman Research Centre, 276-1 idogane-cho, Seto-shi, Aichi-ken, Japan **Local Agent:** Advanced Meditech Suite: 115 (1st Floor), Krishnachura Commercial Complex, 24/B.C Shahid Minar Road, Kallanpur, Dhaka.	**Powered** Lacrossed 2 Coronary Balloon Dilatation Catheter **Class: C**	Coronary Balloon Dilatation Catheter	It is used in percutaneous transluminal coronary angioplasty (PTCA) for the purpose of dilating stenotic lesions within the coronary artery.	**Contraindications:** (1) The product is sterilized, is not re-usable and must not be re- sterilized. Resterilization and/or reuse could result in infection or degradation of product characteristics such as balloon size, shaft strength or lubricity and could result in failure of the product during use. (2) The product should only be used in facilities capable of performing emergency coronary artery bypass grafting (CABG) as a provision against complications that may cause injury or serious complications that could prove to be life threatening. (3) The product is a medical device and should only be used by physicians trained in the procedures of coronary angiography (CAG) and percutaneous transluminal coronary angioplasty (PTCA). (4) Do not disassemble or modify the device to avoid any unexpected incidents such as breakage of the device. **Complications/Side effect:** Possible adverse events include, but are not limited to, the following: Death, myocardial infarction, restenosis following angioplasty, internal hemorrhage, hematoma, ventricular fibrillation including arrhythmia, hypertension, hypotension, hemorrhagic	FSC-Japan	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

					complications, arterial spasm, stroke, distal embolization, arterial or bypass graft occlusion, arterial dissection or perforation or damage, unstable angina, medicinal reaction or allergenic reaction to contrast media, infection, arteriovenous fistula, air embolization, arterial dissection, blood loss from puncture site, ischemia caused by long duration inflation, intravascular thrombosis, nausea or vomiting, palpitation, tachycardia, bradycardia			
140.	**Manufacturer:** Poly Medicure Ltd. Regd Office: 1st Floor, 12 Sant Nagar, East of Kailash, New Delhi-110 065 India. Industry: Plot No. 104-105 & 115-116, Sector-59, HSIIDC Industrial Area, Ballabhagra-121004, Faridbad (HARYANA) India. **Local Agent:** Aglow International, 114, 1st Floor, Aziz Co-operative Super Market, Shahbag, Dhaka.	Catheter Mount **Class: B**	Catheter Mount	It is used for connection between endotracheal tube and breathing circuits. It is necessary to avoid drug & to prevent any chance of accidental situation.	**Contraindications:** Contraindicated in case of air leakage. **Side Effects:** None	FSC-India & EC Certificate	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
141.	**Manufacturer:** Poly Medicure Ltd. Regd Office: 1st Floor, 12 Sant Nagar, East of Kailash, New Delhi-110 065 India. Industry: Plot No. 104-105 & 115-116, Sector-59, HSIIDC Industrial Area, Ballabhagra-121004, Faridbad (HARYANA) India. **Local Agent:**	Polymed Ultra Cath Novo **Class: B**	Sterile, Thoracic Catheter with & without Trocar	It is used for post-operative drainage after cardio thoracic & thoracic surgery.	**Contraindications:** Uncontrolled bleeding diathesis, need for immediate thoractomy. **Side Effects:** None	FSC-India & EC Certificate	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

	Aglow International, 114, 1st Floor, Aziz Co-operative Super Market, Shahbag, Dhaka.							
142.	**Manufacturer:** Meril Life Science PVT. Ltd. India. **Local Agent:** Meril Bangladesh PVT. Ltd. Union Heights-1 13th Floor, Level-14, 55/2, Bir Uttam Qazi Nuruzzaman Saraka, West Panthapath, Dhaka.	**ASPIRON** Aspiration Catheter **Class: D**	Thrombus aspiration catheter	It is used for removal of fresh, soft emboli and thrombi from vessels in the coronary and peripheral vasculature system.	**Contraindications:** 1. It should not be used in vessels <1.5mm in diameter. 2. The catheter is contraindicated in the removal of fibrous, adherent, or calcified material (e.g., atherosclerotic plaque, chronic clot). 3. These components are contraindicated in the following lesion, a. Cerebral vasculature. b. Venous system. **Adverse Events:** Serious adverse events may arise when performing interventional procedures using this device. The operation should be performed in an institution where emergency procedures can be performed. During the use of this device, patient vital signs should be monitored. In case of any abnormalities, the appropriate measures should be taken based on the physician's judgment. As with all percutaneous interventions, adverse events may include: - Access site complications (i.e. Arteriovenous fistula, dissection, hematoma, hemorrhage, pseudoaneurysm, bleeding). Death, - Myocardial infarction, Coronary or bypass graft thrombosis or occlusion, Myocardial ischemia, Stroke/CVA, - Emergent or non-emergent bypass	FSC-India & EC Certificate (Norway)	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj

					graft surgery, - Emboli (air, tissue or			
143.	**Manufacturer:** Cook Incorporated USA **Local Agent:** Nature and Nurture Ltd. House-19, Road-1, Block-A, Banasree, Dhaka	SYDNEY IVF Embryo Transfer Set **Class: B**	Embryo Transfer Catheter	It is used for the atraumatic transfer of embryos to the uterine cavity.	**Contraindications:** None. **Side effects:** None	FSC- USA	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
144.	**Manufacturer:** G. Surgiwear Limited Rasoolpur Jahanganj, Near Hathoda Chauraha, Shahjahanpur-242001, U.P., India **Local Agent:** Inogen Systems Cha-73/2, Progati Sarani, North Badda, Dhaka-1212	SURGIWEAR **Y connector with extra-long Ventricular Catheter** **Class: D**	Sterile Catheter Connector	It is used to connect between two lines during drainage system.	**Contraindications:** None **Side-effects:** none	FSC-India	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
145.	**Manufacturer:** G. Surgiwear Limited Rasoolpur Jahanganj, Near Hathoda Chauraha, Shahjahanpur-242001, U.P., INDIA **Local Agent:** Inogen Systems Cha-73/2, Progati Sarani, North Badda, Dhaka-1212	SURGIWEAR **G-Bone Modified Hydroxyapatite** (Granules**:** 10cc Pack / 1cc Pack; Blocks; Dowel) **Class: D**	Bone Cement	G-Bone is indicated in almost all the places where a bone graft is required. Following are some of the indicative uses specialty wise. • **Neurosurgery:** Cranioplasty, cavity filling and other kind of non-weight bearing reconstructions. • **Orthopedic Surgery:** Cavity filling, gap filling, osteotomy reconstructions and other places of reconstructions, non union fracture and filling the gaps in cases of bone loss due to trauma. • **Plastic Surgery:** Maxillofacial reconstructions, bony augmentation, hand surgery & cavity filling. • **Dental Surgery:** Maxillofacial reconstruction surgery,	**Contraindications:** Presence of infection whether local or general, prohibits all kind of implantation procedures. Open wounds will not respond well with bone grafting surgeries. Poor general health and metabolic deficiencies will not produce desired results. Above mentioned contraindications are general. The surgeon must evaluate the individual patient for the risks involved. **Side Effects:** Complications which may result from the use of this product include the risks associated with the medication and methods utilized in the surgical procedure, as well as patients response, reaction or degree of intolerance to any foreign object implanted into the body.	FSC-India	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

				periodontics, orthodontics and other areas requiring a bone graft. • **Other Specialties:** Where a bone graft is required.				
146.	**Manufacturer:** G. Surgiwear Limited Rasoolpur Jahanganj, Near Hathoda Chauraha, Shahjahanpur-242001, U.P., INDIA **Local Agent:** Inogen Systems Cha-73/2, Progati Sarani, North Badda, Dhaka-1212	SURGIWEAR **G-Bone Calcium Phosphate Cement** **Class: D**	Bone Cement	G-Bone Cement is indicated in almost all the places where a bone graft is required. Following are some of the indicative uses specialty wise. • Neurosurgery: Filling of bur holes, lumber & cervical Fusion, vertebral reconstruction, Cranioplasty, cavity filling and other kind of reconstructions. • Orthopedic Surgery: Cavity filling, gap filling, Lumber & cervical Fusion, vertebral reconstruction, osteotomy reconstructions and other places of reconstructions, non union fracture and filling the gaps in cases of bone loss due to trauma. • Plastic Surgery: Maxillofacial reconstructions, bony augmentation, hand surgery & cavity filling. • Dental Surgery: Maxillofacial reconstruction surgery, periodontics, orthodontics and other areas requiring a bone graft. • Other Specialties: Where a bone graft is required.	**Contraindications:** Presence of infection whether local or general, prohibits all kind of implantation procedures. Open wounds will not respond well with bone grafting surgeries. Poor general health and metabolic deficiencies will not produce desired results. Above mentioned contraindications are general. The surgeon must evaluate the individual patient for the risks involved. **Side Effects:** Complications which may result from the use of this product include the risks associated with the medication and methods utilized in the surgical procedure, as well as patients' response, reaction or degree of intolerance to any foreign object implanted into the body. The medical literature is full of hazards and complications associated with use of Calcium Hydroxyapatite. Some of the major hazards are infection, extrusion of grafted Material, serous discharge, serous collection, hemorrhage, skin erosion and migration.	FSC-India	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
147.	**Manufacturer:** Sree Umiya Surgical Pvt. Ltd. Plot o. 4704, Phase. IV, G.I.D.C, VATA, Ahmedabad. India.	Mediplus Medi Vac set **Class: C**	Medi Vac set made with Blow Container, Butch, Round Butch, Hanger "Y" Connector, Adopter, Closure, S.S.	It is used as an effective device for closed wound drainage.	**Contraindication:** None **Side Effect**: None	FSC India, EC Certificate	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

	Local Agent: Hard Won Power Industries Ltd. House No. 39, Road-12, Shekhertek, Adabor. Dhaka.		Rod, Clamp & N.T. PVC Tubing					
148.	**Manufacturer:** B. Braun Melsungen AG Carl-Barun-StraBe1, Melsungen Germany. **Local Agent:** Asia Pacific Medicals Ltd. 775, Satmasjid Road (2nd Floor), Dhanmodi, Dhaka.	Proxima 2 Plate	Colostomy Base Plate	collection of waste from a surgically diverted biological system (colon, ileum, bladder) and the creation of a stoma	**Contraindication:** None **Side Effect**: None	FSC Germany	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |
149.	**Manufacturer:** B. Braun Melsungen AG Carl-Barun-StraBe1, Melsungen Germany. **Local Agent:** Asia Pacific Medicals Ltd. 775, Satmasjid Road (2nd Floor), Dhanmodi, Dhaka.	Super Filler	Skin barrier Colostomy Paste	pouch leakage, meltdown of skin barrier, parastomal ulceration, irritant dermatitis, allergic dermititis, latex sensitivity and fitting ostomy appliances to paediatrics.	**Contraindication:** None **Side Effect**: None	FSC Germany	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |
150.	**Manufacturer:** Goodman Co. Ltd., Goodman Research Centre, 276-1 idogane-cho, Seto-shi, Aichi-ken, Japan **Local Agent:** Advanced Meditech Suite: 115 (1st Floor), Krishnachura Commercial Complex, 24/B.C Shahid Minar Road, Kallanpur, Dhaka.	GOODTEC Y-Connector Set **Class: B**	Connector Set	It is designed to work in conjunction with guiding catheters, etc to reduce blood loss while assisting in manipulation of catheters, etc., injection of contrast solution from a side port, injection of medicine or saline solution, and measurement of blood pressure.	**Contraindication:** - This product is sterilized, is not reusable and must not be re-sterilized. Re-sterilization could lead to infection of patient or degradation of the material. - Not to be used with products comprising of organic solvent, fat emulsion, or oil-based components. It may incur damage to product. - This product is a medical device and should only be used by physicians who are trained in the procedures of coronary angiography (CAG) and percutaneous transluminal coronary angioplasty (PTCA). **Side Effects:** • Hypotension/hypertension	FSC-Japan	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |

					• Hemorrhagic complications • Infection • Air embolism			
151.	**Manufacturer:** Poly Medicure Ltd. Regd Office: 1st Floor, 12 Sant Nagar, East of Kailash, New Delhi-110 065 India. Industry: Plot No. 104-105 & 115-116, Sector-59, HSIIDC Industrial Area, Ballabhagra-121004, Faridbad (HARYANA) India. **Local Agent:** Aglow International, 114, 1st Floor, Aziz Co-operative Super Market, Shahbag, Dhaka.	Polyseal under water seal drainage system (adult/kid) **Class: B**	Polyseal under water seal drainage system (adult/kid)	It is used to drain the fluid from body collected due to various reasons like larger pleural effusions, post-operative drainage etc. from a patient.	**Contraindications:** Not to be used in case of uncontrolled suction. **Side Effects:** None	FSC India, EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |
152.	**Manufacturer:** G. Surgiwear Limited Rasoolpur Jahanganj, Near Hathoda Chauraha, Shahjahanpur-242001, U.P., India **Local Agent:** Inogen Systems Cha-73/2, Progati Sarani, North Badda, Dhaka-1212	SURGIWEAR **Ceflui Ventricular External Drainage System** **Class: D**	External CSF Drainage System	It is to be used for external drainage of CSF, wherever indicated. Some of the principal indications are: Posterior fosa tumor, hematomas posterior fosa, IV ventricle obstruction, cranial injuries, subdural hematomas, meningitis, ventriculitis and subarachnoid hemorrhage into CSF and cerebral abscess. Reyes syndrome or similar encephalopathy and other cases where monitoring of CSF is required.	**Contraindications:** External CSF drainage and monitoring with a ventricular or lumbar catheter is contraindicated in patients receiving lumbar anticoagulants or who are known to have a bleeding diathesis. The ventricular catheter is contraindicated in scalp infection. Lumbar catheter insertion is contraindicated in non-communicating hydrocephalus, infection in surrounding area and patients with blockage CSF flow due to trauma, hematomas, fracture or tumor. Monitoring and or external drainage in such conditions is at the discretion of the physician. **Side Effects:** The principal complications associated with	FSC-India	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |

					'External Drainage System' implantations are obstruction, functional failure of system and infection.			
153.	**Manufacturer:** G. Surgiwear Limited Rasoolpur Jahanganj, Near Hathoda Chauraha, Shahjahanpur-242001, U.P., India **Local Agent:** Inogen Systems Cha-73/2, Progati Sarani, North Badda, Dhaka-1212	SURGIWEAR **Ventricular External Drainage System** **Class: D**	External CSF Drainage System	It is to be used for external drainage of CSF, wherever indicated. Some of the principal indications are: Posterior fosa tumor, hematomas posterior fosa, IV ventricle obstruction, cranial injuries, subdural hematomas, meningitis, ventriculitis and subarachnoid hemorrhage into CSF and cerebral abscess. Reyes syndrome or similar encephalopathy and other cases where monitoring of CSF is required.	**Contraindications:** External CSF drainage and monitoring with a ventricular or lumbar catheter is contraindicated in patients receiving lumbar anticoagulants or who are known to have a bleeding diathesis. The ventricular catheter is contraindicated in scalp infection. Lumbar catheter insertion is contraindicated in non-communicating hydrocephalus, infection in surrounding area and patients with blockage CSF flow due to trauma, hematomas, fracture or tumor. Monitoring and or external drainage in such conditions is at the discretion of the physician. **Side Effects:** The principal complications associated with 'External Drainage System' implantations are obstruction, functional failure of system and infection.	FSC-India	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
154.	**Manufacturer:** G. Surgiwear Limited Rasoolpur Jahanganj, Near Hathoda Chauraha, Shahjahanpur-242001, U.P., India **Local Agent:** Inogen Systems Cha-73/2, Progati Sarani, North Badda, Dhaka-1212	SURGIWEAR **Lumbar External Drainage System** **Class: D**	External CSF Drainage System	It is to be used for external drainage of CSF, wherever indicated. Some of the principal indications are: Posterior fosa tumor, hematomas posterior fosa, IV ventricle obstruction, cranial injuries, subdural hematomas, meningitis, ventriculitis and subarachnoid hemorrhage into CSF and cerebral abscess. Reyes syndrome or similar encephalopathy and other cases where monitoring of CSF	**Contraindications:** External CSF drainage and monitoring with a ventricular or lumbar catheter is contraindicated in patients receiving lumbar anticoagulants or who are known to have a bleeding diathesis. The ventricular catheter is contraindicated in scalp infection. Lumbar catheter insertion is contraindicated in non-communicating hydrocephalus, infection in surrounding area and patients with blockage CSF flow due to trauma, hematomas, fracture or tumor. Monitoring and or external drainage in	FSC-India	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

				is required.	such conditions is at the discretion of the physician. **Side Effects:** The principal complications associated with 'External Drainage System' implantations are obstruction, functional failure of system and infection.			
155.	Manufacturer: Boston Scientific Corporation, 300 Boston Scientific Way, Marlborough, MA, USA. **Local Agent:** Medi Graphic Trading Ltd, 14, Purana P altan. Dar Us Salam Arcade, 3rd Floor, Dhaka.	Cutting Loop Electrode **Class: B**	Urological Electrosurgical devices	It is used in commercially available RF Generators for the endoscopic surgery for general urological soft tissue.	No contraindications are known for the RF electrode products.	FSC-USA, EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |
156.	**Manufacturer:** Boston Scientific Corporation, 300 Boston Scientific Way, Marlborough, MA, USA. **Local Agent:** Medi Graphic Trading Ltd, 14, Purana P altan. Dar Us Salam Arcade, 3rd Floor, Dhaka.	Contour Embolization Particles **Class: C**	Embolization Particles	It is intended for the embolization of peripheral hypervascular tumors.	Contraindication: Contraindicated in Vascular anatomy or blood flow precledes stable, selective contour embolization particles or catheter placement, presence of vasospasm and hemorrhage. Side effect: None.	FSC USA, EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |
157.	**Manufacturer:** Ningbo Boya Medical Equipment Co. Ltd, No. 102, Jingsan Road, Yaobei Industrial Part, YuYao City, Zhejiang Province, China. **Local Agent:** Pride Medical Appliances Ltd. 19, Indira Road, Farmgate, Dhaka.	TornadoCare HMEF/Bacterial Filter **Class: B**	Breathing System Filter	It is used for bacteria, particle filtration in breathing machine and anesthesia machine and to increase the gas moisture degree.	**Contraindication:** Product should not use after validity period. Do not use if the package is damaged. Not use for different user. **Side Effects:** None	FSC-China	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |
155.	**Manufacturer:** Ningbo Beilun Hengda Medical Dressing Factory. No.78, North Yanshanhe	Soluble Hemostatic Gauze **Class: B**	Soluble hemostatic Gauze	It is used for Adhering and creating pressure to seal the wound.	**Contraindications:** Soluble Hemostatic Gauze should not be used for internal surgical hemostasis (intracranial operation, operation in	FSC China	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |

	Road, Daqi Street, Beilun, Ningbo, Zhejiang Province, China. **Local Agent:** Swadesh. 30, Bijoy Nagar Road (3rd Floor), Dhaka.				chest and abdominal cavities, and pelvic operation), patients with severe disturbances of blood coagulation, large area of wound hemostasis and hemorrhage of large arteries. **Adverse reactions:** No reports on adverse reaction have been issued.			
159.	**Manufacturer:** Nanjing Hong An Medical Appliances Co. Ltd., No.26 Hengguang Road, Nanjing Economic and Technological, District: Nanjing, China. **Local Agent:** Pride Medical Appliances Ltd., 19, Indira Road, Farmgate, Dhaka.	Tornadocare Guedel Airway **Class: B**	Disposable Guedel/ Oropharyngeal Airway	It is used in management of endotracheal clinical issues.	**Contraindication:** Product should not use after validity period. Do not use if the package is damaged. Not use for different user. **Side Effects:** None	FSC China	Abyôgv`b Kiv †h‡Z cv‡i	Abyôgv`b Kiv nj
160.	Manufacturer: Boston Scientific Corporation, 300 Boston Scientific Way, Marlborough, MA, USA. **Local Agent:** Medi Graphic Trading Ltd, 14, Purana P altan. Dar Us Salam Arcade, 3rd Floor, Dhaka.	V-14 Control Wire Guidewire **Class: C**	Peripheral Guide wire	It is indicated to facilitate the place balloon dilatation catheter.	Contraindication: None. Side effect: Allergic reaction, Embolism.	FSC USA, EC Certificate	Abyôgv`b Kiv †h‡Z cv‡i	Abyôgv`b Kiv nj
161.	**Manufacturer:** Meril Healthcare PVT. India. **Local Agent:** Meril Bangladesh PVT. Ltd. Union Heights-1 13th Floor, Level-14, 55/2, Bir Uttam Qazi Nuruzzaman Saraka, West Panthapath, Dhaka.	Latitud Hip Replacement System **Class: C**	Total Hip Replacement System	It is indicated for the following conditions Non-inflammatory degenerative joint diseases including osteoarthritis, post traumatic arthritis and avascular necrosis Rheumatoid arthritis Congenital hip dysplasia Acute traumatic fracture of the femoral head or neck Dislocation of hip Certain cases of Ankylosis Correction of	**Contraindications:** -Active local or systemic infection. - Poor bone quality or Bone stock that is inadequate for support or fixation of the prosthesis. - Loss of musculature, neuromuscular compromise or vascular deficiency in the affected limb rendering the procedure unjustified. - Any condition that may interfere with the survival of the implants such as Charcot's disease, or Paget's disease. - Skeletally immature patients. - Metabolic disorder which may impair	FSC India EC Certificate (Norway)	Abyôgv`b Kiv †h‡Z cv‡i	Abyôgv`b Kiv nj

				functional deformity Revision of failed joint reconstruction or treatment Treatment of nonunion, femoral neck and trochanteric fractures of the proximal femur	bone formation. **Side Effects:** There may be adverse events after surgery, regardless of the type of hip system implanted including: Hip dislocation, when the ball of the thighbone (femur) slips out of its socket in the hip bone (pelvis), Bone fracture, Joint infection, Local nerve damage with numbness/weakness, Device loosening or breakage, Difference in leg lengths, Bone loss (osteolysis).			
162.	**Manufacturer:** St. Jude Medical, CARDIAC RHYTHM MANAGEMENT Division 15900 Valley View Ct Sylmar, CA USA 91342 **Local Agent:** The Spondon Ltd. BSEC Bhaban, Level-9, 102 Kazi Nazrul Islam Avenue, Kawran Bazar, Dhaka.	Implantable Cardioverter/ Defibrillators (Fortify Assura, Ellipse, Quadra Assura, Unify Assura) **Class: D**	Implantable Cardioverter/ Defibrillators	It is used to provide ventricular antitachycardia pacing and ventricular defibrillation for automated treatment of life threatening ventricular arrhythmias.	**Contraindications:** Contraindications for use of the pulse generator system include ventricular tachyarrhythmias resulting from transient or correctable factors. **Side Effects:** None	CFG- USA EC Certificate	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
163.	**Manufacturer:** Poly Medicure Ltd. Regd Office: 1st Floor, 12 Sant Nagar, East of Kailash, New Delhi-110 065 India. Industry: Plot No. 104-105 & 115-116, Sector-59, HSIIDC Industrial Area, Ballabhagra-121004, Faridbad (HARYANA) India. **Local Agent:** Aglow International, 114, 1st Floor, Aziz Co-operative Super Market, Shahbag, Dhaka	Polymed High Pressure Extension Line **Class: B**	Extension Line as an accessory of infusion set	It is used for infusion of fluid such as parenteral nutrition and administration of other drugs and medicines.	**Contraindications:** It is contraindicated in case of highly viscous fluid. Not to be used in patients with known hypersensitivity to any of the materials used. **Side Effects:** None	FSC-India	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
164.	**Manufacturer:** B. Braun Melsungen AG Carl-Barun-StraBe1, Melsungen Germany.	Intrafix Safe set (Auto Lock Infusion Set) **Class: B**	Intrafix Safeset (Auto Lock Infusion Set)	It is used as auto lock infusion set.	**Contraindication:** None **Side Effect**: None	FSC Germany	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

	Local Agent: Asia Pacific Medicals Ltd. 775, Satmasjid Road (2nd Floor), Dhanmodi, Dhaka							
165.	Manufacturer: Meril Endo Surgery PVT. Ltd. India. **Local Agent:** Meril Bangladesh Pvt. Ltd. Union Heights-1 13th Floor, Level-14, 55/2, Bir Uttam Qazi Nuruzzaman Saraka, West Panthapath, Dhaka	**MeriTe Cu 375** Intrauterine Contraceptive device **Class: C**	Copper Intrauterine Contraceptive device	It is used for reversible birth control. It is indicated for intrauterine contraception for up to 5 years.	**Contraindications:** - Pregnancy or suspicion of pregnancy - Abnormalities of the uterus resulting in distortion of the uterine cavity - Acute pelvic inflammatory disease or current behavior suggesting a high risk for pelvic inflammatory disease - Postpartum endometritis or postabortal endometritis in the past 3 months - Anaemia - Valvular heart disease - Coagulation disorders - Anti-inflammatory treatment - Mucopurulent cervicitis - Wilson's disease - Allergy to any component of the device. **Side effects:** none	FSC India & EC Certificate (Norway)	Abyôgv`b Kiv †h‡Z cv‡i	Abyôgv`b Kiv nj
166.	**Manufacturer:** Meril Endo Surgery PVT. Ltd. India. **Local Agent:** Meril Bangladesh PVT. Ltd. Union Heights-1 13th Floor, Level-14, 55/2, Bir Uttam Qazi Nuruzzaman Saraka, West Panthapath, Dhaka	Fiona Hormonal Intrauterine Device **Class: C**	Levonorgesterol 20µgm/24hours Hormonal Intrauterine Device	It is intended for contraception. It can also be used in idiopathic menorrhagia and protection from endometrial hyperplasia during estrogen replacement therapy.	**Contraindications:** • History of Pelvic inflammatory disease or purulent cervicitis. • Recent exposure to sexually transmitted infection. • Existing pregnancy • Between 48 hours or 4 weeks of postpartum. • Uterine abnormality • Gynecological cancer. **Side effects:** Uterine/vaginal bleeding alteration, amenorrhea, Intermenstrual bleeding and spotting, abdominal/pelvic pain, Overain cysts, acne, menorrhagia, headache/migraine, breast	FSC-India & EC Certificate (Norway)	Abyôgv`b Kiv †h‡Z cv‡i	Abyôgv`b Kiv nj

					tenderness/pain.			
167.	**Manufacturer:** Meril Endo Surgery PVT. Ltd. India. **Local Agent:** Meril Bangladesh PVT. Ltd. Union Heights-1 13th Floor, Level-14, 55/2, Bir Uttam Qazi Nuruzzaman Saraka, West Panthapath, Dhaka.	**Erinna** Hormonal Intrauterine Device **Class: C**	Levonorgesterol 20µgm/24hours Hormonal Intrauterine Device	It is intended for contraception. It can also be used in idiopathic menorrhagia and protection from endometrial hyperplasia during estrogen replacement therapy.	**Contraindications:** • History of Pelvic inflammatory disease or purulent cervicitis. • Recent exposure to sexually transmitted infection. • Existing pregnancy • Between 48 hours or 4 weeks of postpartum. • Uterine abnormality • Gynecological cancer. **Adverse reactions:** Uterine/vaginal bleeding alteration, amenorrhea, Intermenstrual bleeding and spotting, abdominal/pelvic pain, Overain cysts, acne, menorrhagia, headache/migraine, breast tenderness/pain.	FSC India EC Certificate (Norway)	Abjgv`b Kiv †h‡Z cv‡i |	Abjgv`b Kiv nj |
168.	**Legal Manufacturer:** Cardis Corporation 14201. N.W. 60th Ave. Miami Lakes. USA **Manufacturering Site:** CARDIS DE Mexico. SA DE CV, Mexico **Local Agent:** Medi Card Ltd. ¼, Paribagh, Mymensingh Road, Dhaka.	Aventi Introducer Sheath **Class: B**	Introducer Sheath	It is used for arterial and venous procedures requiring percutaneous introduction of intravascular devices,	**Contraindication**: None **Side Effects**: None	FSC- USA & EC Certificate	Abjgv`b Kiv †h‡Z cv‡i |	Abjgv`b Kiv nj |
169.	Manufacturer: Boston Scientific Corporation, 300 Boston Scientific Way, Marlborough, MA, USA. **Local Agent:** Medi Graphic Trading Ltd, 14, Purana P altan. Dar Us Salam Arcade, 3rd Floor, Dhaka.	Single Action Pumping System **Class: B**	Urological Irrigation device	It is used with urological and/or endoscopic surgical procedures requiring continuous irrigation.	Contraindication: None. Side effect: Hydronephrosis of the Urinary Tract.	FSC-USA EC Certificate	Abjgv`b Kiv †h‡Z cv‡i |	Abjgv`b Kiv nj |
170.	**Manufacturer:**	Mediplus I.V. Flow	IV Flow Regulator	It is used for Precise flow rate	**Contraindication:** None	FSC-India	Abjgv`b Kiv †h‡Z cv‡i |	Abjgv`b Kiv nj |

	Sree Umiya Surgical Pvt. Ltd. Plot o. 4704, Phase. IV, G.I.D.C, VATA, Ahmedabad. India. **Local Agent:** Hard Won Power Industries Ltd. House No. 39, Road-12, Shekhertek, Adabor. Dhaka.	Regulator **Class: C**	made with Flow Regulator, Tubing & Luer Lock	control of IV fluids.	**Side Effect**: None	EC Certificate		
171.	**Manufacturer:** Meril Healthcare PVT. India. **Local Agent:** Meril Bangladesh PVT. Ltd. Union Heights-1 13th Floor, Level-14, 55/2, Bir Uttam Qazi Nuruzzaman Saraka, West Panthapath, Dhaka.	Destiknee Total Knee system **Class: C**	This Knee system consists of the following components: - Femoral Knee Component PS & CR (Left & Right), - Tibia Base Plate, - Tibia Liner PS & CR / All Poly (Tibial base plate + Tibial liner), - Patellar component. - Instrument set for knee replacement	- Severe knee joint pain, loss of mobility, and disability due to: rheumatoid arthritis, osteoarthritis, traumatic arthritis, polyarthritis - Correction of functional deformities. - Post-traumatic loss of knee joint contour, particularly when there is patellofemoral erosion, dysfunction, or prior patellectomy. - Moderate valgus, varus, or flexion trauma. - Knee fractures untreatable by other methods. - Revision surgery where sufficient bone stock and soft tissue integrity are present.	**Contraindications**: - History of infection in the affected joint. - Compromised skeletal bone quality. - Neuropathic disease that adversely affects prosthetic joints. - Osteoporosis or deficiency of musculature. - Pain-free and stable arthrodesis in an adequate functional position. - Patients with vascular deficiency at the bone site. - Patients with inadequate bone stock to assure a firm press fit and close apposition of the cut bone surfaces to the prosthesis. - Patients with inadequate bone quality (e.g. severe osteoporosis). - Lack of stability of the implanted components throughout a full range of motion. **Adverse reactions:** Long term swelling or infection, no improvement in range of motion, neuropathic disorder, dislocations, bone fractures, aseptic loosening of implant.	FSC India & EC Certificate (Norway)	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

172.	**Manufacturer:** Maxx Orthopedic Inc. USA **Local Agent:** Meril Bangladesh PVT. Ltd. Union Heights-1 13th Floor, Level-14, 55/2, Bir Uttam Qazi Nuruzzaman Saraka, West Panthapath, Dhaka.	Freedom Total Knee system **Class: C**	It is a system of components intended to replace the femoral, tibial and patella articular surfaces of the knee joint. Components are available in many styles and sizes and are manufactured from various types of metals and non-metallic materials.	It is used in the followings: - Severe knee joint pain, loss of mobility, and disability due to: rheumatoid arthritis, osteoarthritis, traumatic arthritis, polyarthritis. - Correction of functional deformities. - Post-traumatic loss of knee joint contour, particularly when there is patellofemoral erosion, dysfunction, or prior patellectomy. - Moderate valgus, varus, or flexion trauma. - Knee fractures untreatable by other methods. - Revision surgery where sufficient bone stock and soft tissue integrity are present. - The Freedom Total Knee System, Freedom Stemmed Tibial Components and Freedom PCK Components are indicated for cemented fixation. Only Cementless Femoral (CR and PS) components with porous coating are additionally indicated for cementless biological fixation application.	**Contraindications:** - History of infection in the affected joint. - Compromised skeletal bone quality. - Neuropathic disease that adversely affects prosthetic joints. - Osteoporosis or deficiency of musculature. - Pain-free and stable arthrodesis in an adequate functional position. - Patients with vascular deficiency at the bone site. - Patients with inadequate bone stock to assure a firm press fit and close apposition of the cut bone surfaces to the prosthesis. - Patients with inadequate bone quality (e.g. severe osteoporosis). Lack of stability of the implanted components throughout a full range of motion. **Adverse Effects:** - Long term swelling or infection. - No improvement in range of motion. - Neuropathic disorders. - Dislocations, bone fractures, and/or joint instability. - Per literature, there is a chance that wear of polyethylene components may result in bone resorption, loosening, and related infection. - Possibility for metal sensitivity reactions. - Venous thrombosis.	FSC India & EC Certificate (Germany)	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj

					- Prolonged and excessive joint pain and/or inflammation. - Aseptic loosening of implant. - Possible detachment of the coating(s) on components with porous coating, potentially leading to increased debris particles -			
173.	Manufacturer: Huaian Wanjia Medical Device Co. Ltd. China. **Local Agent:** Shanto Enterprise, 51, Islampur Road, Dhaka.	Disposable Blood Lancet **Class: B**	Disposable Blood Lancet	Capillary blood sampling, make punctures, such as a fingerstick, to obtain small blood specimens.	**Contraindication:** None **Side Effect**: None	FSC-China	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
174.	**Manufacturer:** Abbott Medical Optics Inc. 1700 East St. Andrew Place Santa Ana, USA. **Local Agent:** S.P Trading House, 24-25 Dilhu8sha, C/A Dhaka.	Tecnis 1 Piece Multifocal IOL (ZMBOO) **Class: C**	Intraocular Lenses	It is used for the visual correction of aphakia in adult patients in whom a cataractous lens has been removed by extra capsular cataract extraction.	May be, Correct the vision problems of cataracts or presbyopia, minor infection,Corneal Adema, Increased Intraocular Pressure, Leakage, IOL Decentration, IOL Power Miscalculation, Retinal Detachment	FSC-USA EC Certificate	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
175.	**Manufacturer:** Abbott Medical Optics Inc. 1700 East St. Andrew Place Santa Ana, USA. **Local Agent:** S.P Trading House, 24-25 Dilhu8sha, C/A Dhaka.	Tecnis 1Piece IOL (ZCBOO) **Class: C**	Intraocular Lenses	It is used for the visual correction of aphakia in adult patients in whom a cataractous lens has been removed by extra capsular cataract extraction.	May be, Correct the vision problems of cataracts or presbyopia, minor infection,Corneal Adema, Increased Intraocular Pressure, Leakage, IOL Decentration, IOL Power Miscalculation, Retinal Detachment	FSC-USA, EC Certificate	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
176.	**Manufacturer:** Abbott Medical Optics Inc. 1700 East St. Andrew Place Santa Ana, USA. **Local Agent:** S.P Trading House, 24-25 Dilhu8sha, C/A Dhaka.	Epoch IOL (EP525A, EP6125A) **Class: C**	Intraocular Lenses	It is used for the visual correction of aphakia in adult patients in whom a cataractous lens has been removed by extra capsular cataract extraction.	May be, Correct the vision problems of cataracts or presbyopia, minor infection,Corneal Adema, Increased Intraocular Pressure, Leakage, IOL Decentration, IOL Power Miscalculation, Retinal Detachment	FSC-USA, EC Certificate	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

177.	**Manufacturer:** Abbott Medical Optics Inc. 1700 East St. Andrew Place Santa Ana, USA. **Local Agent:** S.P Trading House, 24-25 Dilhu8sha, C/A Dhaka.	Sensar 1Piece IOL (AABOO) **Class: C**	Intraocular Lenses	It is used for the visual correction of aphakia in adult patients in whom a cataractous lens has been removed by extra capsular cataract extraction.	May be, Correct the vision problems of cataracts or presbyopia, minor infection,Corneal Adema, Increased Intraocular Pressure, Leakage, IOL Decentration, IOL Power Miscalculation, Retinal Detachment	FSC-USA, EC Certificate	Abyôgv`b Kiv †h‡Z cv‡i	Abyôgv`b Kiv nj
178.	**Manufacturer:** Abbott Medical Optics Inc. 1700 East St. Andrew Place Santa Ana, USA. **Local Agent:** S.P Trading House, 24-25 Dilhu8sha, C/A Dhaka.	Sensar Soft Acrylic IOL (AR40E) **Class: C**	Intraocular Lenses	It is used for the visual correction of aphakia in adult patients in whom a cataractous lens has been removed by extra capsular cataract extraction.	May be, Correct the vision problems of cataracts or presbyopia, minor infection,Corneal Adema, Increased Intraocular Pressure, Leakage, IOL Decentration, IOL Power Miscalculation, Retinal Detachment	FSC-USA, EC Certificate	Abyôgv`b Kiv †h‡Z cv‡i	Abyôgv`b Kiv nj
179.	Valeant Med Sp. Z.o.o. Poland **Local agent :** Janata Traders TCB Bhabon , 1 kawran Bazar,Dhaka	**FOCUSForce** Basic F260 ULTRAflex UF60125 ULTRAflex UF60130 Aspheric AS60125 Aspheric AS60130 Re-vision A100 **Class: C**	Intraocular Lense	FOCUSforce foldable posterior chamber IOLs are indicated for primary implantation for the visual correction of aphakia in adult patients with in whom a cataractous lens has been removed. These lenses are intended for placement in the capsular bag.	**Contraindications :** To date there are no absolute contraindications to Intraocular Lenses' implantation. Relative contra-indications include some forms of: • Chronic active uveitis • Retinal diseases in which the implant may interfere with retinal surgery **Side Effect :** Cataract surgery, with or without lens implantation might be associated with: • Ocular inflammation • Hemorrhage • Intraocular pressure elevation • Post-operative infection • Retinal breaks and detachment • Cystoid macular edema • Corneal edema • Posterior capsule opacity	FSC-Poland EC Certificate (Turkey)	Abyôgv`b Kiv †h‡Z cv‡i	Abyôgv`b Kiv nj
180.	**Manufacturer:** Poly Medicure Ltd.	CVP Manometer	CVP Manometer	It is used for continuous or intermittent monitoring of	**Contraindications:** None	FSC-India	Abyôgv`b Kiv †h‡Z cv‡i	Abyôgv`b Kiv nj

	Regd Office: 1st Floor, 12 Sant Nagar, East of Kailash, New Delhi-110 065 India. Industry: Plot No. 104-105 & 115-116, Sector-59, HSIIDC Industrial Area, Ballabhagra-121004, Faridbad (HARYANA) India **Local Agent:** Aglow International, 114, 1st Floor, Aziz Co-operative Super Market, Shahbag, Dhaka.	**Class: B**		central venous pressure during infusion.	**Side Effects:** None	& EC Certificate		
181.	**Manufacturer:** Ningbo Boya Medical Equipment Co. Ltd, No. 102, Jingsan Road, Yaobei Industrial Part, YuYao City, Zhejiang Province, China. **Local Agent:** Pride Medical Appliances Ltd., 19, Indira Road, Farmgate, Dhaka.	TornadoCare Disposable Nebulizer Mask Class; **Class: B**	Disposable Nebulizer Mask	It is used for bacteria, particle filtration in breathing machine and anesthesia machine and to increase the gas moisture degree	**Contraindication:** Product should not use after validity period. Do not use if the package is damaged. Not use for different user. **Side Effects:** None	FSC China	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
182.	**Manufacturer:** Changzhou Chuangjia Medical Appliance Co. Ltd, Sanhekou Development Zone, Zhenglu, Town, Zhangzhou 213115, China. **Local Agent:** Pride Medical Appliances Ltd., 19, Indira Road, Farmgate, Dhaka.	TornadoCare Oxygen Masks **Class: B**	Oxygen Masks	This product is used for connecting to the oxygen system for medical units with drug for the treatment of patients with respiratory illness.	**Contraindication:** Do not use if the package is damaged. For single use only. Do not use if the package is opened for long time. **Side Effects:** None	FSC China	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
183.	**Manufacturer:** B L Lifesciences Pvt. Ltd. 28-D, Sector-31, Ecotech-I, Greater Noida, Gautam Budh Nagar, U.P., India	**AngiX** Puncture Needle (18G x 7cm / 20G x 4cm / 21G x 7cm)	Accessories of Angio Kit / PTCA Kit	It is indicated for use during Cardiac & Vascular Catherization in order to diagnose the possible blockage in an artery or vessel.	**Contraindication:** This device is not designed, sold, or intended for use except as indicated. **Side Effect**: None	FSC-India EC Certificate (Norway)	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

	Local Agent: Inogen Systems Cha-73/2, Progati Sarani, North Badda, Dhaka-1212	**Class: C**						
184.	**Legal manufacturer:** M/s Becton Dickinson & Company, 1 Becton Drive, Franklin lakes, New Jersey,07417, USA **Manufacturing Site:** *M/s* BD Medical Surgical, 2153, 12th Avenue, Columbus, NE 68601, USA **Local Agent:** M/s Becton Dickinson & Company, 80, Kakrail, Dhaka	BD Blunt Fill Needle, BD Blunt Fill Needle with Filter **Class: B**	Hypodermic Needles	This is a specialized category of needle which is used for aspiration of fluids from vails and ampoules for syringe filling. They are not intended for skin injection. The filter in the liter needles is intended to reduce larger particulates when aspirating, reconstituting, and transferring fluids or when withdrawing medication for glass reservoirs	**Contraindications:** None **Side-effects:** none		Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |
185.	**Manufacturer:** William A Cook Australia Pvt. Ltd., Australia **Local Agent:** Nature and Nurture Ltd. House-19, Road-1, Block-A, Banasree, Dhaka	IVF Needle **Class: B**	ovum aspiration Needle	It is intended for use in the field of assisted reproduction and gynaecology, in particular aspiration of ovarian follicles for the collection of oocytes, and aspiration of pathological ovarian cysts.	**Contraindications:** None. **Side effects:** None	FSC- Australia	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |
186.	**Manufacturer:** Artsana S.P.A Saldarini Catelli 6/10, 22070 Casnate Con Bernate (CO), Italy **Local Agent:** Zas Corporation	PIC Insupen	Sterile disposable pen needle	it is used for subcutaneous insulin therapy with insulin pen.	**Contraindication:** None **Side effect:** None	FSC-Italy	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |

	80/22 Mymensingh Road Nurjehan Tower (3rd Floor) Dhaka-1000. Bangladesh.							
187.	**Manufacturer**: Occlutech GmbH Wildenbruchstrasse 15 07745 Jena Germany. **Local Agent:** Medi Card Ltd. ¼, Paribagh, Mymensingh Road, Dhaka.	Occlutech PDA Occluder **Class: D**	PDA Occluder	It is used as an occlusion system, which is percutaneously implanted through a catheter intervention and intended for the non-surgical occlusion of Patent Ductus Arteriosus (PDA)	**Contraindication**: Contraindicated in case of coagulation disorder. **Side effect**: Hemorrhages. Hemolysis	FSC Germany EC design Examination Certificate	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
188.	Manufacturer: Occlutech GmbH Wildenbruchstrasse 15 07745 Jena Germany. **Local Agent:** Medi Card Ltd. ¼, Paribagh, Mymensingh Road, Dhaka.	Figulla Flex II Occluders (ASD) **Class: C**	Figulla Flex II Occluders (ASD)	It is intended for patients with substantiated left-right-shunt and a clear indication for closure of the defect.	**Contraindication**: Contraindicated: Incase of clotting disorder, atrial tumour and clot. **Side effect:** Clot formation on one of the two discs, infection in the area of the implant.	FSC Germany EC design Examination Certificate	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
189.	Manufacturer: Occlutech GmbH Wildenbruchstrasse 15 07745 Jena Germany. **Local Agent:** Medi Card Ltd. ¼, Paribagh, Mymensingh Road, Dhaka.	Occlutech Delivery Set **Class: C**	Delivery Set	It is used as an accessory for percutaneous catheter Interventions. It is used to implant the oclutech occluder.	**Contraindication**: Contraindicated in case of local skin infection, vascular injury. **Side effect:** Perforation of blood vessels and/or the heart. AV fistula, Infection	FSC Germany EC design Examination Certificate	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
190.	**Manufacturer:** Sree Umiya Surgical Pvt. Ltd. Plot o. 4704, Phase. IV, G.I.D.C, VATA, Ahmedabad. India. **Local Agent:** Hard Won Power Industries Ltd. House No. 39, Road-12, Shekhertek, Adabor. Dhaka.	Mediplus Twin Bore Nasal Oxygen Set **Class: C**	Oxygen Set made with N.T. PVC Tubing "Y" connecter, "T" type parts & cover.	It is used as star lumen tube ensures the supply of oxygen.	**Contraindication:** None **Side Effect**: None	FSC India, EC Certificate	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
191.	**Manufacturer:** G. Surgiwear Limited Rasoolpur Jahanganj, Near Hathoda Chauraha, Shahjahanpur-, INDIA	SURGIWEAR **G-Patch** (Small / Medium / Large)	Patch for repair of Dura	It is indicated for repair of Duramater, repair of peritoneum or places, where ever soft tissue repair is required.	**Contraindications:** Presence of all kinds of infection (meningitis, ventriculitis, skin infection, bacteremia, septicemia and peritonitis etc.) whether local or general, prohibits all kind of	FSC-India	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj

	Local Agent: Inogen Systems Cha-73/2, Progati Sarani, North Badda, Dhaka-1212	**Class: D**			implantation procedures. It is advisable to avoid implantation in such cases. G-Patch is not a strong fabric. It should not be used in places where large tensile forces are expected. In such places stitch may not hold and it may give way and torn. **Side Effects:** The principal complications associated with "Synthetic Fabric Patch" implantation are functional failure and Infection.			
192.	Manufacturer: Poly Medicure Ltd. Regd Office: 1st Floor, 12 Sant Nagar, East of Kailash, New Delhi-110 065 India. Industry: Plot No. 104-105 & 115-116, Sector-59, HSIIDC Industrial Area, Ballabhagra-121004, Faridbad (HARYANA) India. **Local Agent:** Aglow International, 114, 1st Floor, Aziz Co-operative Super Market, Shahbag, Dhaka.	Poly vol Burette Set **Class: B**	Measured volume disposable perfusion set	It is used to administer intravenous fluid and medicines into human circulating system by using intravenous catheter or cannula.	**Contraindications:** Blood transfusion, administration of highly viscous fluids. Not to be used in patients with known hypersensitivity to any of the materials used. **Side Effects:** None.	FSC-India EC Certificate	*Abygv`b Kiv †h‡Z cv‡i |*	*Abygv`b Kiv nj |*
193.	Manufacturer: Changzhou Operson IMP. and Exp. Co. Ltd. China. **Local Agent:** Shanto Enterprise, 51, Islampur Road, Dhaka.	Greenpas **Class: B**	Medical Plaster	Plaster are used in many different ways. Temporary medical casts will immobilize a limb prior to the application of a permanent cast. They are also used as art suppies and can be used to make pregnancy belly	**Contraindication:** None **Side Effect**: None	FSC-China	*Abygv`b Kiv †h‡Z cv‡i |*	*Abygv`b Kiv nj |*

				casts, face masks, dress molds, life casting body molds, and mountains for model train sets				
194.	**Manufacturer:** Artsana S.P.A Saldarini Catelli 6/10, 22070 Casnate Con Bernate (CO), Italy **Local Agent:** Zas Corporation 80/22 Mymensingh Road Nurjehan Tower (3rd Floor) Dhaka-1000. Bangladesh.	AQUABLOC **Class: C**	Sterile waterproof Plaster with antibacterial pad	It is used for protection and treatment of injured skin and mucous.	**Contraindication:** None **Side effect:** None	FSC-Italy	Abjygv`b Kiv †h‡Z cv‡i |	Abjygv`b Kiv nj |
195.	**Manufacturer:** Artsana S.P.A Saldarini Catelli 6/10, 22070 Casnate Con Bernate (CO), Italy **Local Agent:** Zas Corporation 80/22 Mymensingh Road Nurjehan Tower (3rd Floor) Dhaka-1000. Bangladesh.	Soffix Med **Class: C**	Delicate Sterile post operative plaster with antibacterial pad	It is used for all woundcare needs from trauma wounds to post-surgery.	**Contraindication:** None **Side effect:** None	FSC-Italy	Abjygv`b Kiv †h‡Z cv‡i |	Abjygv`b Kiv nj |
196.	**Manufacturer:** B. Braun Melsungen AG Carl-Barun-StraBe1, Melsungen Germany. **Local Agent:** Asia Pacific Medicals Ltd. 775, Satmasjid Road (2nd Floor), Dhanmodi, Dhaka.	Proxima 2 Bag **Class: B**	Colostomy Bag/Drainable Pouch	outlet designed to reduce leakage and easy to connect to a drainage **bag**	**Contraindication:** None **Side Effect**: None	FSC Germany	Abjygv`b Kiv †h‡Z cv‡i |	Abjygv`b Kiv nj |
197.	**Manufacturer:** B L Lifesciences Pvt. Ltd., 28-D, Sector-31,	iPeX **Pressure Monitoring Kit**	Pressure Monitoring Kit	It is used for blood pressure sensing during clinical invasive procedures without interfering	**Contraindications:** Pressure monitoring should not be monitored without using an air-eliminating filter	FSC-India	Abjygv`b Kiv †h‡Z cv‡i |	Abjygv`b Kiv nj |

	Ecotech-I, Greater Noida, Gautam Budh Nagar, U.P., India **Local Agent:** Inogen Systems Cha-73/2, Progati Sarani, North Badda, Dhaka-1212	(Single/Double / Triple) **Class: B**		with blood flow and/or pressure. iPeX Pressure Monitoring Kits are used to continuously monitor Arterial Blood Pressure, Central Venous Pressure, Intra-Cardiac Pressure, Intra-Cranial Pressure, Intra-Uterine Pressure, Pulmonary Artery Pressure or Compartment Pressure.	to clear air passage between solution source and flush device. **Side Effect**: None			
198.	**Manufacturer:** Sree Umiya Surgical Pvt. Ltd. Plot o. 4704, Phase. IV, G.I.D.C, VATA, Ahmedabad. India. **Local Agent:** Hard Won Power Industries Ltd. House No. 39, Road-12, Shekhertek, Adabor. Dhaka.	Mediplus Pressure Monitoring Line **Class: C**	Pressure Monitoring Line made with N.T. PVC Tubing & Luer Lock	It is used as multipurpose extension line suitable for Angiography, Arteriography.	**Contraindication:** None **Side Effect**: None	FSC India, EC Certificate	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
199.	**Manufacturer:** B L Lifesciences Pvt. Ltd., 28-D, Sector-31, Ecotech-I, Greater Noida, Gautam Budh Nagar, U.P., India **Local Agent:** Inogen Systems Cha-73/2, Progati Sarani, North Badda, Dhaka-1212	iPeX **Disposable Pressure Transducer** (With / Without Flush) **Class: B**	Pressure Monitoring Product	It is used for blood pressure sensing during clinical invasive procedures without interfering with blood flow and/or pressure. iPeX Pressure Monitoring Kits are used to continuously monitor Arterial Blood Pressure, Central Venous Pressure, Intra-Cardiac Pressure, Intra-Cranial Pressure, Intra-Uterine Pressure, Pulmonary Artery Pressure or Compartment Pressure.	**Contraindications:** Pressure monitoring should not be monitored without using an air-eliminating filter to clear air passage between solution source and flush device. **Side Effect**: None	FSC-India EC Certificate (Norway)	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
200.	**Manufacturer:** B L Lifesciences Pvt. Ltd. 28-D, Sector-31, Ecotech-I, Greater Noida, Gautam Budh Nagar,	AngiX **Hemostasis Y** (Standard / Click Type) **Class: C**	Accessories of Angio Kit / PTCA Kit	It is indicated for use during Cardiac & Vascular Catherization in order to diagnose the possible blockage in an artery or vessel.	**Contraindication:** This device is not designed, sold, or intended for use except as indicated. **Side Effect**: None	FSC-India EC Certificate (Norway)	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

	U.P., India **Local Agent:** Inogen Systems Cha-73/2, Progati Sarani, North Badda, Dhaka-1212							
201.	**Manufacturer:** B L Lifesciences Pvt. Ltd. 28-D, Sector-31, Ecotech-I, Greater Noida, Gautam Budh Nagar, U.P., India **Local Agent:** Inogen Systems Cha-73/2, Progati Sarani, North Badda, Dhaka-1212	**AngiX** Manifold (2 Core / 3 Core / 4 Core) **Class: C**	Accessories of Angio Kit / PTCA Kit	It is indicated for use during Cardiac & Vascular Catherization in order to diagnose the possible blockage in an artery or vessel.	**Contraindication:** This device is not designed, sold, or intended for use except as indicated. **Side Effect**: None	FSC-India EC Certificate (Norway)	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
202.	**Manufacturer:** B L Lifesciences Pvt. Ltd. 28-D, Sector-31, Ecotech-I, Greater Noida, Gautam Budh Nagar, U.P., India **Local Agent:** Inogen Systems Cha-73/2, Progati Sarani, North Badda, Dhaka-1212	AngiX **Inflation Device** **Class: C**	Accessories of Angio Kit / PTCA Kit	It is indicated for use during Cardiac & Vascular Catherization in order to diagnose the possible blockage in an artery or vessel.	**Contraindication:** This device is not designed, sold, or intended for use except as indicated. **Side Effect**: None	FSC-India EC Certificate (Norway)	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
203.	**Manufacturer:** G. Surgiwear Limited Rasoolpur Jahanganj, Near Hathoda Chauraha, Shahjahanpur-242001, U.P., India **Local Agent:** Inogen Systems Cha-73/2, Progati Sarani, North Badda, Dhaka-1212	SURGIWEAR **Cerebral Catheter Reservoir** (Small / Large) **Class: D**	Cerebral Reservoir	It is used for frequent intraventricular medications and CSF sampling.	**Contraindications:** Presence of all kinds of infection (meningitis, ventriculitis, skin infection, bacteremia, septicemia and peritonitis etc.) whether local or general, prohibits all kind of implantation procedures. It is advisable to avoid implantation in such cases. **Side Effects:** The principal complications associated with 'External Drainage System' implantations are obstruction, functional failure of system and infection.	FSC-India	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
204.	**Manufacturer:** Poly Medicure Ltd.	Poly Respiciser, Respiratory Exerciser	Poly Respiciser, Respiratory Exerciser	This product is used to help patient, to improve the	**Contraindications:** Not to be used without supervision of highly	FSC- India	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

	Regd Office: 1st Floor, 12 Sant Nagar, East of Kailash, New Delhi-110 065 India. Industry: Plot No. 104-105 & 115-116, Sector-59, HSIIDC Industrial Area, Ballabhagra-121004, Faridbad (HARYANA) India. **Local Agent:** Aglow International, 114, 1st Floor, Aziz Co-operative Super Market, Shahbag, Dhaka.	**Class: B**		functioning of their lungs in conditions where the lung functions are compromised due to various respiratory disorders such as "atelectasis (small airway collapse), bronchial asthma"	qualified medical practitioner. **Side Effects:** None			
205.	**Manufacturer:** G. Surgiwear Limited Rasoolpur Jahanganj, Near Hathoda Chauraha, Shahjahanpur-242001, U.P., INDIA **Local Agent:** Inogen Systems Cha-73/2, Progati Sarani, North Badda, Dhaka-1212	SURGIWEAR **Chhabra "Slit N Spring" Hydrocephalus Shunt System** (Regular Reservoir: VPHP / VPMP / VPLP / VAHP / VAMP / VALP; Large Flushing Reservoir: VPHP / VPMP / VPLP) **Class: D**	Hydrocephalus Shunt	It is to be used for treatment of hydrocephalic patients, for shunting of CSF from lateral ventricles of brain to peritoneum or rt. atrium.	**Contraindications:** Presence of all kinds of infection (meningitis, ventriculitis, skin infection, bacteremia, septicemia and peritonitis etc.) whether local or general, prohibits all kind of implantation procedures. It is advisable to avoid shunt implantation in such cases. **Side Effects:** The principal complications associated with cerebrospinal fluid shunting into the peritoneum are shunt obstruction, functional failure of shunt system, infection and intracranial hypotension**.**	FSC-India	*Abygv`b Kiv †h‡Z cv‡i |*	*Abygv`b Kiv nj |*
206.	**Manufacturer:** G. Surgiwear Limited Rasoolpur Jahanganj, Near Hathoda Chauraha, Shahjahanpur-242001, U.P., INDIA **Local Agent:** Inogen Systems Cha-73/2, Progati Sarani, North Badda, Dhaka-1212	SURGIWEAR **Chhabra "Slit N Spring" Valve Only Hydrocephalus Shunt System** (With **/** Without Large Flushing Reservoir**:** High Pressure / Medium Pressure / Low Pressure)	Hydrocephalus Shunt parts & accessories	It is to be used for treatment of hydrocephalic patients, for shunting of CSF from lateral ventricles of brain to peritoneum or rt. atrium.	**Contraindications:** Presence of all kinds of infection (meningitis, ventriculitis, skin infection, bacteremia, septicemia and peritonitis etc.) whether local or general, prohibits all kind of implantation procedures. It is advisable to avoid shunt implantation in such cases. **Side Effects:** The principal complications associated with	FSC-India	*Abygv`b Kiv †h‡Z cv‡i |*	*Abygv`b Kiv nj |*

		Class: D			cerebrospinal fluid shunting into the peritoneum are shunt obstruction, functional failure of shunt system, infection and intracranial hypotension**.**			
207.	**Manufacturer:** G. Surgiwear Limited Rasoolpur Jahanganj, Near Hathoda Chauraha, Shahjahanpur-242001, U.P., INDIA **Local Agent:** Inogen Systems Cha-73/2, Progati Sarani, North Badda, Dhaka-1212	SURGIWEAR **GSL "Dome Valve" Hydrocephalus Shunt System** (High Pressure / Medium Pressure / Low Pressure) **Class: D**	Hydrocephalus Shunt	It is to be used for treatment of hydrocephalic patients, for shunting of CSF from lateral ventricles of brain to peritoneum or rt. atrium	**Contraindications:** Presence of all kinds of infection (meningitis, ventriculitis, skin infection, bacteremia, septicemia and peritonitis etc.) whether local or general, prohibits all kind of implantation procedures. It is advisable to avoid shunt implantation in such cases. **Side Effects:** The principal complications associated with cerebrospinal fluid shunting into the peritoneum are shunt obstruction, functional failure of shunt system, infection and intracranial hypotension**.**	FSC-India	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
208.	**Manufacturer:** G. Surgiwear Limited Rasoolpur Jahanganj, Near Hathoda Chauraha, Shahjahanpur-242001, U.P., INDIA **Local Agent:** Inogen Systems Cha-73/2, Progati Sarani, North Badda, Dhaka-1212	SURGIWEAR **Burr Hole Cover** **Class: D**	Hydrocephalus Shunt parts & accessories	It is to be used for treatment of hydrocephalic patients, for shunting of CSF from lateral ventricles of brain to peritoneum or rt. atrium	**Contraindications:** Presence of all kinds of infection (meningitis, ventriculitis, skin infection, bacteremia, septicemia and peritonitis etc.) whether local or general, prohibits all kind of implantation procedures. It is advisable to avoid shunt implantation in such cases. **Side Effects:** The principal complications associated with cerebrospinal fluid shunting into the peritoneum are shunt	FSC-India	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

					obstruction, functional failure of shunt system, infection and intracranial hypotension**.**			
209.	**Manufacturer:** G. Surgiwear Limited Rasoolpur Jahanganj, Near Hathoda Chauraha, Shahjahanpur-242001, U.P., INDIA **Local Agent:** Inogen Systems Cha-73/2, Progati Sarani, North Badda, Dhaka-1212	SURGIWEAR **Chhabra "Lumber Peritoneal" Hydrocephalus Shunt System** (With / Without Tuohy Needle) **Class: D**	Lumber Peritoneal Shunt	It is to be used for treatment of hydrocephalic patients, for shunting of CSF from lumber region of spinal cord to peritoneum.	**Contraindications:** Presence of all kinds of infection (meningitis, ventriculitis, skin infection, bacteremia, septicemia and peritonitis etc.) whether local or general, prohibits all kind of implantation procedures. It is advisable to avoid shunt implantation in such cases. **Side Effects:** The principal complications associated with cerebrospinal fluid shunting into the peritoneums are shunt obstruction, functional failure of shunt system, infection and intracranial hypotension**.**	FSC-India	*Abygv`b Kiv †h‡Z cv‡i |*	*Abygv`b Kiv nj |*
210.	**Manufacturer:** Chase Medical 885 E Collins Blvd Ste 110, RICHARDSON, TX USA 75081 **Local Agent:** Inogen Systems Cha-73/2, Progati Sarani, North Badda, Dhaka-1212	Chase **Blood Vessel Shunt** (Bulb tip 12 mm shaft: 1.50 mm, 1.75 mm, 2.00 mm, 2.50 mm, 3.00 mm; Bulb tip 30 mm shaft: 2.00 mm, 2.50 mm, 3.00 mm, 4.00 mm; Tapered tip 12 mm shaft: 1.00 mm, 1.25 mm, 1.50 mm, 1.75 mm, 2.00 mm, 2.50 mm) **Class: D**	Coronary Artery Shunts	It is used to internally shunt blood vessels during anastomosis. The shunt valves allow a dry field to be maintained during a vessel anastomosis procedure. The shunt also allows the blood to be delivered distally past the anastomosis during vessel repair. The blood vessel shunt is easily removed prior to the final sutures being tied.	**Contraindications:** This device is not intended for use other than as indicated. **Side-effects:** none	FSC-USA	*Abygv`b Kiv †h‡Z cv‡i |*	*Abygv`b Kiv nj |*

211.	Bausch & Lomb INC, USA **Local agent :** Janata Traders TCB Bhabon , 1 kawran Bazar,Dhaka	Bausch + Lomb renu fresh multi-purpose solution **Class: C**	Contact lens care Solution	It is indicated for use in the daily cleaning, removal of protein deposits, rinsing, chemical (not heat) disinfection, and storage of soft (hydrophilic) contact lenses as recommended by your eye care professional.	**Contraindications :** If you are allergic to any ingredient in this product, do not use. **Side Effect:** The following problems may occur: eyes sting, burn or itch (irritation), comfort is less than when lens was first placed on the eye, feeling of something in the eye (foreign body, scratched area), excessive watering (tearing) of the eye, unusual eye secretions, redness of the eye, reduced sharpness of vision (poor visual acuity), blurred vision, rainbows or halos around objects, sensitivity to light (photophobia), or dry eyes. If you notice any of the above: Immediately remove your lenses. • If the discomfort or problem stops, then look closely at the lens. • If the lens is in any way damaged, do not put the lens back on your eye. Place the lens in the storage case and contact your eye care professional.	FSC-USA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
212.	Manufacturer: Almediko Saglik, Turkey. **Local Agent:** Halifax Traders Ltd. 30/D (Level-6), Satish Sarkar Road, Gandaria, Dhaka.	Memethol Briyer Spray **Class: B**	Memethol Briyer Spray	It is used to resolves pain and itching that the most important symptoms of hemorrhoids.	**Contraindication:** None **Side Effect**: None	FSC Turkey, EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
213.	**Manufacturer:** Meril Endo Surgery PVT. Ltd. India. **Local Agent:** Meril Bangladesh PVT. Ltd. Union Heights-1 13th Floor, Level-14, 55/2, Bir Uttam Qazi	Mirus Endo Cutter Stapler & its cartridges **Class: B**	Sterile endoscopic linear cutter & reloads (Staples)	The Endo Cutter Staplers have applications in abdominal, gynecologic, pediatric and thoracic surgery for resection, transection and creation of anastomosis. Staple shape height depends on the	**Contraindications:** 1. It should not be used on any tissue that compresses to less than .75 mm in thickness, on any tissue that cannot comfortably compress to	FSC India EC Certificate (Norway)	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

	Nuruzzaman Saraka, West Panthapath, Dhaka			specification of single use loading unit (SULU). Always inspect the tissue thickness and select an appropriate staple size prior to application of the Endo cutter stapler.	1.0 mm or on the aorta. 2. Endo Cutter Stapler 2.5 mm staples should not be used on any tissue that compresses to less than l.0 mm in thickness, on any tissue that cannot comfortably compress to 1.5 mm or on the aorta. 3. Endo Cutter Stapler 3.5 mm staples should not be used on any tissue that compresses to less than l.5 mm in thickness, on any tissue that cannot comfortably compress to 2.0 mm or on the aorta. 4. Endo Cutter Stapler 4.8 mm staples should not be used on any' tissue that compresses to less than 2.0mm in thickness, on any tissue that cannot comfortably compress to 2.0 mm or on the aorta. 5. It should not be used on tissue such as liver or spleen where compressibility is such that closure of the instrument would be destructive. 6. These devices are provided STERILE and are intended for use in a Single procedure only. **Side effects:** none			
214.	Manufacturer: Meril Endo Surgery PVT. Ltd. India. **Local Agent:** Meril Bangladesh PVT. Ltd. Union Heights-1 13th Floor, Level-14, 55/2, Bir Uttam Qazi Nuruzzaman Saraka, West Panthapath, Dhaka.	Mirus Linear Stapler (Without dial) **Class: B**	Linear Stapler (Without dial) & Stapler Reloads (Staples)	They have applications in alimentary tract, lungs and bronchi for closure and transection of tissue.	**Contraindications:** 1. It should not be used on any tissue that compresses to less than .75 mm in thickness, on any tissue that cannot comfortably compress to 1.0 mm or on the aorta. 2. Endo Cutter Stapler 2.5 mm staples should not be used on any tissue that compresses to less than	FSC India EC Certificate (Norway)	*Abţgv`b Kiv †h‡Z cv‡i |*	*Abţgv`b Kiv nj |*

					l.0 mm in thickness, on any tissue that cannot comfortably compress to 1.5 mm or on the aorta. 3. Endo Cutter Stapler 3.5 mm staples should not be used on any tissue that compresses to less than l.5 mm in thickness, on any tissue that cannot comfortably compress to 2.0 mm or on the aorta. 4. Endo Cutter Stapler 4.8 mm staples should not be used on any' tissue that compresses to less than 2.0mm in thickness, on any tissue that cannot comfortably compress to 2.0 mm or on the aorta. 5. It should not be used on tissue such as liver or spleen where compressibility is such that closure of the instrument would be destructive. 6. These devices are provided STERILE and are intended for use in a Single procedure only. **Side effects:** none			
215.	Manufacturer: Meril Endo Surgery PVT. Ltd. India. **Local Agent:** Meril Bangladesh Pvt. Ltd. Union Heights-1 13th Floor, Level-14, 55/2, Bir Uttam Qazi Nuruzzaman Saraka, West Panthapath, Dhaka.	Mirus Circular Stapler 3 rows & its cartridges **Class: B**	Circular Stapler & its cartridges	It has applications throughout the alimentary tract for the creation of end-to-end, end-to-side and side-to-side anastomosis in both open and endoscopic surgeries.	**Contraindications:** 1. It should no be used on any tissue that can be compressed to less than 2 mm in thickness, as the staples will not be tight enough to ensure hemostasis. 2. It should not be used on any tissue that cannot be comfortably compressed to 2 mm in thickness. The closure of the cartridge and anvil could crush overly thick inverted tissue and result in failure to create an anastomosis, poor healing, or narrowing of the anastomosis. 3. In those cases that by introducing a cartridge whose diameter is too large for the structure, the tissue may be stretched or thinned out, It should not be used, or it could result in leakage and narrowing of the anastomosis. 4. unless sufficient tissue presents that	FSC India EC Certificate (Norway)	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

					allows proper inversion of tissue edges and secures staples placement. **Side effects:** none			
216.	**Manufacturer:** Balton S.PZ.O.O Nowy Swiat 7/14 Street, 00-496 Warsaw, Poland, **Local Agent:** Advanced Meditech Suite: 115 (1st Floor), Krishnachura Commercial Complex, 24/B.C Shahid Minar Road, Kallanpur, Dhaka.	Peripheral Stent Neptun With Delivery System **Class: D**	Peripheral Stent	Stent for Peripheral Vessel Neptun is used in the following Cases: - Unsatisfactory effect of a procedure (PTA)/residual stenosis, - Internal delamination of tunica intima or vessel lesion, - Recurrent Stenosis or loss of patency following previous intravascular procedures. -Outer presser exerted on the vessel.	**Contraindication:** Insufficient flow-out of blood below the potential stent implantation spot. - Impossibility of a guide wire and balloon catheter passing through the stenosed vessel or arterial occlusion. - Hyper coagulity reported in anamnesis. - Possibility of closing collateral circulation vessels with a stent. - External heavily calcified atherosclerotic lamina , which could damage the stent. - Fresh Thrombus. - Contraindications against taking antithrombotic medicines/ active bleeding from the digestive tract, recently suffered cerebral stroke. - Allergy to contrast media. - Allergy to stainless steel. **Side Effects:** - Vessel puncture - Ablation of tunica intima in a vessel. - Arterial Spasm. - Recent Thrombus inside the implanted stent. - Peripheral thrombosis. - Bleeding and haematoma in the needle insertion site. - False aneurysm in the needle insertion site. - Stent migration. -Allergic reaction to contrast media and stainless steel.	FSC-Poland	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |
217.	**Manufacturer:** Balton S.PZ.O.O Nowy Swiat 7/14 Street, 00-496 Warsaw, Poland **Local Agent:** Advanced Meditech Suite: 115 (1st Floor), Krishnachura Commercial Complex, 24/B.C Shahid Minar Road,	**MER** Carotid self expanding Stent with delivery System **Class: D**	Carotid Stent with delivery System	It is indicated for the treatment of patients who require carotid revascularization and undergoing surgical carotid endarterectomy poses a high risk for adverse events: Patients with neurological symptoms and ≥ 50% stenosis of the common or internal carotid artery visualized by	**Contraindications:** - Contraindications against taking antiplatelet or antithrombotic medicines, - Uncorrected bl eeding disorders, - Hypercoagulability reported in anamnesis, - Active or recent bleeding from the gastrointestinal tract,	FSC-Poland	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |

	Kallanpur, Dhaka.			USG or angiography; Patients without neurological symptoms and ≥ 80% stenosis of the common or internal carotid artery visualized by USG or angiography.	- Recently suffered hemorrhagic or large ischemic cerebral stroke, - Known hypersensitivity to nickel-titanium, - Known hypersensitivity to contrast media, - Severe vascular tortuosity or anatomy that would preclude the safe introduction of a guide catheter, sheath, embolic protection system, or stent system, - Extremely heavily calcified atherosclerotic lamina, which could damage the stent, - Lesions located in the ostium of the common carotid artery, - Fresh thrombus, - Total occlusion of target carotid artery. **Complications/Side effect:** The device is destined exclusively for single use. The device must not be resterilized and/or reused! Do not use if the package is open or damaged. Do not use after expiry date stated on the package. Protect against organic solvents action (e.g. alcohol). While advancing the catheter through the artery, it may become damaged and the life threatening bleeding may occur. The safety and effectiveness of Balton carotid self-expanding stent with delivery system have NOT yet been determined in patients with the following characteristics: Pregnant patients or patients under the age of 18, Prior stenting of the			

					target carotid artery, Myocardial infarction within 72 hours prior to the procedure, CABG or vascular surgery within 30 days prior to the procedure, Major residual neurological deficit (stroke scales: Barthel< 60, NIH > 15 or Rankin > 3) at the periprocedural neurological exam, TIA within 48 hours prior to the procedure, Current radiotherapy of cerebral tumor.			
218.	**Manufacturer:** Balton S.PZ.O.O Nowy Swiat 7/14 Street, 00-496 Warsaw, Poland **Local Agent:** Advanced Meditech Suite: 115 (1st Floor), Krishnachura Commercial Complex, 24/B.C Shahid Minar Road, Kallanpur, Dhaka.	**Nefro** Renal Stent with delivery System **Class: C**	**Nefro** Renal Stent with delivery System	Stent for renal Vessels Nefro is used in the following Cases: - Unsatisfactory effect of a procedure (PTA)/residual stenosis - Stenosis at the ostium of the renal artery - Recurrent Stenosis or loss of patency following previous intravascular procedures. -Revascularisation of an occluded renal artery.	**Contraindications**: - Insufficient flow-out of blood below the potential stent implantation spot. - Impossibility of a guide wire and balloon catheter passing through the stenosed vessel or arterial occlusion. - Hyper coagulity reported in anamnesis. - Possibility of closing collateral circulation vessels with a stent. - External heavily calcified atherosclerotic lamina , which could damage the stent. - Fresh Thrombus. - Contraindications against taking antithrombotic medicines/ active bleeding from the digestive tract, recently suffered cerebral stroke. - Allergy to contrast media. - Allergy to stainless steel. **Complications/Side effect:** - Artery puncture - Ablation of tunica intima in a vessel. - Arterial Spasm. - Recent Thrombus inside the implanted stent. - Peripheral thrombosis. - Bleeding and haematoma in the	FSC-Poland	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

					needle insertion site. - False aneurysm in the needle insertion site. - Stent migration. -Allergic reaction to contrast media and stainless steel.			
219.	**Manufacturer:** Meril Life Science PVT. Ltd. India. **Local Agent:** Meril Bangladesh PVT. Ltd. Union Heights-1 13th Floor, Level-14, 55/2, Bir Uttam Qazi Nuruzzaman Saraka, West Panthapath, Dhaka.	**MeRes 100** Biogradable Coronary stent system **Class: D**	Sirolimus Eluting Bioresorbable vascular scaffold system	It is indicated for improving coronary luminal diameter in patients with symptomatic ischemic heart disease		FSC-India & EC Certificate (Norway)	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
220.	**Manufacturer:** Meril Life Science PVT. Ltd. India. **Local Agent:** Meril Bangladesh Pvt. Ltd. Union Heights-1 13th Floor, Level-14, 55/2, Bir Uttam Qazi Nuruzzaman Saraka, West Panthapath, Dhaka.	**Cogent BMS** **Class: C**	Balloon Expandable Renal & Biliary stent system	It is indicated for use in patients with atherosclerotic disease of the renal arteries following sub-optimal percutaneous transluminal angioplasty (PTRA) of a de novo or restenotic atherosclerotic lesion & for palliation of malignant neoplasms in the biliary tree with reference vessel diameter of 5.00 mm to 7.00mm in patients eligible for Percutaneous Transluminal Angioplasty (PTA) and Stenting procedures.	**Contraindications:** Contraindicated in the following patient types: - Patients in whom anti-platelet and/or anti-coagulant therapy are contraindicated. - Patients judged to have a lesion that prevents complete inflation of an angioplasty balloon. - Transplant patients **Adverse effects:** Abscess, Allergy to cobalt chromium, Arteriovenous fistula, arrhythmia, Death, dialysis, dissection, Drug reaction to antiplatelet agent, bowel infarct, kidney infarct, ischemia, hypertension/hypotension, myocardial infarction, myocardial ischemia.	FSC-India EC Certificate (Norway)	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
221.	**Manufacturer:** Medtronic Ireland, Parkmore Business Park West, Galway Ireland	Endeavor Resolute **Class C**	Zotarolimus-Eluting Coronary Stent System	It is indicated for improving coronary luminal diameter and reducing restenosis in patients with symptomatic ischemic heart disease in de novo	**Contraindications:** It is contraindicated for use in: • Patients with hypersensitivity or allergies to aspirin, heparin, clopidogrel, ticlopidine, drugs such as zotarolimus, tacrolimus, sirolimus or similar drugs or any other	FSC Ireland	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

	Local Agent: Medtronic Bangladesh Pvt. Ltd, Level 6, Shanta Western Towers 186 Gulshan Tejgaon Link Road Dhaka – 1208, Bangladesh			coronary artery lesions in native coronary arteries with a reference vessel diameter of 2.25 mm to 4.0 mm.	analogue or derivative, polymers, cobalt, chromium, nickel, molybdenum, or contrast media. • Patients in whom antiplatelet and/or anticoagulation therapy is contraindicated. • Patients who are judged to have a lesion that prevents complete inflation of an angioplasty balloon or proper placement of the stent or stent delivery system. **Side effects:** None			
222.	**Manufacturer:** Medtronic Ireland, Parkmore Business Park West, Galway Ireland **Local Agent:** Medtronic Bangladesh Pvt. Ltd, Level 6, Shanta Western Towers 186 Gulshan Tejgaon Link Road Dhaka – 1208, Bangladesh	Endeavor Sprint **Class C**	Zotarolimus-Eluting Coronary Stent System	It is indicated for treatment of single and multi-vessel de novo and in-stent restenotic lesions in patients with coronary artery disease – including acute coronary syndrome (acute myocardial infarction or unstable angina) and/or concomitant diabetes mellitus – to improve luminal diameter and reduce restenosis within the stent and at the stent edges in native coronary arteries, with a reference vessel diameter of 2.25 mm to 4.0 mm and a lesion length of ≤ 27 mm.	**Contraindications:** It is contraindicated for use in: • Patients with hypersensitivity or allergies to aspirin, heparin, clopidogrel, ticlopidine, drugs such as zotarolimus, tacrolimus, sirolimus or similar drugs or any other analogue or derivative, polymers, cobalt, chromium, nickel, molybdenum, or contrast media. • Patients in whom antiplatelet and/or anticoagulation therapy is contraindicated. • Patients who are judged to have a lesion that prevents complete inflation of an angioplasty balloon or proper placement of the stent or stent delivery system. **Side effects:** None	FSC Ireland	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
223.	**Manufacturer:** Medtronic Ireland, Parkmore Business Park West, Galway Ireland **Local Agent:** Medtronic Bangladesh Pvt. Ltd, Level 6, Shanta Western	Resolute Onyx **Class C**	Zotarolimus-Eluting Coronary Stent System	It is intended for use in patients eligible for percutaneous transluminal coronary angioplasty (PTCA) with a reference vessel diameter of 2.0 mm to 5.0 mm. The Resolute Onyx™ Stent is indicated for the treatment of	**Contraindications**: It is contraindicated for use in: • Patients with hypersensitivity or allergies to aspirin, heparin, clopidogrel, ticlopidine, mTOR inhibiting drugs such as zotarolimus (tacrolimus, sirolimus, everolimus) or any other mTOR inhibitor analogue or derivative, polymers, cobalt, chromium, nickel, molybdenum,	FSC Ireland	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

	Towers 186 Gulshan Tejgaon Link Road Dhaka – 1208, Bangladesh			the following patient and lesion subsets: Diabetes Mellitus, Multivessel Disease, Acute Coronary Syndrome (ACS), Acute Myocardial Infarction (AMI), Unstable Angina (UA), Bifurcation Lesions, In-Stent Restenosis (ISR), Chronic Total Occlusions (CTO), Total Occlusions (TO) The Resolute Onyx™ Stent is intended to improve coronary luminal diameters of either single or multiple vessels as an adjunct to coronary interventions and to reduce restenosis. The stent is intended as a permanently implanted device.	platinum and iridium, or contrast media. • Patients in whom antiplatelet and/or anticoagulation therapy is contraindicated. • Patients who are judged to have a lesion that prevents complete inflation of an angioplasty balloon or proper placement of the stent or stent delivery system. **Side effects:** None			
224.	**Manufacturer:** Medtronic Ireland, Parkmore Business Park West, Galway Ireland **Local Agent:** Medtronic Bangladesh Pvt. Ltd, Level 6, Shanta Western Towers 186 Gulshan Tejgaon Link Road Dhaka – 1208, Bangladesh	Resolute Integrity **Class C**	Zotarolimus-Eluting Coronary Stent System	It is intended for use in patients, including those with diabetes mellitus, eligible for percutaneous transluminal coronary angioplasty (PTCA) with a reference vessel diameter of 2.25 mm to 4.0 mm. The Resolute Integrity Stent is intended to improve coronary luminal diameters of either single or multiple vessels as an adjunct to coronary interventions and to reduce restenosis. The stent is intended as a permanently implanted device.	**Contraindications:** It is contraindicated for use in: • Patients with hypersensitivity or allergies to aspirin, heparin, clopidogrel, ticlopidine, mTOR inhibiting drugs such as zotarolimus (tacrolimus, sirolimus, everolimus) or any other mTOR inhibitor analogue or derivative, polymers, cobalt, chromium, nickel, molybdenum, or contrast media. • Patients in whom antiplatelet and/or anticoagulation therapy is contraindicated. • Patients who are judged to have a lesion that prevents complete inflation of an angioplasty balloon or proper placement of the stent or stent delivery system.	FSC Ireland	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
225.	**Manufacturer:** Medtronic Ireland, Parkmore Business Park West, Galway	Integrity BMS **Class C**	Zotarolimus-Eluting Coronary Stent System	It is indicated for treatment of coronary occlusive disease. The Integrity Stent is intended	**Contraindications:** • Patients in whom antiplatelet	FSC Ireland	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

	Ireland **Local Agent:** Medtronic Bangladesh Pvt. Ltd, Level 6, Shanta Western Towers 186 Gulshan Tejgaon Link Road Dhaka – 1208, Bangladesh			for use in patients eligible for Percutaneous Transluminal Coronary Angioplasty (PTCA) with reference vessel diameter of 2.25 – 4.0mm. The stent is intended as a permanently implanted device. Stents may be deployed either singly or in a multiple tandem manner to maintain vessel patency.	and/or anticoagulation therapy is contraindicated. • Patients who are judged to have a lesion that prevents complete inflation of an angioplasty balloon. **Side effects:** None			
226.	**Manufacturer:** Poly Medicure Ltd. Regd Office: 1st Floor, 12 Sant Nagar, East of Kailash, New Delhi-110 065 India. Industry: Plot No. 104-105 & 115-116, Sector-59, HSIIDC Industrial Area, Ballabhagra-121004, Faridbad (HARYANA) India. **Local Agent:** Aglow International, 114, 1st Floor, Aziz Co-operative Super Market, Shahbag, Dhaka.	Polyway **Class: B**	Three way stop cock	To administer two drugs or fluids at the same time	**Contraindications:** - Not to be use in patients with known hypersensitivity to any of the amterials used. - Not to be used in case of highly viscous fluid **Side Effects:** None	FSC-India	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
227.	**Manufacturer:** B. Braun Melsungen AG Carl-Barun-StraBe1, Melsungen Germany. **Local Agent:** Asia Pacific Medicals Ltd. 775, Satmasjid Road (2nd Floor), Dhanmodi, Dhaka.	Discofix **Class: B**	Stop cock system for infusion therapy	It is used for infusion therapy.	**Contraindication:** None **Side Effect**: None	FSC Germany	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
228.	**Manufacturer:** Sree Umiya Surgical Pvt. Ltd. Plot o. 4704, Phase. IV, G.I.D.C, VATA, Ahmedabad. India. **Local Agent:**	Mediplus Three Way Stopcock **Class: C**	Three Way Stop cock made with Male Luer Lock & N.T. PVC. Tubing	It is used to facilitates multiple line thought single IV access.	**Contraindication:** None **Side Effect**: None	FSC India, EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

	Hard Won Power Industries Ltd. House No. 39, Road-12, Shekhertek, Adabor. Dhaka.							
229.	**<u>Legal Manufacturer:</u>** **Becton Dickinson Infusion Therapy** AB, Florettgatan 29C, PO Box 631, SE-251 06 Helsingborg, Sweden **<u>Manufacturing Site:</u>** **Becton Dickinson Infusion Therapy S**ystems Inc., S.A. de C.V., Periferico Luis Donaldo Colosio #579, Nogales, Sonora, C.P. 84048 **Local Agent:** M/s Becton Dickinson & Company, 80, Kakrail, Dhaka	BD Connecta **Class: B**	Stopcock	This is a specialized category of Connecta is used for the general purpose of supping the circulatory system with fluids or medication from one or two different sources via an IV cannula or extension tube. It is also used in infusion therapy and hemodynamic pressure monitoring.	**Contraindications:** None **Side-effects:** none		Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj	
230.	**Manufacturer:** Poly Medicure Ltd. Regd Office: 1st Floor, 12 Sant Nagar, East of Kailash, New Delhi-110 065 India. Industry: Plot No. 104-105 & 115-116, Sector-59, HSIIDC Industrial Area, Ballabhagra-121004, Faridbad (HARYANA) India. **Local Agent:** Aglow International, 114, 1st Floor, Aziz Co-operative Super Market, Shahbag, Dhaka.	Yankaur suction set with standard tip **Class: B**	Yankaur suction set with standard tip	- To clear the airway for intubation or removal of secretion. - To clear the blood secretion	**Contraindications:** Contraindication: can damage the soft tissue during insertion **Side Effects:** None	FSC India, & EC Certificate	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj	
231.	**Manufacturer:** Poly Medicure Ltd.	Poly VAC set Close wound suction unit	Poly VAC set Close wound suction unit	It is used for closed wound drainage or collection of fluid	**Contraindications:** Fistulas to organs or body cavities.	FSC-India	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj	

	Regd Office: 1st Floor, 12 Sant Nagar, East of Kailash, New Delhi-110 065 India. Industry: Plot No. 104-105 & 115-116, Sector-59, HSIIDC Industrial Area, Ballabhagra-121004, Faridbad (HARYANA) India. **Local Agent:** Aglow International, 114, 1st Floor, Aziz Co-operative Super Market, Shahbag, Dhaka.	**Class: B**		from surgical wound under negative pressure through collection tube and collection device (bellow) with an option to operate on or two catheters simultaneously.	**Side Effects:** None			
232.	**Manufacturer:** Sree Umiya Surgical Pvt. Ltd. Plot o. 4704, Phase. IV, G.I.D.C, VATA, Ahmedabad. India. **Local Agent:** Hard Won Power Industries Ltd. House No. 39, Road-12, Shekhertek, Adabor. Dhaka.	Mediplus Vaccu Suck Set **Class: C**	Vaccue Suck Set made with N.T. PVC Tubing PVC, Part, Vaccu Part, PVC Chamber & Catheter	It is used for Yankaur Suction	**Contraindication:** None **Side Effect**: None	FSC- India EC Certificate	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
233.	**Manufacturer:** Avent S. de R.L de C.V. Carretera Internacional Salida Norte No. 1053 Magdlena, Sonora Mexico. **Local Agent:** Medi Sensor Technology Incorporated Head Office. House 500A/1 (2nd Floor), Road, 8 Dhanmondi R/A, Dhaka.	Halyard Closed Suction System for Adults **Class: B**	Halyard Closed Suction System for Adults	It is used to Reduces the risk for contamination from outside pathogen, to protect from ventilator associated pneumonia (VAP).	**Contraindication:** None **Side Effect**: None	FSC Mexico	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
234.	Manufacturer: Huaian Wanjia Medical Device Co. Ltd. China.	Catgut **Class: B**	Sterilised Surgical Suture	Used to hold body tissues together after an injury or surgery. Application	**Contraindication:** None **Side Effect**: None	FSC-China	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

	Local Agent: Shanto Enterprise, 51, Islampur Road, Dhaka.			generally involves using a needle with an attached length of thread.				
235.	**Manufacturer:** Poly Medicure Ltd. Regd Office: 1st Floor, 12 Sant Nagar, East of Kailash, New Delhi-110 065 India. Industry: Plot No. 104-105 & 115-116, Sector-59, HSIIDC Industrial Area, Ballabhagra-121004, Faridbad (HARYANA) India. **Local Agent:** Aglow International, 114, 1st Floor, Aziz Co-operative Super Market, Shahbag, Dhaka.	Polymed 3 way stopcock with Extension tube **Class: B**	3 way stopcock with Extension tube (as an accessory of I.V. Canula/Cathter/ Perfusion set)	Infusion of multiple I. V fluids, parenteral nutrition and administration of other drugs.	**Contraindications:** It is contraindicated in case of highly viscous fluid. Not to be used in patients with known hypersensitivity to any of the materials used. **Side Effects:** None	FSC India, & EC Certificate	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
236.	**Manufacturer:** Natasha Lin, Fortune Medical Instrument Corporation. 6FI, NO: 23, Sec. 2, JhongJheng E. Road, Danshuei Dist. New Taipei City. Taiwan. **Local Agent:** MicroMed, 218/C Dr. Kudrat-E-Khuda Road (Elephant Road), Dhaka.	Fortune Silicone Stomach (Gastric) Tube **Class: B**	Silicon Stomach Tube	It is designed for nutrition supplement to stomach and may be recommended for various purposes: for patients who cannot take food or swallow, take enough food by month to keep nutrition, congenital defects of month, esophagus, or stomach.	**Contraindication**: None **Side effect**: None	FSC-Taiwan	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
237	**Manufacturer:** Natasha Lin, Fortune Medical Instrument Corporation. 6FI, NO: 23, Sec. 2, JhongJheng E. Road, Danshuei Dist. New Taipei City. Taiwan. **Local Agent:** MicroMed, 218/C	Fortune C. W. V Drain system **Class: B**	Drainage Tube	It is designed to collect fluid from different parts of body.	**Contraindication**: None **Side effect**: None	FSC Taiwan	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

	Dr. Kudrat-E-Khuda Road (Elephant Road), Dhaka.							
238.	**Manufacturer:** Sree Umiya Surgical Pvt. Ltd. Plot o. 4704, Phase. IV, G.I.D.C, VATA, Ahmedabad. India. **Local Agent:** Hard Won Power Industries Ltd. House No. 39, Road-12, Shekhertek, Adabor. Dhaka.	Mediplus Nasagestric Tube **Class: C**	Nasagestric Tube made with X-Ray Opaque Line N.T. PVC Tubing, Stopper & S.S. Boll	It is used for gastro intestinal feeding and aspiration.	High risk of aspiration, Gastric stasis, Gastro-oesophageal reflux, Upper gastrointestinal stricture	FSC India, EC Certificate	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
239.	Manufacturer: Welford Manufacturng (M) Sdn. Bhd 12, Jalan Angkasawan U1/39, Section U1, Hicom Glenmarie Industrial P ark, Selangor Darul Ehsan, Malaysia **Local Agent:** Ajanta Trading Corporation. 137 Lake Circus, Kalabagan, Dhaka.	Endotracheal Tube **Class: B**	Endotracheal Tube (Cuff/Un-cuffed)	Deliver oxygen in higher concentrations than found in air, or to administer other gases such as helium, nitric oxide, nitrous oxide, xenon, or certain volatile anesthetic agents such as desflurane, isoflurane, orsevoflurane, route for administration of certain medications such as salbutamol, airway management in the settings of general anesthesia, critical care, mechanical ventilation, and emergency medicine	**Contraindication:** None **Side Effect**: None	Malaysia	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
240.	**Manufacturer:** Shandong chengwu Medical Products Factory China **Local Agent** Tropical Eurasia internaqtional Ltd. 149/A New Airport Road, Farmgat, Dhaka	Disposable Blood Specimen collection tube (Heaprin Sodium) **Class: B**	Disposable Blood Specimen collection tube (Heaprin Sodium)	It is used for Blood Specimen collection.	**Contraindication:** None **Side effect:** None	FSC-China & EC Certificate	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
241.	**Manufacturer:** Shandong chengwu Medical Products Factory China	Disposable Blood Specimen collection tube (Heaprin Lithium)	Disposable Blood Specimen collection tube (Heaprin Lithium)	It is used for Blood Specimen collection.	**Contraindication:** None **Side effect:** None	FSC-China & EC Certificate	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

	Local Agent Tropical Eurasia internaqtional Ltd. 149/A New Airport Road, Farmgat, Dhaka	**Class: B**						
242.	**Manufacturer:** Shandong chengwu Medical Products Factory China **Local Agent** Tropical Eurasia internaqtional Ltd. 149/A New Airport Road, Farmgat, Dhaka	Disposable Blood Specimen collection tube (ESR 1:4) **Class: B**	Disposable Blood Specimen collection tube (ESR 1:4)	It is used for Blood Specimen collection.	**Contraindication:** None **Side effect:** None	FSC-China & EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |
243.	**Manufacturer:** Shandong chengwu Medical Products Factory China **Local Agent** Tropical Eurasia internaqtional Ltd. 149/A New Airport Road, Farmgat, Dhaka	Disposable Blood Specimen collection tube (Fluride) **Class: B**	Disposable Blood Specimen collection tube (Fluride)	It is used for Blood Specimen collection.	**Contraindication:** None **Side effect:** None	FSC-China & EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |
244.	Croma-Pharma Gesellschaft m.b.H Austria **Local agent :** Janata Traders TCB Bhabon , 1 kawran Bazar,Dhaka	**VISICROM 2%** **2.5ml** **Class: C**	Hydroxy Propyl Methyl Cellulose	The product is an ophthalmic viscosurgical device (OVD) intended for intraocular application during ophthalmic anterior segment surgeries. It is highly dispersive with especially good adherence to tissues.	**Contraindications:** The product must not be administered to patients who are known to be hypersensitive to HPMC or other components of the solution. **Side Effect:** Hypotension: transient episodes of hypotension have been observed after phacoemulsification and intraocular lens implantation. Anaphylaxis: a case of severe Anaphylaxis, probably caused by an	FSC-Austria	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |

					equivalent HPMC product, was reported.			
245.	Croma-Pharma Gesellschaft m.b.H Austria **Local agent :** Janata Traders TCB Bhabon , 1 kawran Bazar,Dhaka	**EYEFILL S.C. 2% 0.9ml** **Class: C**	Sodium Hyaluronate	The product is a medical device intended for intraocular use during ophthalmic anterior and posterior segment surgeries. It protects the intraocular tissue and prevents adhesions and synechia formation during the procedure.	**Contraindications:** The product must not be administered to patients who are known to be hypersensitive to HPMC or other components of the solution. Hypertension: transient postoperative elevations of intraocular pressure (IOP), most probably caused by the viscoelastic device, were reported. **Side Effect:** . Hypotension: transient episodes of hypotension have been observed after phacoemulsification and intraocular lens implantation. Anaphylaxis: a case of severe Anaphylaxis, probably caused by an equivalent HPMC product, was reported.	FSC-Austria	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj
246.	Croma-Pharma Gesellschaft m.b.H Austria **Local agent :** Janata Traders TCB Bhabon , 1 kawran Bazar,Dhaka	**NEOCROM cohesive 1.4%** **1ml** **Class: C**	Sodium Hyaluronate	It is intended for intraocular use during ophthalmic anterior and posterior segment surgery.It protects the intraocular tissue and prevents adhesions and synechia formation during the procedure.	**Contraindications:** The product must not be administered to patients who are known to be hypersensitive to sodium hyaluronate or other components of the solution. There are incompatibilities between sodium hyaluronate and quaternary ammonium compounds such as benzalkonium chloride solutions. The product should therefore not come into contact with operating instruments rinsed with these solutions or with ophthalmic preparations containing ammonium compounds as a preservative. No interactions with other substances have been reported yet. **Side Effect:** Postoperative elevations of intraocular pressure (IOP) have been reported with sodium hyaluronate viscoelastics. The IOP elevations are usually	FSC-Austria	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj

					transient, peaking at 4 to 7 hours postoperatively and returning to baseline within a few days. However, since their maximum may exceed 30 mmHg, the IOP should be carefully monitored. IOP-lowering therapy may be necessary, especially in patients with a compromised outflow facility. The IOP elevations may be caused by a reduction of aqueous outflow due to blockage of the trabecular meshwork.			
247.	Croma-Pharma Gesellschaft m.b.H Austria **Local agent :** Janata Traders TCB Bhabon , 1 kawran Bazar,Dhaka	**CORNEA PROTECT** **Class: C**	2.0ml viscoelastic solution	It is intended for intraocular application during ophthalmic anterior segment surgeries. It is highly dispersive with especially good adherence to tissues	**Contraindications:** The product must not be administered to patients who are known to be hypersensitive to HPMC or other components of the solution. Hypertension: transient postoperative elevations of intraocular pressure (IOP), most probably caused by the viscoelastic device, were Reported. **Side Effect :** . Hypotension: transient episodes of hypotension have been observed after phacoemulsification and intraocular lens implantation. Anaphylaxis: a case of severe Anaphylaxis, probably caused by an equivalent HPMC product, was reported.	FSC-Austria	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
248.	Croma-Pharma Gesellschaft m.b.H Austria **Local agent :** Janata Traders TCB Bhabon , 1 kawran Bazar,Dhaka	**MEGACROM 1.8% 1ml** **Class: C**	Sodium Hyaluronate	The product is a medical device intended for intraocular use during ophthalmic anterior and posterior segment surgery. It protects the intraocular tissue and prevents adhesions and synechia formation during the procedure.	**Contraindication:** The product must not be administered to patients who are known to be hypersensitive to sodium hyaluronate or other components of the solution. There are incompatibilities between sodium hyaluronate and quaternary ammonium compounds such as	FSC-Austria	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

					benzalkonium chloride solutions. The product should therefore not come into contact with operating instruments rinsed with these solutions or with ophthalmic preparations containing ammonium compounds as a preservative. No interactions with other substances have been reported yet. **Side Effect :** Postoperative elevations of intraocular pressure (IOP) have been reported with sodium hyaluronate viscoelastics. The IOP elevations are usually transient, peaking at 4 to 7_hours postoperatively and returning to baseline within a few days. However, since their maximum may exceed 30_mmHg, the IOP should be carefully monitored. IOP-lowering therapy may be necessary, especially in patients with a compromised outflow facility. The IOP elevations may be caused by a reduction of aqueous outflow due to blockage of the trabecular meshwork.			
249.	**Manufacturer:** **GE Healthcare** GE Medical Systems Information Technologies, Inc., USA **local Agent:** Wipro GE Health care Shanta Western Tower, Level-8, Tejgaon, Dhaka	CARESCAPE V100 Vital Signs Monitor	Vital Signs Monitor	It is intended to monitor a single adult, pediatric or neonatal patient's vital signs at the bedside or during intra-hospital transport. Vital signs parameters include non-invasive blood pressure (systolic, diastolic, and mean arterial pressure), pulse rate, and/or oxygen saturation (pulse oximetery) and/or temperature. The portable device is designed for use in numerous	**Contraindication:** None **Side Effects:** None	FSC-USA	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

				clinical settings in various hospital department such as emergency, radiology, recovery, medical/surgical, labor and delivery, endoscopy, cardiac step-down. The CARESCAPE V100 Vital Signs Monitor can also be used in satellite areas, physicians' office, or alternative care settings.				

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bs	c¯ZKvi‡Ki bvg	‡gwW‡Kj wWfvB‡mi bvg		†Rwbwi K bvg	wb‡`kbv/e¨envi	Contraindication & Side-effect	wWmwm-244 Zg mfvi wm×vš	FSC/CPP	‡UKwbK¨vj mve-Kwgwi 64 Zg mfvi wm×vš	mfvi wm×vš
250.	Devon Innovations Pvt. Ltd., India [Zain International Medical & Surgical Export Import Ltd.)	a)	Ureteral Indwelling double pigtail stent/set (DJ Stent) **Class: C**	Ureteral Indwelling double pigtail stent/set (DJ Stent) 2.5F to 10F : 8-30 cm	Used for temporary internal drainage from the Ureteropelvic junction to the bladder. Supplied sterile in peel open pouch. Intended for one time use.	Contraindication: None Side Effects: None	DbZ mvZwU †`‡ki g‡a¨ †h †Kvb GKwU †`‡ki FSC/KvbvWv-Gi FSC/EC Certificate `wLj mv‡c‡ÿ cieZx mfvq we‡ePbv Kiv n‡e|	FSC-India, EC Certificate	Abyôvb Kiv †h‡Z cv‡i|	Abyôvb Kiv nj|
251.	Devon Innovations Pvt. Ltd., India [Zain International Medical & Surgical Export Import Ltd.)	b)	Ureteral Catheter **Class: B**	Ureteral Catheter 3F to 8F: 70 cm	Used for dilation of the Ureter prior to Ureteroscopy and /or stone manipulation. Supplied sterile in peel open packages. Used with 0.038" guide-wire	Contraindication: None Side Effects: None	**H**	FSC-India, EC Certificate	Abyôvb Kiv †h‡Z cv‡i|	Abyôvb Kiv nj|
252.	Devon Innovations Pvt. Ltd., India [Zain International Medical & Surgical Export Import Ltd.)	c)	Percutaneous Pigtail Suprapubic Catheter/Set **Class: B**	Percutaneous Pigtail Suprapubic Catheter/Set 6F to 16F : 22-30 cm	Used to provide bladder drainage by percutaneous placement of a pigtail catheter. Supplied sterile in peel open package. Intended for one time use	Contraindication: None Side Effects: None	**H**	FSC-India, EC Certificate	Abyôvb Kiv †h‡Z cv‡i|	Abyôvb Kiv nj|

bs	c¯ZKvi‡Ki bvg	‡gvW‡Kj wWfvB‡mi bvg		†RvbwiK bvg	wb‡`kbv/e¨envi	Contraindication & Side-effect	wWmwm-244 Zg mfvi wm×vš	FSC/CPP	‡UKwbK¨vj mve-KwgwUi 64 Zg mfvi wm×vš	mfvi wm×vš
253.	Devon Innovations Pvt. Ltd., India [Zain International Medical & Surgical Export Import Ltd.)	d)	Percutaneous Pigtail Nephrostomy Catheter/Set **Class: B**	Percutaneous Pigtail Nephrostomy Catheter/Set 5F to 16F : 30 cm	Sterile tube inserted into the bladder to drain urine	Contraindication: None Side Effects: None	**H**	FSC-India, EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj	
254.	Devon Innovations Pvt. Ltd., India [Zain International Medical & Surgical Export Import Ltd.)	e)	Dilator Sets **Class: B**	Dilator Sets 5F to 16F : 22-70 cm	Used for dilation of the Ureter prior to Ureteroscopy and or stone manipulation.	Contraindication: None Side Effects: None	**H**	FSC-India, EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj	
255.	Devon Innovations Pvt. Ltd., India [Zain International Medical & Surgical Export Import Ltd.)	f)	Percutaneous Malecot naphrostomy Catheter **Class: B**	Percutaneous Malecot naphrostomy Catheter 8F to 30F: 30 cm	Used for temporary or permanent drainage of urine from the kidney by percutaneous placement	Contraindication: None Side Effects: None	**H**	FSC-India, EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj	
256.	Devon Innovations Pvt. Ltd., India [Zain International Medical & Surgical Export Import Ltd.)	g)	Guidewires **Class: B**	Guidewires 0.018 to 0.038 150 cm	Urological Product	Contraindication: None Side Effects: None	**H**	FSC-India, EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj	
257.	Sahajanand Medical Technologies Pvt Ltd., India [Zain International Medical & Surgical Export Import Ltd.)	a)	SUPRAFLEX (Drug Eluting Coronary Stent) **Class: D**	Sirolimus eluting polimer free Coronary Stent system	Use for coronary blockage.	Contraindication: None Side Effects: None	**H**	FSC-India, EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj	

Annex-G

Proposed Product for Locally Manufacture (Herbal):

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
1.	Incepta Herbal & Nutricare Ltd.	Cranberry Extract 300mg Soft Capsule Each soft gelatin capsule contains 300 mg Cranberry Extract USP	Cranberry is a small, evergreen shrub grown throughout North America. Cranberry has a long history of use among native American Indian tribes, primarily for treating urinary conditions. Juice and extracts from the fruit (berry) are used as medicine. Cranberry is most commonly used for prevention and treatment of urinary tract infections (UTIs). Cranberry JUICE seems to help prevent UTIs, but so far it doesn't seem to be effective in treating UTIs. Cranberry is also used for neurogenic bladder (a bladder disease), as well as to deodorize urine in people with urinary incontinence (difficulty controlling urination). Some people use cranberry to increase urine flow, kill germs, speed skin healing, and reduce fever.	**Contraindications:** Cranberry has a record of safety, although specific long-term safety data are lacking. No significant herb-drug interactions have been reported. A single study found that cranberry may increase the absorption of vitamin B_{12} in patients who also are taking proton pump inhibitors and that it may allow the kidneys to metabolize weakly alkaline drugs (such as antidepressants and opioids) more rapidly, thus reducing their effectiveness. A small study found a significant rise in urinary oxalate levels, prompting a caution that regular use of cranberry may increase the risk of kidney stone formation in patients with a history of oxalate calculi. **Side Effects:** Cranberry is LIKELY SAFE for most people. Cranberry juice and cranberry extracts have been used safely in research. Cranberry juice is LIKELY SAFE for children. But drinking too much cranberry juice can cause some side effects such as mild stomach upset and diarrhea. Drinking more than 1 liter per day for a long period of time might increase the chance of getting kidney stones.	New	Mosby's Drug Consult (Page: ||| - 29)	Abygv`b Kiv ‡h‡Z cv‡i|	Abygv`b Kiv হল|

bs	c¯ZKvi‡Ki bvg	JI‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
2.	Radiant Nutraceuticals Limited (Herbal Division)	Cranberry *Vaccinium macrocarpon* 400 mg Capsule	Reduction in UTI recurrence , Kidney stones, Treatment of UTI	Contraindicated for patient with renal insufficiency or developing uric acid or calcium oxalate stones. Generally well tolerated. At high dosages diarrhea or mild gastrointestinal upset may occur	New	The ABC clinical guide to herbs; p-76 The Handbook of Clinically Tested Herbal Remedies; p-265	Abyjgv`b Kiv ‡h‡Z cv‡i	Abyjgv`b Kiv হল|
3.	M/s. Drug International Ltd. (Herbal Division)	Cranberry Vaccinium macrocarpon (Standardized extracts) 140mg Capsule	▪ Prevent urinary tract Infection ▪ Cranberry has also been investigated for numerous other medicinal uses, include prevention of H. polyori infection and dental plaque.	Do	New	1. PDR for Herbal medicines 4th edition Page n. 234-240	Abyjgv`b Kiv ‡h‡Z cv‡i	Abyjgv`b Kiv হল|
4.	Incepta Herbal & Nutricare Ltd.	Kava Extract (Piper methysticum) 100mg Capsule Each capsule contains 100 mg of Kava Kava (Piper methysticum)	◘ Nervous anxiety and stress ◘ Restlessness, tension, and agitation Kava Kava is used for nervous tension, stress, and agitation. Unproven uses: In folk medicine, the herb is used as a sleeping agent and sedative; for asthma, rheumatism, dyspeptic symptoms, chronic cystitis, syphilis, gonorrhea, and weight reduction. Homeopathic uses: Kava Kava is used for states of excitement and exhaustion. It is also used for gastritis and pain in the urethra.	**Contraindications:** Kava and kava-containing products are not recommended for use in children younger than 12 years of age, or in patients with renal disease, thrombocytopenia, or neutropenia. 1 Additionally, patients with depression, liver disease, and Parkinson disease should avoid using kava. **Side Effects:** Kava has also been reported to cause mild and reversible gastrointestinal complaints, CNS complaints, including dizziness and headache, and various hypersensitivity/dermatological reactions. Pupil dilatation, near vision abnormalities, and eye movement coordination abnormalities have been reported. Kava dermopathy (a reversible darkening or yellowing of the skin with whitish scaling and flaking) has been reported with long-term use of higher doses. Regular administration of Kava for longer than 3 months is not recommended.	New	PDR for Herbal Medicines (Page: 494)	Abyjgv`b Kiv ‡h‡Z cv‡i	Abyjgv`b Kiv হল|

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
5.	Radiant Nutraceuticals Limited (Herbal Division)	Kava *Piper methysticum* 100 mg Capsule	Anxiety disorder, Sleep disorder, Stress and restlessness, Muscle relaxant	Not for use by persons under 18 years of age. Persons who have liver disease or depression should take with precaution Adverse effects with recommended doses of kava are relatively rare. Large doses may cause a scaly, yellowing skin condition, which resolves when use is discontinued.	New	The ABC clinical guide to herbs; p-259	Abyjgv`b Kiv ‡h‡Z cv‡i|	Abyjgv`b Kiv হল|
6.	Incepta Herbal & Nutricare Ltd.	Granules: Ispaghula Husk 0.110gm+Ispaghula Seed 2.6gm+Senna Fruits 0.62gm Sachet Granules: Ispaghula 54.2%, tinnevelly senna fruits 12.4%.(5 g granules contains 2.6 g Ispaghula Seed BP, 0.11 g Ispaghula Husk BP and 0.34-0.66 g Tinnevelly Senna Pods BP)	• Constipation • Constipation in bed-ridden patients • Constipation in pregnancy • Pain-free bowl evacuation in cases of hemorrhoids	**Contraindications:** • Blockage of the gut (intestinal obstruction) • Children under 5 years of age • Inflammation of the bowel and back passage (ulcerative colitis) • Inflammation of the mucous membrane of both small and large intestines (enterocolitis) **Side Effects:** Medicines and their possible side effects can affect individual people in different ways. The following are some of the side effects that are known to be associated with this medicine. Just because a side effect is stated here, it does not mean that all people using this medicine will experience that or any side effect. • Excess gas in the stomach and intestines (flatulence or wind). • Abdominal pain or cramps. • Abdominal swelling and risk of obstruction of the intestines if the medicine is not taken with sufficient fluid (see the instructions on how to take above). • Excessive use can cause diarrhea and low levels of potassium in the blood (see warning above). You should not exceed the recommended dose.	New	BNF 68 (Page: 72)	Abyjgv`b Kiv ‡h‡Z cv‡i|	Abyjgv`b Kiv হল|

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
7.	Popular Pharmaceuticals Ltd.	Ispaghula Husk (as effervescent Powder) Plantago Ovata 100gm/100gm	Gently relives Constipation Reduces High Blood Cholesterol Level Relieves Irritable Bowel Syndrome Restores Bowel Regularity Prevents Haemorrhoids Stabilizes Glucose Level	Do	New	British Herbal pharmacopeia Martindale extra Pharmacopeia Pharmacognosy Treas & Evabs' WHO Monographs on selected medicinal plants	Abygv`b Kiv ‡h‡Z cv‡i	Abygv`b Kiv হল|
8.	Radiant Nutraceuticals Limited (Herbal Division)	Psyllium Husk Powder + Standardize Senna Leaf Dry Extract Ispaghula Powder 3.5g USP (Plantago Ovata) + Cinsidering 5.5% Sennoside –B content 2.54g BP(Cassia angustifolia)	Psyllium Husk & Senna leaf Powder is the best laxative in Constipation associated with a)irritable bowel syndrome b)Fissures c)Haemorrhoids d)Pregnancy and e)Pre and post-operative conditions	Pathological narrowing in the gastrointestinal tract, intestinal obstructions, difficult-to-control diabetes mellitus, acutely inflamed intestinal diseases, e.g., Crohn's disease, ulcerative colitis, appendicitis, abdominal pain of unknown origin. Children under 12 years of age. Allergic reactions may occur. Chronic use/abuse: loss of electrolytes, especially loss of potassium, albuminuria and hematuria, pigment implantation into the intestinal mucosa, which is harmless and usually is reversed upon discontinuation of the drug.	New	The complete German commission E monograph	Abygv`b Kiv ‡h‡Z cv‡i	Abygv`b Kiv হল|
9.	Radiant Nutraceuticals Limited (Herbal Division)	Psyllium Husk powder Ispaghula Husk USP 3.5gm (Plantago Ovata)	Psyllium Husk is used in Constipation associated with a)irritable bowel syndrome b)Fissures c)Haemorrhoids d)Pregnancy and e)Pre and post-operative conditions	Contraindicated in patients who have pathological narrowing in the gastrointestinal tract, obstruction or threatening obstruction of the bowel (ileus), or difficulties in regulating diabetes mellitus. No significant side effect has been observed	New	USP, BP	Abygv`b Kiv ‡h‡Z cv‡i	Abygv`b Kiv হল|

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
10.	Radiant Nutraceuticals Limited (Herbal Division)	Psyllium Husk Dispersible powder Ispaghula Husk USP 3.5gm (Plantago Ovata)	Psyllium Husk is used in Constipation associated with a)irritable bowel syndrome b)Fissures c)Haemorrhoids d)Pregnancy and e)Pre and post-operative conditions	Contraindicated in patients who have pathological narrowing in the gastrointestinal tract, obstruction or threatening obstruction of the bowel (ileus), or difficulties in regulating diabetes mellitus. No significant side effect has been observed	New	USP, BP & Ph. Eur	Abyjgv`b Kiv ‡h‡Z cv‡i	Abyjgv`b Kiv হল|
11.	M/s. Square Herbal & Nutraceuticals Ltd.	Ispaghula Husk + Lactitol Monohydrate Ispaghula Husk 3.5gm (Plantago Ovata) + Lactitol Monohydrate 10gm/ Sachet	▪ Treatment of chronic idiopathic constipation and hepatic encephalopathy.	Contraindications: Appendicitis, Unexplained abdominal pain, Galactosemia Side effect: Abdominal distension, Cramp and flatulence	New	British Herbal Pharmacopoea p-113 PDR for Herbal medicines 4th edition P-669 ▪ WHO monographs Vol.1 p.202 USP 32-NF27 P. 1263	Kw¤‡bkb wnmv‡e bv _vKvq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i	Kw¤‡bkb wnmv‡e bv _vKvq Av‡e`b bv gÄyi Kiv nj
12.	M/s. Square Herbal & Nutraceuticals Ltd.	Echinacea+ Andrographis+ Bical Skullcap Echinacea Purpurea 350mg+Andrographis Paniculata 25mg+ Scutellaria Baicalensis 125mg Capsule	▪ Cold, Cough, Fever, Sore throat, Tendency to infection, Upper respiratory tract infection, Viral fever and infection, Dengue fever, Immune enhancement & Prevent antibiotic resistance	Contraindications: Multiple Sclerosis , Rheumatoid arthritis and lupus. Side effect: Very rarely allergic reactions, nausea, hypotension.	New	PDR for Herbal P-267 PDR for Herbal medicines 4th edition P-739. ▪ Mosby's Handbook of Herbs & Natural Supplements 3rd edition. P-399 & 38 ▪ WHO monographs Vol.2 p.2002, p-12	Abyjgv`b Kiv ‡h‡Z cv‡i	Abyjgv`b Kiv হল|

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
13.	Incepta Herbal & Nutricare Ltd.	Echinaceae 200mg + Goldenseal 100mg Capsule Each capsule contains 200mg of *Echinacea purpurea* aerial parts, *Echinacea angustifolia* root BP & 100mg goldenseal root and rhizome BP	Echinacea is one of the most commonly prescribed botanicals for the immune system. Echinacea (E. angustifolia and purpurea) is considered a "blood purifier," primarily due to its anti-inflammatory and immune stimulating properties. Researchers have found the extract of Echinacea to have anti-viral activity against the influenza pathogen. It increases the number of phagocytes, which are the type of white blood cells that "engulf" microbial invaders such as the flu virus.	**Contraindications:** Do not use if pregnant or breastfeeding. If you take immunosuppressive medication, do not take Echinacea. Echinacea is contraindicated if you are taking antibiotics, corticosteroids, or chemotherapy. None well defined for Goldenseal. **Side Effects:** Goldenseal is <u>POSSIBLY SAFE</u> for most adults when taken by mouth as a single dose. There is not enough reliable information to know if goldenseal is safe for long-term use. Echinacea is considered to be safe when taken in recommended dosages. When taken by mouth, it can cause nausea, sore throat and numbness and tingling in the mouth. If you are allergic to other plants in the daisy family, you may not be able to take echinacea. If you have multiple sclerosis, diabetes, HIV/AIDS or liver disease, you should only take it under your doctor's supervision. If you take immunosuppressive medication, do not take echinacea.	New	PDR for Herbal Medicines (Page: 395) (Page: 267)	Abygv`b Kiv ‡h‡Z cv‡i|	Abygv`b Kiv হল|

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
14.	Incepta Herbal & Nutricare Ltd.	Probiotic Powder (Lactobacillus reteri NCIMB 30242 80mg Capsule Each capsule contains 80 mg powder containing at least 2.0 x 10^9 (2 billion) colony-forming units (CFU) of live bacteria up to the date of expiration	• *Lactobacillus reuteri* is a natural supplement that contains the probiotic lactobacillus reuteri NCIMB 30242, the first and only probiotic that safely supports healthy cholesterol in adults with moderately elevated cholesterol but within the normal range. • *Lactobacillus reuteri* the natural bacteria may help to promote overall digestive health and help maintain a healthy digestive system. • *Lactobacillus reuteri* is used to improve digestion and restore normal flora. • *Lactobacillus reuteri* have been used to treat bowel problems (such as diarrhea, irritable bowel), eczema, vaginal yeast infections, lactose intolerance, and urinary tract infections.	**Contraindications:** *Lactobacillus* preparations are contraindicated in persons with a hypersensitivity to lactose or milk. *Lactobacillus* preparations have not been studied during pregnancy, in breastfeeding women, or in women trying to become pregnant. If you are pregnant, breastfeeding or trying to become pregnant, you should check with your healthcare provider before using it to ensure it is right for you. **Side Effects:** A significant number of human clinical trials have reported no adverse effects associated with consumption of Lactobacillus reuteri (L. reuteri). In the present study, the clinical safety and toxicology of oral ingestion of supplement capsules containing L. reuteri NCIMB 30242 was investigated. A randomized group of 131 subjects received a dose of 2.9 x 109 CFU L. reuteri NCIMB 30242 capsules (n = 67) or placebo capsules (n = 64) twice daily for 9 weeks. Clinical chemistry and hematological parameters of safety were analyzed. The frequency, duration and intensity of adverse events (AE)s and clinical significance of safety parameters were recorded for both groups. No clinically significant differences between the probiotic capsule and placebo capsule treated groups were detected in either the blood clinical chemistry or hematology results. The frequency and intensity of AEs was similar in the two groups. These results demonstrate that administration of a twice daily dose of 2.9 x 109 CFU was safe and well tolerated in the population evaluated over 9 weeks.	New	PDR for Herbal Medicines, 4th edition (Page: 996-1001)	Abyjgv`b Kiv ‡h‡Z cv‡i|	Abyjgv`b Kiv হল|

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
15.	Incepta Herbal & Nutricare Ltd.	Lactobacillus acidophilus 0.5 billion CFU + Bifidobacterium bifidum 0.5 billion CFU (Total Probiotic Cultures 1 Billion) Each serving (Choc ball) contains Lactobacillus acidophilus 500 million cfu and Bifidobacterium lactis 500 million cfu.	Probiotic Choc Balls are the delicious way to help maintain digestive health. Probiotics are the good bacteria essential for digestive health and wellbeing. Often it is the imbalance of good to bad bacteria in our gastrointestinal system that can cause digestive upset. An imbalance in the intestinal bacteria can result in tummy trouble including bloating, gas and upset stomach.	**Contraindications:** *Lactobacillus* preparations are contraindicated in persons with a hypersensitivity to lactose or milk. No contraindications are listed for bifidobacteria, since most species are considered nonpathogenic and nontoxigenic. **Side Effects:** Lactobacillus is LIKELY SAFE for most people, including babies and children. Side effects are usually mild and most often include intestinal gas or bloating.	New	PDR for Herbal Medicines, 4th edition (Page: 996-1001)	Abygv`b Kiv ‡h‡Z cv‡i|	Abygv`b Kiv হল|
16.	Incepta Herbal & Nutricare Ltd.	Lactobacillus rhamnosus GR-1 & Lactobacillus reuteri RC-14 blend 51.1111mg eq. to 2.3 billion Each capsule contains 2.3 billion organisms of *Lactobacillus rhamnosus*, GR-1 and *Lactobacillus reuteri*, RC-14	• Vaginal irritation • Vaginal discomfort • Reduction of colonization of bad bacteria and yeast in the vagina. • Protection of vagina against imbalances • Re-establishment and maintenance of good bacteria in the vagina.	**Contraindications:** Known hypersensitivity to any component of the formulation. **Side Effects:** Probiotics are LIKELY SAFE for most people**.**	New	PDR for Herbal Medicines, 4th edition (Page: 996-1001)	Abygv`b Kiv ‡h‡Z cv‡i|	Abygv`b Kiv হল|

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
17.	Radiant Nutraceuticals Limited (Herbal Division)	Fermented Soy (Probiotic) *Soy fermented by Lactobacillus delbrueckii* 500 mg Tablet	Acute and chronic gastritis, Rise of gastric juice after alcohol or tobacco abuse	European Food Safety Authority (EFSA) Journal 2011;9(5):2136 Technical data sheet: Fermented Soy, Lallemand Health Ingredients	New	European Food Safety Authority (EFSA) Journal 2011;9(5):2136 Technical data sheet: Fermented Soy, Lallemand Health Ingredients	Abyjgv`b Kiv ‡h‡Z cv‡i|	Abyjgv`b Kiv hvq|
18.	M/s. Square Herbal & Nutraceuticals Ltd.	Probiotic Fructo-Oligosaccharides 50mg Lactobacillus acidophilus 1 billion (13.33mg) per 1 gram powder in sachet	▪ Helps to maintain a favorable balance of intestinal microflora in babies and children. ▪ Lactose intolerance and ▪ bowel discomfort	Contraindications: Hypersensitivity Side effect: Flatulence and constipation	New	PDR for Herbal Medicines; 4th edition. p. 996 – 1001 ▪ Dietary supplements; Second edition by Pamela Mason. p. 180 – 185.	Abyjgv`b Kiv ‡h‡Z cv‡i|	Abyjgv`b Kiv hvq|
19.	M/s. Square Herbal & Nutraceuticals Ltd.	Probiotic blend, Papaya extract, & soy fibre Fructo-Oligosaccharides 100.00 mg Soy fiber & papaya extract 140.00 mg Probiotics blend 300mg Lactobacillus acidophilus 2.25 billion, Bifidobacterium bifidum 450 million Lactobacillus bulgaricus 75 million Lactobacillus casei 2.25 billion Lactobacillus lactis 2.1 billion Bifidobacterium infantis 1.5 billion Bifidobacterium longum 75 million Streptococcus thermophilus 300 million per 1gram powder in Capsule	▪ Helps to maintain the intestinal microflora ▪ Aids the digestion ▪ Antibiotic related diarrhea & illness ▪ Lactose intolerance ▪ IBS etc	Contraindications: Hypersensitivity Side effect: Flatulence and constipation	New	▪ PDR for Herbal Medicines; 4th edition. p. 996 – 1001 ▪ Dietary supplements; Second edition by Pamela Mason. p. 180 – 185.	Abyjgv`b Kiv ‡h‡Z cv‡i|	Abyjgv`b Kiv hvq|

bs	c¯ZKvi‡Ki bvg	JI‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
20.	M/s. Square Herbal & Nutraceuticals Ltd.	Probiotic for children Fructo-Oligosaccharides (FOS) 50.00 mg Bifidobacterium bifidum 13.33 mg (Equivalent to 1 billion) per 1gram powder in sachet	▪ Rotavirus diarrhea, ▪ Antibiotic-associated diarrhea, ▪ Clostridium difficile diarrhea ▪ Boost gastrointestinal immunity	Contraindications: Hypersensitivity Side effect: Flatulence and constipation	New	▪ Dietary supplements; Second edition by Pamela Mason. p. 180 – 185. Martindale 35. Page # 2113.	Abygv`b Kiv ‡h‡Z cv‡i	Abygv`b Kiv হল|
21.	M/s. Square Herbal & Nutraceuticals Ltd.	Probiotic Preparation Combination of Lactobacillus acidophilus- 2 billion, Lactobacillus bulgaricus- 1 billion, Bifidobacterium bifidum- 1 billion & fructo-oligosaccharides- 100 mg /Sachet	▪ prevention and treatment of ▪ Rotavirus diarrhea, ▪ Antibiotic-associated diarrhea, ▪ Clostridium difficile diarrhea, ▪ Traveler's diarrhea. ▪ Lactobacillus and bifidobacterium significantly reduce the events of Irritable Bowel Syndrome (IBS).	Contraindications: Hypersensitivity Side effect: Flatulence and constipation	New	▪ PDR for Herbal Medicines; 4th edition. p. 996-1001	Abygv`b Kiv ‡h‡Z cv‡i	Abygv`b Kiv হল|
22.	Renata Limited (Herbal Division) Kashor, Hobirbari, Bhaluka, Mymensingh.	Probiotic 0.1 billion (*Lactobacillus reuteri)* *Lactobacillus reuteri* 0.1 billion viable bacteria per sachet take 1 sachet mixed with water or liquid.	Excessive crying and colic in infants.	None reported.		Dietary supplements (4^{th}Edition); Page:378	Abygv`b Kiv ‡h‡Z cv‡i	Abygv`b Kiv হল|

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
23.	Renata Limited (Herbal Division) Kashor, Hobirbari, Bhaluka, Mymensingh.	Probiotic 1 billion (*Lactobacillus reuteri)* *Lactobacillus reuteri* 1 billion viable bacteria per sachet. take 1 sachet mixed with water or liquid.	Excessive crying and colic in infants.	None reported.	New	Dietary supplements (4thEdition); Page:378	Abyjgv`b Kiv ‡h‡Z cv‡i|	Abyjgv`b Kiv হল|
24.	M/s. Total Herbal & Neutraceuticals	Lactobacillus acidophilus +Lactobacillus plantarum+ Lactobacillus reuteri+Lactobacillus casei +Lactobacillus rhamnosus+Bifidobacterium animalis (B.lactis)+Bifidobacterium longum susp.longum (B.longum)+ Bifidobacterium bifidum Lactobacillus acidophilus 2.8 Billion+Lactobacillus plantarum 4 Billion+ Lactobacillus reuteri 2 Billion +Lactobacillus casei 1.6 Billion + Lactobacillus rhamnosus 3 Billion +Bifidobacterium animalis (B.lactis) 4.4 Billion +Bifidobacterium longum susp.longum (B.longum)2 Billion+ Bifidobacterium bifidum 0.2 Billion	Anti – Constipation Reproductive Health	Consult a health care practitioner prior to use if you have fever, vomiting, bloody diarrhea or severe abdominal pain. Discontinue use and consult a health care practitioner if symptoms of digestive upset (e.g. diarrhea) occur, worsen, or persist beyond 3 days	New	a. PDR for Herbal Medicines; 3rd edition. - P.965 b. Irit Chermesh and Rami Eliakim; Probiotics and the gastrointestinal Tract: Where are we in 2005?; World j Gastroenter	Abyjgv`b Kiv ‡h‡Z cv‡i|	Abyjgv`b Kiv হল|

						lo 2006 February 14; 12(6): 853-857		

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
25.	M/s. Total Herbal & Neutraceuticals	Lactobacillus acidophilus Bifidobacterium lactis Lactobacillus rhamnosus Lactobacillus acidophilus 3 Billion+Bifidobacterium lactis 4 Billion + Lactobacillus rhamnosus 3 Billion	1. Gut halth 2. Anti- diarrhea 3. Immune Boost	Consult a health care practitioner prior to use if you have fever, vomiting, bloody diarrhea or severe abdominal pain. Discontinue use and consult a health care practitioner if symptoms of digestive upset (e.g. diarrhea) occur, worsen, or persist beyond 3 days	New	a. PDR for Herbal Medicines; 3rd edition. - P.965	Abyjgv`b Kiv ‡h‡Z cv‡i|	Abyjgv`b Kiv হল|
26.	M/s. Total Herbal & Neutraceuticals	*Lactobacillus acidophilus Lactobacillus rhamnosus Bifidobacterium animalis (B.lactis) Bifidobacterium bifidum* Lactobacillus acidophilus 2 Billion+ Lactobacillus rhamnosus 2 Billion+ Bifidobacterium animalis (B.lactis) 4 Billion+ Bifidobacterium	1) Grwoth boost 2) Immune Boost 3) Alimentation Improvement	Consult a health care practitioner prior to use if you have fever, vomiting, bloody diarrhea or severe abdominal pain. Discontinue use and consult a health care practitioner if symptoms of digestive upset (e.g. diarrhea) occur, worsen, or persist beyond 3 days	New	PDR for Herbal Medicines; 3rd edition. - P.965	Abyjgv`b Kiv ‡h‡Z cv‡i|	Abyjgv`b Kiv হল|

		bifidum 2 Billion						

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
27.	M/s. Kemiko Pharmaceuticals Ltd. (Herbal Division)	Extract of blackcurrant fruit, Lactobacillus acidophilus, Bifidobacterium lactis, Thiamine Nitrate, Riboflavin, Nicotinamide, Pantothenic, Pyridoxin, Cyanocobalamin, Ascorbic Acid, Anhydrous Caffeine , Zinc Sulfate. Ribes nigrum 200 mg, Lactobacillus acidophilus 2 billion, Bifidobacterium lactis 2 billion, Thiamine Nitrate 1.4 mg, Riboflavin 1.6 mg, Nicotinamide 18 mg, Pantothenic acid 6 mg, Pyridoxin 6 mg,Cyanocobalamin 1.0 mcg, Ascorbic Acid 500.0 mg, Anhydrous Caffeine 50.0 mg, Zinc Sulfate 20.661 mg **Powder (5 gm).**	Antibiotic-associated diarrhea, Travelers' diarrhea, Lactose intolerance,Vaginal candidiasis, Irritable bowel syndrome, strengthens Immune system.	<u>Side-effects:</u> No health hazards or side-effects are known in conjunction with the proper administration of designated therapeutic dosage.	New	For *Ribes nigrum* (Blackcurrant) -PDR for Herbal Medicine-4th edition-P/100-101Research at Massey University, New Zealand For Probiotics-PDR for Herbal Medicine-4th edition-P/996-1000	wfUwgb Gi c‡qvRbxq †idv‡iÝ bv _vKvq Av‡e`b bv gÄyÝKiv hvB‡Z cv‡i|	wfUwgb Gi c‡qvRbxq †idv‡iÝ bv _vKvq Av‡e`b bv gÄyÝKiv nj|

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
28.	M/s. Kemiko Pharmaceuticals Ltd. (Herbal Division)	Extract of blackcurrant fruit, Lactobacillus acidophilus, Bifidobacterium lactis, Thiamine Nitrate, Riboflavin, Nicotinamide, Pyridoxin, Ascorbic Acid, Zinc Sulfate. Ribes nigrum 75 mg, Lactobacillus acidophilus 2 billion, Bifidobacterium lactis 2 billion, Thiamine Nitrate 1.122 mg, Riboflavin 1.571 mg, Nicotinamide 22.222 mg, Pyridoxin 2.222 mg, Ascorbic Acid 144.737mg, Zinc Sulfate 27.548 mg **powder(3.5 gm).**	Antibiotic-associated diarrhea, Travelers' diarrhea, Lactose intolerance,Irritable bowel syndrome, strengthens Immune system.	<u>Side-effects:</u> No health hazards or side-effects are known in conjunction with the proper administration of designated therapeutic dosage.	New	For *Ribes nigrum* (Blackcurrant) - PDR for Herbal Medicine-4th edition-P/100-101Research at Massey University, New Zealand For Probiotics- PDR for Herbal Medicine-4th edition-P/996-1000	**wfUwgb Gi c‡qvRbxq †idv‡iÝ bv _vKvq Av‡e`b bv gÄýKiv hvB‡Z cv‡i|**	**wfUwgb Gi c‡qvRbxq †idv‡iÝ bv _vKvq Av‡e`b bv gÄýKiv nj|**

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix Kwgw U (Jla wbqšY Kwgw Ui †UKwbK¨vj mve Kwgw Ui mvfvi wm×vš	mfvi wm×š
29.	Radiant Nutraceuticals Limited (Herbal Division)	Selenium yeast 30mg + Vitamin D yeast 50mg + Activated yeast ß-glucans 270mg (Probiotic) Selenium yeast, Vitamin D yeast & Activated yeast ß-glucans 350mgTablet	Antioxidant, Natural defenses, Immunomodulation with supply of essential nutrients including vitamins, minerals, amino acids	Mosby's Drug Consult Page-99 The European Food Safety Authority Journal (2008) 766, 1-42 Technical data sheet: Yeast based formulation, Lallemand Health Ingredients	New	The EFSA Journal (2008) 766, 1-42 The EFSA Journal (2009) 1148, 1-6 The EFSA Journal (2011) 9(5):2137 Technical data sheet: Yeast based formulation, Lallemand Health Ingredients	Kwc‡bkb wnmv‡e †idv‡iÝ bv_vKvq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i	Kwc‡bkb wnmv‡e †idv‡iÝ bv_vKvq Av‡e`b bv gÄyi Kiv nj|
30.	Incepta Herbal & Nutricare Ltd.	Refined Evening Primrose Oil 1000mg Soft Capsule Each soft gelatin capsule contains 1000 mg of Refined Evening Primrose Oil BP (*Oenothera biennis L.*)	• Premenstrual syndrome symptoms (PMS) • Cyclical mastalgia • Lactation	**Contraindications:** Evening Primrose Oil has few adverse effects, causing occasional headache, nausea if taken on an empty stomach, and diarrhea only in high doses. It may exacerbate temporal lobe epilepsy and mania, so should be avoided in these cases.Paradoxically, high doses of linoleic acid and arachidonic acid may reduce T-cell function. **Side Effects:** Evening primrose oil was previously suspected to lower the seizure threshold in schizophrenic patients; however, this is now disputed.	Rose 500 (General), Eprim 500mg (Square)	Mosby's Drug Consult™ (Page: ||| - 44)	Abyjgv`b Kiv ‡h‡Z cv‡i|	Abyjgv`b Kiv হল|

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
31.	M/s. Drug International Ltd. (Herbal Division)	Evening Primrose Oil Evening Primrose Oil Soft Cap. 1000mg (Oenothero Bicnnis (Extract)	▪ Premen Strual Syndron ▪ Breast Pain	Previously it was not recommended for patients diagnosed with schizophrenia. However, a recently published analysis of clinical trials involving polyunsaturated fatty acids in the treatment of schizophrenia did not indicate a clear therapeutic or adverse effect of Evening Primrose Oil supplements on schizophrenic patients. Side effects are rare at recommended dosages.Overdose may cause loose stool and abdominal pain.	New	1) PDR for Herbal Medicines 4th edition, Page: 311-313 2. Bangladesh National unini Formulary Page: 682-683	Abygv`b Kiv ‡h‡Z cv‡i|	Abygv`b Kiv হল|
32.	Radiant Nutraceuticals Limited (Herbal Division)	Elderflower 36mg, Primrose 36mg, Common sorrel 36mg, European vervain 36mg & Gentian 12mg Extracts of *Sambucus nigra, Primula officinalis, Rumex acetosa, Verbena officinalis,Gentiana lutea* 156 mg Tablet	Acute and chronic inflammation of the paranasal sinuses and the upper respiratory tract	Hypersensitive (allergic) to the ingredients. Tablets should not be used by children younger than 12 years old. Safely used in millions of doses over 35 years. Reported adverse side effects include gastrointestinal (GI) disorders and hypersensitivity (allergy) reactions.	New	American Botanical Council (ABC), Scientific & Clinical Monograph	Abygv`b Kiv ‡h‡Z cv‡i|	Abygv`b Kiv হল|

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
33.	Incepta Herbal & Nutricare Ltd.	Grape Seed Extract 50mg Soft Capsule Each soft gelatin capsule contains 50 mg Standardized Grapeseed extract	• Treatment of varicose veins and chronic venous • insufficiency • Reduce swelling due to surgery or injury • Treat and prevent macular degeneration • To reduce the risk for cancer and heart disease • Treat diabetic retinopathy and neuropathy	**Contraindications:** Grape seed extract is contraindicated in patients with known hypersensitivity to grape seed. **Side Effects:** Grape seed extract is generally considered safe. Side effects may include headache, itchy scalp, dizziness, and nausea. If you have a bleeding disorder or high blood pressure, talk to your doctor before you start using grape seed extract.	New	PDR for Herbal Medicines (Page: 409)	Abyjgv`b Kiv ‡h‡Z cv‡i|	Abyjgv`b Kiv হল|
34.	Incepta Herbal & Nutricare Ltd.	Grapeseed Extract 100mg Softgel Capsule Each soft gelatin capsule contains 100 mg Standardized Grapeseed extract	• Treatment of varicose veins and chronic venous • insufficiency • Reduce swelling due to surgery or injury • Treat and prevent macular degeneration • To reduce the risk for cancer and heart disease • Treat diabetic retinopathy and neuropathy	**Contraindications:** Grape seed extract is contraindicated in patients with known hypersensitivity to grape seed. **Side Effects:** Grape seed extract is generally considered safe. Side effects may include headache, itchy scalp, dizziness, and nausea. If you have a bleeding disorder or high blood pressure, talk to your doctor before you start using grape seed	New	PDR for Herbal Medicines (Page: 409)	Abyjgv`b Kiv ‡h‡Z cv‡i|	Abyjgv`b Kiv হল|

				extract.				

bs	c¯ZKvi‡Ki bvg Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
35.	Incepta Herbal & Nutricare Ltd. Black Cohosh 100mg Capsule Each capsule contains Black Cohosh BP 100mg standardized extract	*Possibly Effective for:* Menopausal symptoms: Research shows that taking some black cohosh products can reduce some symptoms of menopause. But the benefits are only modest. Black cohosh might lessen the frequency of hot flashes. Most of this research is for a specific commercial black cohosh product, Remifemin. The benefits may not occur with all products that contain black cohosh. Research using black cohosh products other than Remifemin have not always shown benefits for menopausal symptoms. Some of these studies show that these other black cohosh products do not reduce hot flashes or menopausal symptoms any better than a sugar pill ("placebo"). Some women take black cohosh for hot flashes related to breast cancer treatment. Women with breast cancer should not use black cohosh without talking to their cancer specialist or other health provider. Some early research suggested that black cohosh might reduce hot flashes in breast cancer patients, but more recent and higher quality research shows that black cohosh does not reduce hot flashes in women with breast cancer. Also, there is some question as to whether black cohosh is safe for women with breast cancer. It is important for a woman with breast cancer to discuss any use of black cohosh with her health providerbefore using it.	**Contraindications:** Black cohosh is contraindicated during pregnancy because of its potential ability to stimulate uterine contraction. The safety of black cohosh in breastfeeding mothers and the degree of transmission of black cohosh in breast milk are unknown. Controversy remains regarding the safety of black cohosh in women with a personal history or strong family history of breast cancer. **Side Effects:** Black cohosh is POSSIBLY SAFE when used appropriately by adults. Black cohosh can cause some mild side effects such as stomach upset, cramping, headache, rash, a feeling of heaviness, vaginal spotting or bleeding, and weight gain. There is also some concern that black cohosh may be associated with liver damage. It is not known for sure if black cohosh actually causes liver damage. Researchers are studying this. Until more is known, people who take black cohosh should watch for symptoms of liver damage. Some symptoms that may suggest liver damage are yellowing of the skin and eyes (jaundice), unusual fatigue, or dark urine. If these symptoms develop, black cohosh should be stopped and a health provider should be contacted. People who take black cohosh should talk with their health provider about getting tests to make sure their liver is working well.	Menocare 40mg (Acme)	PDR for Herbal Medicines (Page: 99)	Abyjgv`b Kiv ‡h‡Z cv‡i|	Abyjgv`b Kiv হল|

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
36.	M/s. Kemiko Pharmaceuticals Ltd. (Herbal Division)	Black cohosh root, Blue cohosh root, Chasteberry fruit , Cramp bark, Curcumin extract, Dong quai root, Wild yam, Ginger rhizome extract, Peony root extract, Calcium D-glucarate, Taurine 50 mg, Choline citrate, Rutin, Magnesium citrate, Pyridoxal 5 phosphate, Ascorbic. Cimicifuga racemosa 40 mg, Caulophyllum thalictroides 40 mg, Vitex agnus-castus 40 mg, Viburnum opulus 40 mg, Curcuma longa 40 mg, Angelica sinensis 40 mg, Dioscorea villosa 40 mg, Zingiber officinale 25 mg, Paeonia lactiflora 25 mg, Calcium D-glucarate 33 mg, Taurine 50 mg, Choline citrate 50 mg, Rutin 100 mg, Magnesium citrate 65 mg, Pyridoxal 5 phosphate 8 mg, Ascorbic acid 40 mg **Capsule.**	Premenstrual syndrome, dysmenorrhoea, sleep disturbances, uterine cramps, enhances estrogen regulation, supports nervous system.	<u>Side-effects:</u> No serious side-effects observed**.** Not recommended for pregnant and lactating mother.	New	Capsule **BioFem PMS**, a product of Biogenesis Nutraceuticals Inc., USA. PDR for Herbal Medicine 4th Ed. Vol.1, P# 95-100	wfUwgb Gi c‡qvRbxq †idv‡iÝ bv_vKvq Av‡e`b bv gÄyKiv hvB‡Z cv‡i|	wfUwgb Gi c‡qvRbxq †idv‡iÝ bv_vKvq Av‡e`b bv gÄyKiv nj|

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
37.	Incepta Herbal & Nutricare Ltd.	Standardized Spirulina Extract 750mg Soft Capsule Each capsule contains 750 mg standardized Spirulina Extract	▪ Spirulina's most powerful health benefits lie in its ability to stimulate and maintain immune system activity. ▪ As a food additive, Spirulina maxima contain beta-carotene, tocopherols and phenolic acids, which are proven to exhibit antioxidant properties. The antioxidant activity of Spirulina can be attributed to phycocyanin, the blue pigment found in blue-green algae that contain phytochemicals with scavenging properties. ▪ Spirulina has ability to cause a significant change in vascular tone by increasing the synthesis and release of nitric oxide and by decreasing the synthesis and release of a vasoconstricting substance from the endothelial cells.	**Contraindications:** Spirulina is contraindicated in patients in whom the risk of heavy metal poisoning is not outweighed by benefit of use. **Side Effects:** Hepatic: increased serum alkaline phosphatase level. Metabolic: increased serum calcium level.	Pirulin® 450 mg & Pirulin® 500 capsule, Navit Tab 500mg (Square), Spirucap 500mg (Drug International),	Mosby's Drug Consult™ (Page: ||| - 106)	Abyôgv`b Kiv ‡h‡Z cv‡i	Abyôgv`b Kiv হল|
38.	Radiant Nutraceuticals Limited (Herbal Division)	Spirulina *Spirulina platensis* 500mgTablet	Antioxidant with proteins, vitamins and minerals. Helps to combat many health problems as diabetes, arthritis and anemia & hyperlipidemic	Spirulina is contraindicated in those who are hypersensitive to any component of Spirulina containing supplement. Adverse effects: not known.	New	Spirulina in Human Nutrition and Health Dietary	Abyôgv`b Kiv ‡h‡Z cv‡i	Abyôgv`b Kiv হল|

						Supplements, 3rd edition, p-298		

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
39.	M/s. Bexter Herbal & Nutraceuticals	Spirulina Spirulina plankton 250mg Tablet	Mal nutrition, Immune eficincy diabetic, High cholesterol, Allergic reaction, Skin disorder, hair loss, Decreased milk Supply in lactating mothers.	Spirulina is a potent super food in the family of blue-green algae that packs a real nutritional punch. There are, however, certain contraindications and precautions to know about before purchasing and taking spirulina.	New	1.PDR for herbal medicine 2nd edition 2. WHO Monographs on Selected medicinal plants	Abygv`b Kiv ‡h‡Z cv‡i	Abygv`b Kiv হল|
40.	M/s. Total Herbal & Neutraceuticals	Spirulina +Green Tea Tablet Spirulina Platensis 400mg+Camellia Sinensis 100mg	General weakness, Immune enhancer, Lipid Lowring, Rheumatoid arthritis		New	Mani UV, Desai S, Lyer U Studies on the log term evect of Spirulina supplemenation on serum lipid profile and glycated proteins in NIDDM patiens, J. nutraceut funct med Foods, 2000; 2, 25-32	Kw¤‡bkb wnmv‡e †idv‡iÝ bv_vKvq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i	Kw¤‡bkb wnmv‡e †idv‡iÝ bv_vKvq Av‡e`b bv gÄyi Kiv nj|
41.	Incepta Herbal & Nutricare Ltd.	St. John's Wort Extract 23gm/100ml Topical Oil Each ml contains: Fresh organic and wild St. John's Wort (*Hypericum perforatum*) herb extract BP 230 mg which is obtained from the extract of 920 mg of aerial parts and flowers of *Hypericum perforatum* (Extraction ratio 1:4)	• Used as an anti-inflammatory, St Johns wort oil can help to ease the discomfort of hemorrhoids and skin irritations such as sun burn, although ironically in some rare cases St John's wort herb is known to cause photo dermatitis. •	**Contraindications:** This product should not be taken internally by those on sedatives or tranquilizers. May cause photosensitivity and skin irritations with internal and external use. **Side Effects:** St. John's wort is LIKELY SAFE for most people when taken by mouth short-term. It can cause some side effects such as trouble sleeping, vivid dreams, restlessness, anxiety, irritability, stomach upset, fatigue, dry mouth, dizziness,	New	PDR for Herbal Medicines (Page: 811)	Abygv`b Kiv ‡h‡Z cv‡i	Abygv`b Kiv হল|

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
			• This means that the skin can become sensitive to sunlight and have an adverse reaction such as flaking skin or inflamed areas. For sunburn, it is recommended to keep the oil refrigerated and apply directly to the skin. • It is also used in some countries to help heal the skin on people suffering from first degree burns, as it is known to speed up the healing process.	headache, skin rash, diarrhea, and tingling. Take St John's wort in the morning or lower the dose if it seems to be causing sleep problems. St. John's wort seems to be safe when used in children under 12 years of age for up to 6 weeks.However, St. John's wort is POSSIBLY UNSAFE when taken by mouth in large doses. It might cause severe reactions to sun exposure. Wear sun block outside, especially if you are light-skinned.				
42.	M/s. Kemiko Pharmaceuticals Ltd. (Herbal Division)	St. John's Wort flower extract, Rhodiola root extract, Lithium orotate, L-theanine, 5-HTP, GABA, N-acetyl-L-tyrosine, DMAE bitrate, Methylcobalamin, Folate, Pyridoxal 5 phosphate. Hypericum perforatum 75 mg, Rhodiola rosea 50 mg, Lithium 2.5 mg, L-theanine 25 mg, 5-HTP 25 mg, GABA 50 mg, N-acetyl-L-tyrosine 100 mg, DMAE bitrate 100 mg,Methylcobalamin 100 mcg, Folate 200 mcg,	Enhances work performance, decrease physical and mental fatigue, increase stamina and support depressed moods.	Side-effects: No serious side-effects observed.	New	Capsule Eskaloft, a product of Biogenesis Nutraceuticals Inc., USA PDR for Herbal Medicine 4th Ed., Vol. 2, P# 797-811	wfUwgb Gi c‡qvRbxq †id‡i Ý bv_vKvq Av‡e`b bv gÄy Kiv hvB‡Z cv‡i	wfUwgb Gi c‡qvRbxq †id‡i Ý bv_vKvq Av‡e`b bv gÄy Kiv nj	

		Pyridoxal 5 phosphate 12.5 mg						

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
43.	Incepta Herbal & Nutricare Ltd.	Black (Cumin) Seed oil 500mg Softgel Capsule Each soft gelatin capsule contains Black (Cumin) Seed Oil 500 mg	Improve Immune System Black seed oil capsule stimulates our immune system & thus prepares our body to fight against invading organisms. Reduce Blood Sugar Level By lowering blood sugar level, it helps diabetic patient's keep the control on their diabetic condition. Reduce blood pressure The active ingredients in black seed oil have been found to be effective in reducing high blood pressure. Allergies Black Seed Oil acts as an antihistamine which helps to reduce the common symptoms of allergies (watery eyes, sneezing). Simply take half a teaspoon of oil twice a day when allergy symptoms begin to appear. Flu and Fever Black Seed Oil has been found to help reduce fever, by inducing perspiration which helps the body cool and stimulate the release of toxins. Take one teaspoon of Black Seed Oil once a day when flu/fever systems occur. Also be sure to drink plenty of fluids throughout the day.	**Contraindications:** Not known. **Side Effects:** There are no known side effects of Black Seed Oil if it is taken moderately. However, it is not recommended for use during pregnancy.	New	Opsonin markets it as consumer product.	c‡qvRbxq †idv‡iÝ bv _vKvq Av‡e`b bv gÄyÿ Kiv †h‡Z cv‡i|	c‡qvRbxq †idv‡iÝ bv _vKvq Av‡e`b bv gÄyÿ Kiv nj|

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
44.	Radiant Nutraceuticals Limited (Herbal Division)	Black seed oil *Nigella sativa* 500 mg Capsule	Respiratory and digestive problems, parasites, inflammation. It also used for a variety of health conditions including, colds, infections, headaches, toothaches and digestive aid after large meals	Contraindicated during pregnancy. Safety in young children has not been established. Patients with liver or kidney disease are advised not to use without medical supervision. In general, black seed extract is not associated with serious side effects.	New	Gale encyclopedia of alternative medicine, 3rd edition; p-271 Journal of Herbal Medicine and Toxicology 4 (2) 1-8 (2010) ISSN : 0973-4643	c‡qvRbxq †idv‡iÝ bv _vKvq Av‡e`b bv gÄÿ Kiv †h‡Z cv‡i|	c‡qvRbxq †idv‡iÝ bv _vKvq Av‡e`b bv gÄÿ Kiv nj|
45.	Incepta Herbal & Nutricare Ltd.	Stevia (Leaf) Sachet 30mg Each sachet contains 30mg of Stevia (leaf) extract powder INN.	Stevia sachets are all-natural with no calories or carbs. When mixed in your favorite hot liquid, Stevia powder begins dissolving instantly, deliciously supplementing your coffee or tea. Stevia powder is extracted from the leaves of the stevia plant, and it has been described as the sweetest supplement on earth. What's truly extraordinary about stevia is that even though it is so sweet, it has no calories and it's actually beneficial to health!	**Contraindications:** Anyone with an allergy or sensitivity to stevia should avoid this product. **Side Effects:** A 2009 review summarized the basic research in which steviosides and related compounds are being tested for possible antidisease actions, with no effect yet demonstrated in humans. A 2011 review found that the use of stevia sweeteners as replacements for sugar might benefit diabetic patients because it is a non-caloric additive.	New	PDR for Herbal Medicines, 4th edition (Page: 789)	Abÿgv`b Kiv †h‡Z cv‡i|	Abÿgv`b Kiv হল|

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
46.	Incepta Herbal & Nutricare Ltd.	Stevia (Leaf) Tablet 30mg Each tablet contains 30mg of Stevia (leaf) extract INN.	Stevia Tabs are all-natural with no calories or carbs. When dropped in your favorite hot liquid, SteviaTabs begin dissolving instantly, deliciously supplementing your coffee or tea. Stevia is extracted from the leaves of the stevia plant, and it has been described as the sweetest supplement on earth. What's truly extraordinary about stevia is that even though it is so sweet, it has no calories and it's actually beneficial to health!	**Contraindications:** Anyone with an allergy or sensitivity to stevia should avoid this product. **Side Effects:** A 2009 review summarized the basic research in which steviosides and related compounds are being tested for possible antidisease actions, with no effect yet demonstrated in humans. A 2011 review found that the use of stevia sweeteners as replacements for sugar might benefit diabetic patients because it is a non-caloric additive.	New	PDR for Herbal Medicines, 4th edition (Page: 789)	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv হল|
47.	M/s. Square Herbal & Nutraceuticals Ltd.	Stevia Stevia standardized leaf extract 62.5 mg powder in sachet	▪ As a sweetening agent ▪ Hypertension ▪ Inflammation ▪ Weight loss ▪ Diabetes	Contraindications: hypotension and hypoglycemia	New	▪ PDR for Herbal Medicines, 4th Edition, Page: 789	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv হল|
48.	Radiant Nutraceuticals Limited (Herbal Division)	Valerian tablet 600mg *Valeriana officinalis* 600mg Tablet	For the symptomatic relief of insomnia, restlessness, and spasms due to nervous tension	Contraindicated for children under 12 years without medical supervision. Valerian is considered generally safe. Adverse effects may include headache and stomach upset, but these effects are rare.	New	The ABC clinical guide to herbs; p-351 American Herbal Pharmacopoeia and Therapeutic Compendium;	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv হল|

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix Kwgw U (Jla wbqšY Kwgw Ui †UKwbK¨vj mve Kwgw Ui mvfvi wm×vš	mfvi wm×š
						Analytical, Quality Control,and Therapeutic Monograph; *Valeriana officinalis*		
49.	Radiant Nutraceuticals Limited (Herbal Division)	valerian Valeriana Officinalis (Standardized Root extract) 450mg Tablets	Use for the symptomatic relief of insomnia, restlessness and spasms due to nervous tension.	Contraindicated for children under 12 years without medical supervision. Valerian is considered generally safe. Adverse effects may include headache and stomach upset, but these effects are rare.	New	The ABC clinical guide to herbs P-351	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv হল|
50.	Renata Limited (Herbal Division) Kashor, Hobirbari, Bhaluka, Mymensingh.	Valerian root capsule Valerian officinalis 500 mg per capsule Dosage for sleep-500mg once 1-2 hours before bedtime	Anxiety, insomnia, sleep disorders, restlessness based on nervous disorders.	WHO contradicts use of valerian for children under the age of 12 years, without medical supervision. However, German authorities allow clinical use as long as valepotriate and baldrinal-free preparations are used. The Herbal PDR be associated with headaches, restless states, sleeplessness, mydriasis and vague cardiac disturbance with long-time use, but no data on this.	New	The ABC Clinical Guide to Herbs; page 354	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv হল|
51.	Radiant Nutraceuticals Limited (Herbal Division)	Valerian 500mg + Hop 65mg Fixed combinations of *Valeriana officinalis* L., radix (valerian root) and *Humulus lupulus* L., flos (hop strobiles) 565mg Tablet	Sleeping disorders, mental stress	Contraindicated to any patient sensitive to the API or any other excipients of the dosage form. Safety during pregnancy has not been established clinically. There is no known side effect.	New	The ABC clinical guide to herbs; p-351 The complete German commission E monograph	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv হল|

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
52.	M/s. Drug International Ltd. (Herbal Division)	Valerian Root Extract + Chamomile Herbs Valeriana officinalis 40mg+Chamaemelium nobile 20mg Capsule	▪ Treatment of insomnia		New	1. AFP (American Family Physician, M fam physician 2003 Apr 15,67(8), 1755-1758	c‡qvRbxq †idv‡iÝ bvB wea‡q Av‡e`b bv gÄy Kiv †h‡Z cv‡i	c‡qvRbxq †idv‡iÝ bv _vKvq Av‡e`b bv gÄy Kiv nj
53.	M/s. Drug International Ltd. (Herbal Division)	Valerian Extract (Root) + Chamomile flower Extract+ Lemon Balm Extract (leaf) + Hops Extract (Stroble) + Passion Flower Extract (Aerial) Valeriana officinalis 500mg+Chamaemelium nobile 50mg+ Humulus Lupulus 120mg+ Melissa officinalis 300mg + Passiflora incarnate 50mg Tablet	▪ Treatment of insomnia		New	1. AFP (American Family Physician, M fam physician 2003 Apr 15,67(8), 1755-1758	c‡qvRbxq †idv‡iÝ bvB wea‡q Av‡e`b bv gÄy Kiv †h‡Z cv‡i	c‡qvRbxq †idv‡iÝ bv _vKvq Av‡e`b bv gÄy Kiv nj
54.	M/s. Square Herbal & Nutraceuticals Ltd.	Valerian root extract +Hops flower extract+ Passion Flower extract Valeriana Officinalis 250mg+Humulus lupulus 50mg+ Passiflorea herba 50mg Tablet	▪ Anxiety, insomnia, sleep disorders, restlessness based on nervous disorders	Contraindications: Medical Supervision needed for below 12 years old. Side effect: Little morning sleepiness	New	▪ The complete German commission E Monographs p 305 ▪ The ABC clinical Guide to herbs 1st	Kw¤‡bkb wnmv‡e †idv‡iÝ bvB wea‡q Av‡e`b bv gÄy Kiv †h‡Z cv‡i	Kw¤‡bkb wnmv‡e †idv‡iÝ bvB wea‡q Av‡e`b bv gÄy Kiv nj

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix Kwgw U (Jla wbqšY Kwgw Ui †UKwbK¨vj mve Kwgw Ui mvfvi wm×vš	mfvi wm×š
						▪ Edition p.351-364 ▪ PDR for Herbal medicines 4th edition P-448,634-635, 872-876 ▪ WHO monographs Vol.1 p.267-276		
55.	Radiant Nutraceuticals Limited (Herbal Division)	Aloe Vera *Aloe vera* 1000 mg Capsule	Occasional constipation	Aloe should not be used in patients with intestinal obstruction or stenosis, atony, severe dehydration. PREGNANCY & LACTATION: No teratogenic or fetotoxic effects were seen in rats after oral treatment with aloe extract (up to 1000 mg/kg). Abdominal spasms and pain may occur.	available in other strength	WHO monographs on selected medicinal plants; p-33 PDR for herbal medicines; p-16	Abyjgv`b Kiv †h‡Z cv‡i|	Abyjgv`b Kiv হল|
56.	M/s. Bexter Herbal & Nutraceuticals	Aloe Vera Aloe barbadensis 400mg Capsule	Aloe Bera helps in proper functioning of the digestive system. The adaptogenic properties of the plant are beneficial for proper digestive function.	Ingestion is contraindicated in pregnant and breast-feeding women, children younger than 12 years of age, patients with inflammatory bowel disease, and elderly patients with suspected intestinal obstruction.	New	1. The complete German commission E monographs 2. WHO Monographs on commission E Monographs 3. PDR for herbal medicine 2nd edition	Abyjgv`b Kiv †h‡Z cv‡i|	Abyjgv`b Kiv হল|

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
57.	M/s. Drug International Ltd. (Herbal Division)	Aloe Vera Extract Aloe Vera Extract (Aloe Barbadensis) 100mg Soft Gelatine Capsule	▪ Colic, Skin disease, amenorrhoea, worm infestation and infection, Fungal diseases, digestion Problem		New	1. Physician desk reference for Herbal medicines 4th Edition Page: 19-26 2. WHO monographs Vol.1 p. 33-40, 3. Europenan Meicines Agency 4. USA Ogarnacioeua	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv হল।
58.	M/s. Square Herbal & Nutraceuticals Ltd.	Aloe + Fennel Aloe (*Aloe ferox*) 250 mg and Fennel (*Foeniculi fructus*) 280 mg Capsule	▪ Constipation ▪ Dyspepsia ▪ Flatulence	Contraindications: Intestinal obstruction, Crohn's disease, ulcerative colitis, appendicitis. Side effect: very rarely Gastrointestinal cramp	New	PDR for Herbal Medicine 4th Edition; Pg: 19, 317 ▪ German Commission E 1st Edition; Pg: 80, 129 ▪ British Pharmacopoeia 2009; Pg: 6817	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv হল।
59.	Radiant Nutraceuticals Limited (Herbal Division)	Feverfew *Tanacetum parthenium* 50 mg/ Capsule	Migraine prophylaxis, Nausea and vomiting associated with migraine	Feverfew is contraindicated for persons who are allergic to feverfew Not recommended for children under two years of age. Not for use in pregnancy and lactation. Allergic contact dermatitis can result from handling fresh feverfew. Mouth ulceration and swelling of the tongue, lips, and oral mucosa have also been reported.	New	The ABC clinical guide to herbs ; p-135	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv হল।

bs	c¯ZKvi‡Ki bvg Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
60.	M/s. Square Herbal & Nutraceuticals Ltd.	Feverfew Feverfew (*Tanacetum parthenium)* 100 mg Capsule	▪ Migraine headaches prophylaxis ▪ Nausea and vomiting associated with migraine	Contraindications: Hypersensitivity Side effect: Abdomenal blotting, indigestion, heartburn and digestive upset.	New	PDR for Herbal Medicine 4th Edition; Pg: 321 The ABC clinical guide to herbs 1st Edition; Pg: 135 ▪ British Pharmacopoeia 2009; Pg: 6982	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv হল|
61.	Radiant Nutraceuticals Limited (Herbal Division)	Flaxseed oil *Linum usitatissimum* 1000 mg Capsule	Hyperlipidemia, Atherosclerosis, Risk of breast cancer and metastasis, Constipation	Contraindicated in cases of ileus of any origin. PREGNANCY & LACTATION: No known restrictions. Essential fatty acid (EFA) supplementation during pregnancy and nursing is beneficial for fetal and infant brain development and visual function. No significant side effect has been observed	New	The ABC clinical guide to herbs ; p-143	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv হল|
62.	Radiant Nutraceuticals Limited (Herbal Division)	Ginger *Zingiber officinale* 500mg Capsule	Motion sickness, Chemotherapy-induced nausea, Morning sickness, Nausea	Patients with gallstones should consult a healthcare provider before using ginger. PREGNANCY & LACTATION: traditionally been used to prevent morning sickness during the first trimester. Caution is advised against using excessive dosages of dried ginger during pregnancy. No significant side effect has been	New	The ABC clinical guide to herbs; p-171	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv হল|

				observed				

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
63.	M/s. Total Herbal & Neutraceuticals	Ginger 500mg Tablet *Zingiber officinale* 500mg	**a)** Colic, Flatulent dyspepsia, and Specifically for flatulent intestinal Colic vomiting, Particularly motion (travel) sickness, as a digestive aid Blood sugar and cholesterol management, b) Chemotherapy induced nausea, C) carminative		New	*British Herbal Pharmacopoeia, 1996*. Exeter: British Herbal Medicine Association, 1996, Page-87 *c. United States Pharmacopeia 29 and National Formulary 24 and Supplements*. Rockville, Maryland, US, United States Pharmacopeial Convention, 2006. *d.* American Herbal Pharmacopoeia – Page- *e.* Herbal Medicine , Third Edition, Page no 293-298 WHO Monograph on Selected Medicinal Plant, Volume -1, Page-277-287	Abyjgv`b Kiv †h‡Z cv‡i|	Abyjgv`b Kiv হল|
64.	Radiant Nutraceuticals Limited (Herbal Division)	Turmeric Powdered Extract (Contains NLT 20% of total Curcuminoids) *Curcuma longa* 500mg Capsule	Indigestion including sensation of fullness, flatulence and slow digestion	Hypersensitivity to the active substance(s). PREGNANCY AND LACTATION: Safety during pregnancy and lactation has not been established. Mild symptoms of dry mouth, flatulence and gastric irritation may occur.	New	European Medicines Agency (EMA), Community herbal monograph on *curcuma longa* PDR for herbal medicines; p-775	Abyjgv`b Kiv †h‡Z cv‡i|	Abyjgv`b Kiv হল|

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
65.	Radiant Nutraceuticals Limited (Herbal Division)	Curcuminoids *Curcuma longa* 500mgTablet	Osteoarthritis, dyspepsia	Contraindicated in patients having hypersensitivity to Curcuminoids or any other excipients of formulation. This product usually has very few side effects.	New	WHO monographs on selected medicinal plants United States Pharmacopeia	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv হল|
66.	Acme Laboratories (Herbal Division)	Turmeric 500mg Capsule Turmeric Powder (Curcuma Longa Linn 500mg	Dyspeptic Complaints, Loss of appetite	Contraindication: No contraindications have been determined. Side effects: Clinical trials report few adverse reactions. Rare cases of contact dermatitis and anaphylaxis have been reported. An increased risk of kidney stones is possible in susceptible individuals.	New	PDR for herbal Medicines 2nd Edition Page 775-777 The Complete German Commission E Monograph Page 222-223	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv হল|
67.	Acme Laboratories (Herbal Division)	Turmaric Plus Sandal Oil Turmeric Extract (Curcuma Longa Linn 1.60gm+Sandal Wood Oil (Santalum Album) 0.05gm 0.05gm/10gm	This Cream Prevents and cures skin infections, Inflammations, Blemishes Wounds and others skin disorder, It soothes boils, Pimples, acne and burns. It removes rashes, Cures dermatitis, allergic eruption, nappy rashes and eczema.	Contraindication: Caution should be taken in-patient with cardiovascular diseases or diabetes. Hypertension resulting from ginseng abuse syndrome is associated with prolonged high dose ginseng with concomitant use of caffeine. General adverse effect includes insomnia, epistaxis, headache, nervousness and vomiting. Side Effects: No known (at therapeutic doses), But massive over dosages can bring about ginseng abuse syndrome, which is characterized by hypertension, insomnia, hypertonia and edema.	available at cosmetics	-	c‡qvRbxq †idv‡iÝ bvB wea©vq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i	c‡qvRbxq †idv‡iÝ bvB wea©vq Av‡e`b bv gÄyi Kiv nj|

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
68.	Radiant Nutraceuticals Limited (Herbal Division)	Devil's claw *Harpagophytum procumbens* *210*mgTablet	Dyspeptic complaints, Loss of appetite, Rheumatism including arthritis or joint diseases and regenerating cartilage	Contraindicated for patients with stomach or duodenal ulcers There is no known side effect in case of therapeutic dosages.	New	PDR for herbal medicines; p-247 Technical date sheet on extract of *Harpagophytum procumbens*	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv হল|
69.	Radiant Nutraceuticals Limited (Herbal Division)	Soy Isoflavones *Glycine max* 200mgTablet	Hypercholesterolemia, Menopause related symptoms, Cardiovascular disorders, Hot flashes and Osteoporosis	No known Contra-indication There is no evidence of side effect from soy protein. Rare side effect including soy allergies	New	Herbal Medicine: Expanded Commission E Monographs Soy Isoflavones, Journal of Alternative Medicine Review Monographs Technical data sheet on soy extract	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv হল|
70.	M/s. Total Herbal & Neutraceuticals	Isoflavones Capsule Isoflavones (Soy Isoflaqvones) 150mg	CVD Cardiovascular Disease, Cancer, Osteoporosis, Menopausal Symptoms	Do	New	ABC clinical Guide to Herbs Page 153-170,WHO herbal Medicine Monograph, Vo. 2 Herbal medicine A Guide for Health care Professionals Page-129-133	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv হল|

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix Kwgw U (Jla wbqšY Kwgw Ui †UKwbK¨vj mve Kwgw Ui mvfvi wm×vš	mfvi wm×š
71.	Radiant Nutraceuticals Limited (Herbal Division)	Chlorella *Chlorella regularis* 500mgTablet	Antioxidant & detoxifier ,helps to prevent diabetes, asthma & indigestion, regulates blood pressure and PH	Chlorella is contraindicated in patients having hypersensitivity to Chlorella or any other excipients of the tablet Side effects are rare and may cause occasional gastrointestinal side effects. These side effects include nausea and diarrhea.	New	Dietary Supplements, 3rd edition, Pharmaceutical Press, UK	c‡qvRbxq †idv‡iÝ bvB weavq Av‡e`b bv gÄy Kiv †h‡Z cv‡i|	c‡qvRbxq †idv‡iÝ bvB weavq Av‡e`b bv gÄy Kiv nj|
72.	Radiant Nutraceuticals Limited (Herbal Division)	Fufang Danshen Diwan or Danshen Pill Extracts of *Radix Salviae Miltiorrhizae, Radix Notoginseng & Borneolum syntheticum/* 250mg capsule	For preventing and treating coronary atherosclerosis, angina pectoris & hyperlipidemia	Contraindicated to any patient sensitive to the API or any other excipients of the dosage form. Danshen appears to be a generally well tolerated agent. May be increased risk of bleeding when used with anticoagulants or agents that inhibit platelet aggregation and adhesion.	New	1. Chinese Pharmacopoeia Commission, *Pharmacopoeia of The People's Republic of China*, Edition 2005, Volume 1.page 416-417 2. Herbal Experts Working Group of the Pan-European Federation of TCM Societies (PEFOTS), High quality traditional Chinese herbal medicinal products intended to be registered as a traditional herbal medicinal product in the European Union, Edition 2004, *Danshenform.* page 1- 254 3. Product data sheet : Danshen Pill	Abyjgv`b Kiv †h‡Z cv‡i|	Abyjgv`b Kiv হল|

bs	c¯ZKvi‡Ki bvg	JI‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
73.	Radiant Nutraceuticals Limited (Herbal Division)	Huoxiang Zhengqi Shui or Summer Comfort Pill Extracts of *Rhizoma Atractylodis,Pericarpium citri reticulatae, Cortex Magnoliae officinalis (Processed with ginger), Radix Angelicae Dahuricae, Poria, Pericarpium Arecae, Rhizoma Pinelliae, Radix Glycyrrhizae, Oleum Pogostemonis* (Herba Patchouli oil) *Oleum Folii Perillae* (Folium Perillae oil)/ 2.6gm Sachet	Used for improving fever, cold sore (calenture), headache, abdominal distention, vomiting and gastrointestinal discomfort	Huoxiang Zhengqi Shui Pill is contraindicated to any patient sensitive to the API or any other excipients of the dosage form. No significant side effects were observed.	New	1. Chinese Pharmacopoeia Commission, *Pharmacopoeia of The People's Republic of China*, Edition 2005, Volume 1. page 474-475 2. Product data sheet : Summer Comfort Pill	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv hvB
74.	Radiant Nutraceuticals Limited (Herbal Division)	CerebralCare Granules Extract of *Radix Angelicae Sinensis, Rhizoma Chuanxiong, Radix Paeoniae Alba, Radix Rehmanniae Preparata, Ramulus Uncariae Cum Uncis, Caulis Spatholobi, Spica Prunellae, Semen Cassiae, Concha Margaritifera, Rhizoma Corydalis, Herba Asari/* 4gm Sachet	Improve blood circulation & enhance energy	Cerebral Care granules is contraindicated to child, pregnant woman or lactating women & patients with hepatic and renal insufficiency Possible side-effects are Nausea after medication occasionally, but with no impact on continuous medication, and it can disappear of itself	New	1. *An Illustrated Chinese Materia Medica.* Page 84,376,454,548,648, 604,520,166,222,112 2. Product data sheet : CerebralCare Granules	c‡qvRbxq †idv‡iÝ bvB wea‡q Av‡e`b bv gÄy Kiv †h‡Z cv‡i	c‡qvRbxq †idv‡iÝ bvB wea‡q Av‡e`b bv gÄy Kiv nj

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix Kwgwu (Jla wbqšY Kwgwui †UKwbK¨vj mve Kwgwui mvfvi wm×vš	mfvi wm×š
75.	Radiant Nutraceuticals Limited (Herbal Division)	Pearlium Pearlium (Soluble pearl extract) 325mg capsule	Calcium, Magnesium and Vitamin D supplement for normal development and maintenance of healthy bones and teeth. Helps to prevent calcium deficiency. Adequate Calcium , Vitamin D as part of a healthy diet, along with physical activity helps prevent bone loss/Osteoporosis in pre- and postmenopausal women.	Contraindicated to any patient sensitive to the API or any other excipients of the dosage form. No significant side effects were observed.	New	1.Calcium, Magnesium, Vitamin D, Zinc Monograph, Natural Health Products Directorate, Canada 2.The Role of nutrients in bone health, Critical Reviews in Food Science and Nutrition , 2006, 46;page 621-628	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv হল|
76.	Radiant Nutraceuticals Limited (Herbal Division)	Hops Hops Cone Standardized Extract Capsule 85mg (Humulas Lupulus)	For the treatment of symptomatic relief of menopausal symptom like hot flashes, insomnia, restlessness and spasms due to nervous tension.	Hops should not be taken by individuals suffering from depressive illness. During pregnancy and lactation should be avoided. No health hazards or side effects are known in conjunction with the proper administration of designated therapeutic dosages. Drowsiness or sedation may occur.	New	1) PDR for Herbal Medicine, 4th Edition; Page-400 2) The Journal of American Botanical Council - 2010	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv হল|

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
77.	Radiant Nutraceuticals Limited (Herbal Division)	Fraxinus Fraxinus Standardized Extract Capsule 500mg (Fraxinus excelsior)	For the treatment of symptomatic relief of arthritis, gout, bladder complaints, as well as laxative and diuretic.	Contraindicated to any patient sensitive to the API or any other excipients of the dosage form. No significant side effects were observed.	New	1) PDR for Herbal Medicine, 4th Edition; Page-50 2) The Journal of American Botanical Council -2010	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv hvB
78.	Radiant Nutraceuticals Limited (Herbal Division)	Green Coffee Beans Green Coffee Beans Standardized Extract Capsule 400mg (Coffea Arabica)	For management of Diarrhea, inflammation of the mouth and pharynx, used to treat hypotonia and as a constituent of analgesics.	Pregnant women should avoid caffeine, under no circumstance exceeding a dosage of 300 mg per day. Side effects include hyperacidity, stomach irritation, and diarrhea and reduced appetite. But side effects are not recorded at therapeutic dosages.	New	PDR for Herbal Medicine, 4th Edition; Page-202 The Journal of American Botanical Council -2010	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv hvB
79.	M/s. Kemiko Pharmaceuticals Ltd. (Herbal Division)	Valerin root extract, Chamomile flower, Hops extract, Passion flower extract, Skullcap leaf, Wild lettuce leaf, California poppy extract, Jamaican dogwood bark, Jujube seed extract, Melatonin , Calcium amino acid chelate, Magnesium amino acid chelate, GABA, Inositol, L-theanine, 5-HTP.	Supports for normal sleep patterns. Acts as muscle relaxant.	<u>Side-effects:</u> No serious side-effects observed. Not recommended for pregnant and lactating mother.	New	Capsule **Sleep Factors**, a product of Biogenesis Nutraceuticals Inc., USA PDR for Herbal Medicine 4th Ed., Vol. 2, P# 872-875	c‡qvRbxq †idv‡iÝ bvB weavq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i	c‡qvRbxq †idv‡iÝ bvB weavq Av‡e`b bv gÄyi Kiv nj

bs	c¯ZKvi‡Ki bvg	JI‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
		Valeriana officinalis 60 mg , Matrecaria recutita 20 mg, Humulus lupulus 20 mg, Passiflora incarnata 20 mg, Scutellaria lateriflora 20 mg, Lactuca virosa 20 mg, Eschscholzia california 12.5 mg, Piscidia piscipula 12.5 mg, Ziziphus spinosa 12.5 mg, Melatonin 1.5 mg, Calcium amino acid chelate 20 mg, Magnesium amino acid chelate 20 mg, GABA 50 mg,						
80.	M/s. Kemiko Pharmaceuticals Ltd. (Herbal Division)	Saw palmetto berry extract, Nettle leaf extract, Pygeum bark extract 2.5% phytosterol, Pumpkin seed, Beta sitosterol (40% sterols), Lycopene (5%), Asparate. Serenosa serrulata	Benign Prostatic Hyperplasia, Encourages healthy testosterone-dihydrotestosterone (DHT) balance, Supports enhanced control of bladder function, Supplies targeted nutrition for bladder and prostate tissues.	<u>Side-effects:</u> No serious side-effects observed. Not recommended for pregnant and lactating mother.	New	Capsule **BioProstate** , a product of Biogenesis Nutraceuticals Inc., USA PDR for Herbal Medicine 4th Ed., Vol. 2, P# 725-729	c‡qvRbxq †idv‡iÝ bvB weavq Av‡e`b bv gÄy Kiv †h‡Z cv‡i	c‡qvRbxq †idv‡iÝ bvB weavq Av‡e`b bv gÄy Kiv nj	

		315 mg, Urtica diolica 75 mg, Pygeum africanum						
bs	**c¯ZKvi‡Ki bvg**	**JI‡ai bvg I †R‡bwiK bvg**	**wb‡`kbv**	**Contra-indication & Side effect**	**Status (New Molecule/ Existing)**	**Reference**	*nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš*	**mfvi wm×š**
		62.50 mg, Cucurbita pepo 40 mg, Beta sitosterol (40% sterols) 7.5 mg, Lycopene (5%) 100 mcg, Zinc 5 mg **Capsule.**						
81.	M/s. Square Herbal & Nutraceuticals Ltd.	Saw Palmetto standardized extract with Ginseng & Pygeum extract Saw Palmetto berries extract 160.00 mg +Panax Ginseng 3.30 mg+ Pygeum bark Extract 1.70 mg + Zinc Methionine 83.45mg + Copper Gluconate 14.28 mg Liquid Filled Hard gelatin capsule	▪ Benign Prostatic Hyperplasia	Contraindications: No known contraindication. Side effect: minor GI complaints.	New	The ABC Clinical Guide to Herbs, 1st edition; P. 311-319 British Herbal Compendium; Vol-2; P.345-352 PDR for Herbal Medicine (4th edition); P. 384-389, 679-681	*Kw¤‡bkb wnmv‡e c‡qvRbxq †idv‡iÝ bvB weavq Av‡e`b bv gÄy Kiv †h‡Z cv‡i |*	*Kw¤‡bkb wnmv‡e c‡qvRbxq †idv‡iÝ bvB weavq Av‡e`b bv gÄy Kiv nj |*

82.	M/s. Drug International Ltd. (Herbal Division	Milk thistle Sylimarin Marianum 140mg Capsule	▪ Treatment of Jaundice, Chronic inflammatory liver conditions i.e. Hepatitis, alcoholic liver damage and hepatic cirrhosis.	Contraindications: Chemotherapy, immunosuppressant, allergic Side effect: No known side effect.	New	1) PDR for Herbal medicines 4th edition 2) The complete german commission-E-Monograph Therapeutic Guid to Herbal Medicines 3) British Herbal Pharmacopoeia	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv* হল|

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
83.	M/s. Kemiko Pharmaceuticals Ltd. (Herbal Division)	milk thistle seed powder, black radish, Beet root, Turmeric, Green tea, Dandelion, Ox bile (Bovine) , L-taurine, Betaine (HCL), Inositol, DL-methionine, Liver extract 50 mg, Choline (Bitrate), Magnesium (Citrate), Cyanocobalamin, Calcium folinate, Pyridoxal 5-phosphate. Silybum marianum 17 mg, Raphanus sativus 8 mg, Beta vulgaris 8 mg, Curcuma longa 17 mg, Camellia sinensis 17 mg, Taraxacum officinale 17 mg, Ox bile (Bovine) 17 mg , L-taurine 17 mg, Betaine (HCL) 17 mg, Inositol 33 mg, DL-methionine 33 mg, Liver extract 50 mg, Choline (Bitrate) 67 mg, Magnesium (Citrate) 9 mg, Cyanocobalamin 3 mcg, Calcium folinate 36 mcg, Pyridoxal 5-phosphate 1.5 mg **Cap**	Supports bile flow, detoxifies liver, provides liver protection.		New	Capsule **BioLiv**, a product of Biogenesis Nutraceuticals Inc., USA PDR for Herbal Medicine 4th Ed., Vol. 2, P# 578-581	c‡qvRbxq †idv‡iÝ bvB wea‡q Av‡e`b bv gÄy Kiv †h‡Z cv‡i|	c‡qvRbxq †idv‡iÝ bvB wea‡q Av‡e`b bv gÄy Kiv nj|

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix Kwgw U (Jla wbqšY Kwgw Ui †UKwbK¨vj mve Kwgw Ui mvfvi wm×vš	mfvi wm×vš
84.	M/s. Square Herbal & Nutraceuticals Ltd.	Combination of milk thistle, Brahmi, Ashwagandha, Green tea and Turmeric Milk Thistle Extract USP 225 mg + Bacopa Extract USP 150mg + Ashwagandha Extract BP 150mg + Green Tea Extract USP 75mg + Turmeric Extract BP 75mg Tablet	▪ Coronary Heart Disease ▪ High Blood Pressure ▪ High Cholesterol ▪ Multiple Sclerosis ▪ Muscular Dystrophy ▪ Parkinson's disease ▪ Asthma/Emphysema ▪ Kidney Failure ▪ Rheumatoid Arthritis ▪ Lupus disease ▪ Skin Cancer ▪ Non-alcoholic Fatty Liver Disease	Contraindications: Chemotherapy, immunosuppressant, allergic Side effect: No known side effect.	New	▪ Milk Thistle (Silybum marianum) USP 36-NF31; Page:1538 Bacopa (Bacopa monniera) USP 36-NF31; Page: 1343 ▪ Ashwagandha (Withania somnifera) - BP 2014; Vol: IV; Page: 395 ▪ Green Tea (Camellia sinensis) USP 36-NF31; Page: 1500 ▪ Turmeric (Curcuma longa) - BP 2014; Vol: IV; Page: 382 Milk Thistle- The ABC Clinical Guide to Herbs, page-285 Green Tea- The ABC Clinical Guide to Herbs, page-335 Curcumin- WHO monographs on selected medicinal plants V-1, P-115	Kwc‡bkb wnmv‡e c‡qvRbxq †idv‡iÝ bv _vKvq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i|	Kwc‡bkb wnmv‡e c‡qvRbxq †idv‡iÝ bv _vKvq Av‡e`b bv gÄyi Kiv nj|

bs	c¯ZKvi‡Ki bvg	JI‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
85.	M/s. Kemiko Pharmaceuticals Ltd. (Herbal Division)	Colchicine, extract of chicken sternum (Collagen II) Colchicum autumnale equivalent to 0.5 mg Colchicine & extract of chicken sternum equivalent to 693 mg Collagen II **Tablet.**	Gout, Osteoarthritis, Osteoporosis, Rheumatoid arthritis, degenerative bone diseases .	No known Contra-indication & Side effect	New	For *Colchicum autumnale*- PDR for Herbal Medicine-4th edition-P/214-215The Complete German Commission E Monograph-P/86 For Collagen II –Research at Harvard University School of Medicine- www.autumnleavesnutrition.com Research at Anhui University in Hefei, China- http://collageena.com/info_study_results.html	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv হল|

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×vš
86.	M/s. Square Herbal & Nutraceuticals Ltd.	Rosehip+ Glucosamine+ Chondroitin Sulfate Rosa Canina 150mg+ Zea mays 250mg+ Shark Cartilage 200mg Tablet	▪ Osteoarthritis (OA) Rheumatoid arthritis (RA) Degenerative change in bone and Joint. Prophylaxis of Muscle and tendon injury.	Contraindications: Caution in Diabetes, G6PD deficiency, thalassemia, sickle cell anemia. Side effect: Diarrhea, constipation	New	PDR for Herbal medicines 4th edition P-258,955,967 Mosby's Handbook of Herbs & Natural Supplements 3rd edition. P-506	Kwª‡bkb wnmv‡e c‡qvRbxq †idv‡iÝ bv _vKvq Av‡e`b bv gÄy Kiv †h‡Z cv‡i|	Kwª‡bkb wnmv‡e c‡qvRbxq †idv‡iÝ bv _vKvq Av‡e`b bv gÄy Kiv nj|
87.	M/s. Square Herbal & Nutraceuticals Ltd.	Lutein+ Zeaxanthin Chlorella Sorokiniana 20mg+ Lycium Barbarum 5mg Capsule	▪ Cataracts ▪ Age related macular degeneration ▪ Rethinitis Pigmentosa	Contraindications: Wilson's disease Side effect: No side effect found	New	PDR for Herbal medicines 4th edition P-980-983	Abyjgv`b Kiv †h‡Z cv‡i|	Abyjgv`b Kiv হল|
88.	M/s. Square Herbal & Nutraceuticals Ltd.	Methi + Gurmar Fenugreek Standardized Extract 300 mg + Gymnema Standardized Extract 200 mg Capsule	▪ Type-2 diabetes mellitus ▪ Obesity	Contraindications: Caution in low blood suger. Side effect: No side effect	New	▪ The complete German commission E (GCE) p- 130 American botanical council, HC 060392-387, October 30, 2009	Abyjgv`b Kiv †h‡Z cv‡i|	Abyjgv`b Kiv হল|
89.	M/s. Square Herbal & Nutraceuticals Ltd.	Gripe water Mouri BP 4 mg + Ada BP 5 mg + Sodium Bicarbonate 15mg /5ml Syrup	▪ Dyspeptic complains ▪ Abdominal pain, windy colic in infant ▪ Halitosis ▪ Worm infestation ▪ Respiratory and urinary tract complications.	Contraindications: No known contraindication. Side effect: GI tract irritation, rear allergic reaction	Banned in 1982	British Herbal Compendium; Volume-2; Page-147-150 Complete German Commission E; Page – 128 BP 2010, Page# 3572	Banned Item weavq Av‡e`b bv gÄy Kiv †h‡Z cv‡i|	Banned Item weavq Av‡e`b bv gÄy Kiv nj|

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
90.	M/s. Square Herbal & Nutraceuticals Ltd.	Lutein + Zeaxanthin Lutein (*Chlorella sorokiniana*) 25 mg and Zeaxanthin (*Lycium barbarum*) 5 mg Capsule	▪ Cataracts ▪ Age related macular degeneration ▪ Retinitis pigmentosa	Contraindications: Wilson's disease Side effect: No side effect found	New	PDR for Herbal Medicine 4th Edition; Pg: 980-983 ▪ Judy et al., Lutein and Zeaxanthin and Their Potential Roles in Disease Prevention, Journal of the American College of Nutrition (2004), 23 (6): 567S–587S	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv হল|
91.	M/s. Square Herbal & Nutraceuticals Ltd.	Proanthocyanidins 100mg Capsule French Maritime Pine Bark 100 mg Capsule	▪ Improving vascular health ▪ Improving blood circulation and blood pressure ▪ Platelet function normalization ▪ Coronary artery disease ▪ Venous insufficiency ▪ Other potential indications are: thrombosis, diabetes, hypertension, asthma, attention deficit hyperactivity disorder (ADHD), endometriosis, dysmenorrheal and osteoarthritis.	Contraindications: No known contraindications. Side effect: GI discomfort , dizziness .	New	The ABC clinical guide to herbs 1st Edition; Pg: 369 ▪ American Botanical Council, Proprietary Botanical Ingredients, Scientific and clinical monograph for Pycnogenol.	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv হল|

bs	c¯ZKvi‡Ki bvg	JI‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
92.	M/s. Total Herbal & Neutraceuticals	Biotin Tablet Biotin 1000mcg	Hair loss., Diabetes:, Diabetic nerve pain: Brittle fingernails and toenails:	It is not known if this product interacts with any medicines. Before taking this product, talk with your health professional if you take any medications	New	1. USP – Dietary Supplement Compendium, USP32- NF27, Page -447-448 2. Dietary reference intakes: vitamin A, vitamin K, arsenic, boron, chromium, copper, iodine, iron, manganese, molybdenum, nickel, silicon, vanadium and zinc: J Am Diet Assoc.2001;101:294-300.	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv হল|
93.	M/s. Total Herbal & Neutraceuticals	Cinnamon+ Alpha Lopic Acid Capsule	Diabetes: Liver disease Diabetes mellitus, Glaucoma,	Contact with cinnamon bark or oil may cause an allergic reaction. Cinnamon oil is stated to be a	New	World Health Organization. *WHO Monographs on*	c‡qvRbxq †idv‡iÝ bvB wea‡q Av‡e`b bv gÄyi Kiv †h‡Z cv‡i	c‡qvRbxq †idv‡iÝ bvB wea‡q Av‡e`b

		Cinnamomum zeylanicum 400mg + Alpha Lopic Acid 100mg	Hypertension and Alzheimer's disease.	dermal and mucous membrane irritant, and a dermal sensitiser.It is a hazardous oil and should not be used on the skin. The oil should not be taken internally.		*Selected Medicinal Plants*, vol 1. Geneva: World Health Organization,		*bv gÄy Kiv nj* |

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
						1999. Page-95-104 a. *British Herbal Pharmacopoeia, 1996*. Exeter: British Herbal Medicine Association, 1996. , Page-59 *The Complete German Commission E Monographs*. Austin, Texas: American Botanical Council, 1998. Page-111 *British Herbal Pharmacopoeia*. Keighley: British Herbal Medicine Association, 1983.		
94.	M/s. Total Herbal & Neutraceuticals	Melatonin Capsule Melatonin 3mg Capsule	Jet lag, Sleeping difficulties Cancer prevention Regulation of sleep	No data are available, but in theory melatonin may be additive with medication that causes CNS depression. In addition, beta-blockers inhibit melatonin release, and this may be the mechanism by which beta-blockers cause sleep disturbance. Other drugs, including fluoxetine, ibuprofen and indomethacin, may also reduce nocturnal melatonin secretion. Melatonin may influence the effects of warfarin	New	USP – Dietary Supplement Compendium. USP32-NF27 Page1556 1. Dietary Supplements. Third Edition.	c‡qvRbxq †idv‡iÝ bvB wea‡q Av‡e`b bv gÄy Kiv †h‡Z cv‡i	c‡qvRbxq †idv‡iÝ bvB wea‡q Av‡e`b bv gÄy Kiv nj

						Page. 108-114		

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix Kwgw U (Jla wbqšY Kwgw Ui †UKwbK¨vj mve Kwgw Ui mvfvi wm×vš	mfvi wm×š
95.	M/s. Bexter Herbal & Nutraceuticals	Melatonin Avena Sativa 10mg Tablet	Sleep disorders in blind people. Sleeping problems in chindren with autism and ental retardation	Before taking melatonin, tell your doctor or pharmacist if you are allergic to it; or to other hormones; or if you have any other allergies.	New	1. British Herbal Pharmacopoea 2012. 2. WHO Monographs on commission E Monographs	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv হল |
96.	M/s. Total Herbal & Neutraceuticals	Neem Capsule Azadirachta indica 500mg Capsule	Anthelmintic, Antifungal, Antidiabetic, Antibacterial, Antiviral, Sedative		New	1. *The Ayurvedic pharmacopoeia of India. Part I. Vol. II*. New Delhi, Ministry of Health and Family Welfare, Department of Indian System of Medicine and Homeopathy, 1999. 3. *Indian medicinal plants. Vol. I*. New Delhi, Orient Longman, 1971.	c‡qvRbxq †idv‡iÝ bvB wea vq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i |	c‡qvRbxq †idv‡iÝ bvB wea vq Av‡e`b bv gÄyi Kiv nj |
97.	M/s. Total Herbal & Neutraceuticals	Bee pollen Capsule Bee pollen 500mg	Bee pollen may have antioxidant and anti-inflammatory activity, Produce antimicrobial effect, Renew skin. Boost immunity. Decrease allergy symptoms		New	Steben RE, Boudreaux P-1978; 18:221-226	c‡qvRbxq †idv‡iÝ bvB wea vq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i |	c‡qvRbxq †idv‡iÝ bvB wea vq Av‡e`b bv gÄyi Kiv nj |

bs	c¯ZKvi‡Ki bvg	JI‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
98.	M/s. Total Herbal & Neutraceuticals	Guto kola Hydrocotyle Capsule Centella Asiatica 500mg	possess mild diuretic, Antirheumatic, Dermatological, Peripheral vasodilator and vulnerary properties, Rheumatic conditions, Leprous ulcers and cicatrisation after surgery	Avoid use if hypersensitive to any of the ingredients of C. asiatica or gotu kola.	New	herbal drugs and phytopharmaceuticals Stuttgart medpharm . British Herbal Pharmacopoeia . W HO Monographs on selected Medicinal plants	Abyjgv`b Kiv †h‡Z cv‡i|	Abyjgv`b Kiv হল|
99.	M/s. Bexter Herbal & Nutraceuticals	Gotu Kola Centella asiatica 100mg Tablet	anti-inflammatory activity, exhibit anti-anxiety activity and cnhance mental function, stimulate blood vessel grouth in concetive tissue, increase the strength of the dermis and increaded keratinisation of the epidermis of the skin;.	Avoid use if hypersensitive to any of the ingredients of C. asiatica or gotu kola.	New	1.PDR for herbal medicine 2nd edition 2. WHO Monographs on Selected medicinal plants	Abyjgv`b Kiv †h‡Z cv‡i|	Abyjgv`b Kiv হল|
100.	M/s. Total Herbal & Neutraceuticals	Garcinia Cambogia+ Gymnema Sylvestre Capsule	**Exercise performance:** Hydroxycitric acid, a constituent in garcinia, may increase fat	Garcinia may lower blood sugar levels. Caution is advised in patients with diabetes (high blood sugar) or hypoglycemia (low blood sugar),	New	*A INDIAN PHARMACOPEAI*	Abyjgv`b Kiv †h‡Z cv‡i|	Abyjgv`b Kiv হল|

		Garcinia Cambogia (Garcinia Cambotia (Fruit) 750mg+ Gymnema	metabolism and enhance exercise performance. Additional study is needed to confirm these results 3**) Weight loss:** Evidence supporting	and in those taking drugs, herbs, or supplements that affect blood sugar. Serum glucose levels may need to be monitored by a		*2007 - Page no1397-1398* *b. United States*		

bs	c¯ZKvi‡Ki bvg	JI‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix Kwgw U (Jla wbqšY Kwgw Ui †UKwbK¨vj mve Kwgw Ui mvfvi wm×vš	mfvi wm×š
		Sylvestre (Gymnema Sylvestre Extract 134mg	hydroxycitric acid, the active ingredient in *Garcinia cambogia*, for weight loss is mixed. Additional study is warranted to clarify early findings	qualified healthcare professional, including a pharmacist, and medication adjustments may be necessary.		*Pharmacopeia 29 and National Formulary 24 and Supplements.* . c. PDR for *Herbal Medicines, Fourth Edition*. by Thomson Healthcare, Inc. 863,796		
101.	M/s. Bexter Herbal & Nutraceuticals	Lavender Lavendula angustifolia 400mg Capsule	Asthma, Brating bronechtis cough depression, anxiety and insomnia, muscle relaxation and agitation.	Lavender is contraindicated in Epilepsy, High blood pressure, Hypoglycemia & Kidney problems.	New	1. British Herbal Pharmacopoea 2. The complete German commission E monographs 3. Who Monographs on commission E	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv হল|

						Monographs		
102.	M/s. Bexter Herbal & Nutraceuticals	Chamomile Asteraceae (Daisy) 500mg Capsule	Boost the immune system, eliminate bad bacteria, anti-inflammatory, antimicrobial lowers blood cholesterol.	A relatively low percentage of people are sensitive to chamomile and develop allergic reactions. People sensitive to ragweed and chrysanthemums or other members of the *Compositae* family are more prone to develop contact allergies to chamomile, especially if they	New	1. British Herbal Pharmacopoea 2. The complete German commission E	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv* হল|

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
				take other drugs that help to trigger the sensitization.		monographs 3. Who Monographs on commission E Monographs		
103.	M/s. Bexter Herbal & Nutraceuticals	Goldenrod Solodago Virgauria 400mg Capsule	Gastroenteritis, Arthritis, Allergies	Contraindications are few for Goldenrod, although those who are pregnant or have serious heart or kidney problems should steer clear. There are no known side effects either alone or with other drugs or herbs, but this has not been studied extensively. Minor allergic reactions could occur in some people	New	1. British Herbal Pharmacopoea 1983 2. The complete German commission E monographs 3. Who Monographs on commission E Monographs 4. Oxford University press 2003	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv হল|
104.	M/s. Bexter Herbal & Nutraceuticals	Buchu Agathosma Betulina 500mg Tablet	Antiseptic, Aromatic, Carminative, Diaphoretic, Digestive Tonic	Avoid in kidney infections and diseases. Buchu's essential oil contains *diosmin* and *pulegone* which can cause irritation.	New	1.PDR for herbal medicine 2nd edition 2. WHO Monographs on Selected medicinal plants	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv হল|
105.	M/s. Bexter Herbal & Nutraceuticals	Coenzyme Coenzyme 100mg Tablet	Coenzyme Q-10 is a vitamin-like substance found throuhtour th especially in the heart, liver, kidney and pancreas. it is eaten in small amounts in meats and seafood. Coenzyme Q-10 can also e made in a laboratory. It is used	We have no information for Quenzyme Q10 contraindications.	New	1.PDR for herbal medicine 2nd edition 2. WHO Monographs on	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv হল|

			as medicine, Chest pain (angina) High blood Pressure & Hair l			Selected medicinal plants		

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
106.	M/s. Bexter Herbal & Nutraceuticals	Sage Salvia Officinalis 600mg Tablet	Calming and stimulating the nevous system, candia, colds, cought, dental abscesses	Contraindications have not been identified.	New	1.PDR for herbal medicine 2nd edition 2. WHO Monographs on Selected medicinal plants	Abšygv`b Kiv †h‡Z cv‡i	Abšygv`b Kiv hvq|
107.	M/s. Hamdard Laboratories (Waqf) Bangladesh Ltd.	Salvia officinalis *Salvia officinalis 250mg Capsule*	Indigestion due to stomach disorder, diarrhoea, loss of appetite, bloating, flatulence, inflammation of mouth & pharynx, excessive perspiration, Alzheimer's disease, hyperlipidemia and profuse lactation	Contraindicated during pregnancy.	New	PDR for Herbal Medicines, 2nd Edition, USA. pp. 655-657.	Abšygv`b Kiv †h‡Z cv‡i	Abšygv`b Kiv hvq|
108.	M/s. Bexter Herbal & Nutraceuticals	Bone Meal Bone Meal 500mg Tablet	Treating bone loss and osteoporosis, taking calcium for 30years after menopause might result in a 10% improvement in bone strength and a 50% overall reduction in bone break rates.	Arteriosclerosis with Occlusion of the Arteries, Kidney Stone, Kidney Disease, Sarcoidosis, High Amount of Phosphate in the Blood, High Amount of Calcium in the Blood, Excessive Amount of Vitamin D in the Body.	New	1.PDR for herbal medicine 2nd edition 2. WHO Monographs on Selected medicinal plants	Av‡e`b bv gÄwÿ Kiv †h‡Z cv‡i	Av‡e`b bv gÄwÿ Kiv nj	
109.	M/s. Bexter Herbal & Nutraceuticals	Parsley Petroselium Crispum 450mg Capsule	Obesity, Urnary tract infections. Digestive problems, Menstrual Problems	Not recommended for water retention due to heart or kidney condition. Renal inflammation Pregnancy (potential abortifacient).	New	1. The complete German commission E monographs 2. WHO Monographs on commission E Monographs 3. British Herbal Pharmacopoea 1996	Abšygv`b Kiv †h‡Z cv‡i	Abšygv`b Kiv hvq|

bs	c¯ZKvi‡Ki bvg	JI‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
110.	M/s. Hamdard Laboratories (Waqf) Bangladesh Ltd.	Pausinystalia yohimbe Pausinystalia yohimbe 10mg Capsule	Impotence, erectile dysfunction, debility and exhaustion.	It is contraindicated in liver and kidney diseases.	New	PDR for Herbal Medicines, 2nd Edition, USA. pp. 843-846.	Abšygv`b Kiv †h‡Z cv‡i	Abšygv`b Kiv হল|
111.	M/s. Hamdard Laboratories (Waqf) Bangladesh Ltd.	Mucuna pruriens Mucuna pruriens 500mg Capsule	Nervous debility, sexual debility, low sperm count, premature ejaculation, loss of libido, hypertension, diabetes, hypercholesterolemia, amenorrhoea, parkinson's disease and bell's palsy.	There is no known contraindication.	New	PDR for Herbal Medicines, 2nd Edition, USA. pp. 230-231.	Abšygv`b Kiv †h‡Z cv‡i	Abšygv`b Kiv হল|
112.	M/s. Hamdard Laboratories (Waqf) Bangladesh Ltd.	*Berberis vulgaris* *Berberis vulgaris 500mg Capsule*	Liver malfunctions, splenopathy, jaundice, gallbladder disease, indigestion, tuberculosis, piles, renal disease, urinary tract disorders, gout, rheumatism, arthritis, lumbago, malaria, leishmaniasis and opium & morphine withdrawal.	There is no known contraindication.	New	PDR for Herbal Medicines, 2nd Edition, USA. pp. 61-62.	Abšygv`b Kiv †h‡Z cv‡i	Abšygv`b Kiv হল|
113.	M/s. Hamdard Laboratories (Waqf) Bangladesh Ltd.	*Passiflora incarnata* *Passiflora incarnate 400mg Capsule*	Nervous excitement, insomnia, depressive states such as hysteria, general nervous agitation, and nervous gastrointestinal complaints.	There is no known contraindication.	New	PDR for Herbal Medicines, 2nd Edition, USA. pp. 573-575.	Abšygv`b Kiv †h‡Z cv‡i	Abšygv`b Kiv হল|
114.	M/s. Hamdard Laboratories (Waqf) Bangladesh Ltd.	Aesculus hippocastanum Aesculus hippocastanum 300mg	Venous insufficiency, chronic venous insufficiency (such as pain and a sensation of heaviness in the legs, nocturnal cramps in the claves, pruritis and swelling of the legs), post-	There is no known contraindication.	New	PDR for Herbal Medicines, 2nd Edition, USA. pp.	Abšygv`b Kiv †h‡Z cv‡i	Abšygv`b Kiv হল|

		Capsule	traumatic and post-operative soft tissue swelling, hemorrhoids, lumber and low back pain, venous back pressure, varicositis, phlebitis			403-407.		

bs	c¯ZKvi‡Ki bvg	JI‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
115.	M/s. Hamdard Laboratories (Waqf) Bangladesh Ltd.	Crataegus oxycanthus Crataegus oxycanthus 300mg Capsule	Cardiac insufficiency, senile cardiac insufficiency, dysrrhythmia, chronic cor pulmonale, angina pectoris, heart insufficiency due to old age, myocarditis, arteriosclerosis, nervous heart complaints, mental excitement, nervous debility, hypertension and stresses.	It is contraindicated in children under 12 years old and during the first trimester of pregnancy.	New	PDR for Herbal Medicines, 2nd Edition, USA. pp. 271-275.	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv হল|
116.	M/s. Hamdard Laboratories (Waqf) Bangladesh Ltd.	*recutita* *Matricaria recutita 350mg Capsule*	Cough, cold, bronchitis, fevers, inflammation of skin, mouth, pharynx, stomach, duodenum, colon, rectum & sexual organs, irritation of upper respiratory tract, wounds and burns	It should not be taken by anyone with a known allergy to its components.	New	PDR for Herbal Medicines, 2nd Edition, USA. pp. 331-335.	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv হল|
117.	M/s. Hamdard Laboratories (Waqf) Bangladesh Ltd.	*UncaPria tomentosa* *Uncaria tomentosa 500mg Capsule*	Rheumatic complaints, gastritis, inflammation, asthma, wounds, menstrual irregularity and crohn's disease.	There is no known contraindication.	New	PDR for Herbal Medicines, 2nd Edition, USA. pp. 160-162.	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv হল|
118.	M/s. Hamdard Laboratories (Waqf) Bangladesh Ltd.	Solidago virgaurea *(‡Mv‡ìb iW)* Solidago virgaurea (‡Mv‡ìb iW) 450mg Capsule	Urinary calculi, infections of urinary tract, burning urination, oliguria, renal insufficiency and liver disorders, inflammation of kidney, kidney and bladder stones.	There is no known contraindication.	New	PDR for Herbal Medicines, 2nd Edition, USA. pp. 289-291.	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv হল|
119.	M/s. Hamdard Laboratories	Bitter Melon	The special preparation of standardized fruit extract of *Momordica charantia* makes the	Generally there is no known contraindication.	New	*Mosby's Drug Consult*	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv

	(Waqf) Bangladesh Ltd.	Bitter Melon 500mg Capsule	product as a perfect natural remedy for balancing the body. It acts as a rejuvenator in pancreatic activity and restores pancreatic functions. It promotes endogenous insulin production and augments alternative modes of glucose handling and eventually reduces blood			(2006), Section III, Herbal and Supplement Information. p.		হল।

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
			sugar. It also acts as a blood purifier for beautiful & healthy skin, combats acne and pimples, boosts up immune system and lowers blood cholesterol level. It acts as carminative, laxative and stomachic and very effective in stomach disorders			III-13.		
120.	Acme Laboratories (Herbal Division)	Peppermint oil 187mg Capsul Mentha piperita Linn.	In addition to use as a seasoning and flavoring, peppermint is used to treat irritable bowel syndrome (IBS) and other GI conditions. Menthol is available in numerous commercial preparations used to treat respiratory tract infections and topically for its cooling and warming action to relieve pain.	Contraindicated in patients with achlorhydria. Also Contraindicated for infants due to the potential risk of spasm of the tongue or respiratory arrest. Side effect : None known according to commission E.	New	PDR for herbal Medicines	Abyjgv`b Kiv †h‡Z cv‡i|	Abyjgv`b Kiv হল|
121.	Acme Laboratories (Herbal Division)	Zinger Powder 250mg Capsule Zingiber Officinale Linn 250mg	Loss of appetite, Travel Sickness, Dyspeptic Complaints	Contraindication: Because of its cholagogic effect, the drug should not be taken in presence of gallstone condition. Side effects: Ginger can cause hypersensitivity reaction resulting in dermatitis. Large overdoses can causes central nervous system depression and cardiac arrhythmias.	New	PDR for herbal Medicines 2nd Edition Page 339-340 The Complete German Commission E Monograph Page 135-136	Abyjgv`b Kiv †h‡Z cv‡i|	Abyjgv`b Kiv হল|
122.	Acme Laboratories	Ginseng Syrup	1. Adaptogen & general tonic, 2. Increase athlette per forman &	Contraindication: Caution should be taken in patients with cardiovascular disease or	New	The Complete German	Abyjgv`b Kiv †h‡Z cv‡i|	Abyjgv`b Kiv

	(Herbal Division)	Ginseng Standardized Concerted Liquid Extract (Panax Ginseng) 300mg/5ml	endurance, 3. Vitality, 4. Fatique & Devility, 5. Endancement of physical & mental capacity	diabetes. Side effects: No known (at therapeutic doses), but massive over dosage can bring about ginseng abuse syndrome which is		Commission E monographs,Herbal Medicine Expanded E Monographs,		হল।

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
				characterized by hypertension, insomnia and hypertonia		British Herbal Pharmacopoeia 1996 PDR for herbal Medicines		
123.	Renata Limited (Herbal Division) Kashor, Hobirbari, Bhaluka, Mymensingh.	Plantain Plantain powder 250 mg per capsule Days 1-3: 1 capsule daily Days 4-7: 2	Manage tobacco cravings, reduce tension associated with smoking cessation, and facilitates the release of accumulated tobacco toxins from the body.	Should be used cautiously or is contradicted by Elderly people, children, and people with compromised immune systems.	New	PDR for Herbal Medicines (4th edition)	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv হল|
124.	Drug International Ltd (Herbal Division)	**Mucolytic-Cough Relieving** (Ivy Leaf Dry Extract 0.750gm + Thyme Dry Extract 1.00gm) / 100ml Syrup	Mucolytic and expectorant, helps to relief symptoms of catarrh (inflammation) of the respiratory tract accompanied by coughing.	It is contraindicated in patients with known hypersensitivity to any other components of this syrup. In rare cases, this cough syrup may have a laxative effect. In rare cases skin allergy or gastric disturbance may occur.	New	PDR for Herbal Medicines 4th edition.	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv হল|
125.	Incepta Herbal & Nutricare Ltd.	Chitosan hydrochloride 500mg Capsule Each capsule contains 500 mg Chitosan Hydrochloride BP	*Possibly Effective for:* Patients with kidney failure who are on long-term hemodialysis. When taken by these patients, chitosan may reduce high cholesterol; help to correct anemia; and improve physical strength, appetite, and sleep. Treating periodontitis, a dental condition. Applying chitosan ascorbate directly to the	**Contraindications:** None well established. **Side Effects:** Some studies have proven chitosan causes gas, bloating, and diarrhea (due to fermentation of the fat in the large intestine).more long term and short term studies are required to establish side effects.	New	Pharma Euro	c‡qvRbxq †idv‡iÝ bvB wea vq Av‡e`b bv gÄyi Kiv †h‡Z cv‡i	c‡qvRbxq †idv‡iÝ bvB wea vq Av‡e`b bv gÄyi Kiv nj	

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
			gums seems to help in the treatment of periodontitis. Helping to remake tissue after plastic surgery. Applying N-carboxybutyl chitosan directly seems to help donor site tissue rebuild in plastic surgery.					
126.	Incepta Herbal & Nutricare Ltd.	Softgel Capsule: Phytosterols 400mg+Fish Oil Omega-3 Fatty Acids equivalent to 270mg Eicosapentiaenoic Acid and 180mg Docosahexaenoic Acid Each soft gelatin capsule contains 900mg of fish oil including 450mg heart-healthy omega-3 fatty acids EPA (Eicosapentaenoic acid) 270mg & DHA	• Maintenance of normal blood pressure. • Support the body's natural anti-inflammatory response. • Support normal, healthy cholesterol Levels. • Reduce the risk of coronary heart disease. • Support joint & heart health. • Natural anti-inflammatory. • Improvement of brain function.	**Contraindications:** Known hypersensitivity to any component of the formulation. **Side Effects:** • Mild to severe gastrointestinal upset or diarrhea (caused by fishy taste/fishy smell of the formulation) • Acid reflux • Heartburn • Indigestion • Bloating	New	Mosby's Drug Consult (Page: ||| - 82)	Kwc‡qvRbxq †idv‡iÝ bvB weavq Av‡e`b bv gÄwÿ Kiv †h‡Z cv‡i	Kwc‡qvRbxq †idv‡iÝ bvB weavq Av‡e`b bv gÄwÿ Kiv nj	
127.	Incepta Herbal & Nutricare Ltd.	Vasak 5gm+Tulsi 3gm+Sunthi 3gm+Karkat Sringi 3gm+Mulethi 3gm+Draksha 3gm+Baheda 2gm+Pippali 1gm+Dalchini 1gm+Laung 0.7gm+Kali Maricha 0.2gm+Pudina Sat 0.1gm / 100mL Syrup Each 5 ml syrup contains	It liquefies phlegm. It is effective for all kinds of cough such as dry irritable cough, allergic & smokers cough. It soothes the throat irritation and relieves hoarseness.	**Contraindications:** There is no evidence available on contraindication but it may happen in -patients who are hypersensitive to any of its ingredients. **Side Effects:** Herbal medicine is clinically proven as	New	N/A	c‡qvRbxq †idv‡iÝ bvB weavq Av‡e`b bv gÄwÿ Kiv †h‡Z cv‡i	c‡qvRbxq †idv‡iÝ bvB weavq Av‡e`b bv gÄwÿ Kiv nj	

		extracts of Vasak (*Adhatoda vasica*) 250mg, Tulsi (*Ocimum sanctum*) 150mg,		safe & well tolerated. In the recommended doses, side effects are rare.				

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix Kwgw U (Jla wbqšY Kwgw Ui †UKwbK¨vj mve Kwgw Ui mvfvi wm×vš	mfvi wm×š
		Sunthi (*Zingiber offcinale*) 150mg, Karkatshringi (*Pistacia integerrima*) 150mg, Mulethi (*Glycerrhiza glabra*) 150mg, Draksha (*Vitis vinifera*) 150mg, Baheda (Terminalia belerica) 100mg, Pippali (Piper longnum) 50mg, Dalchini (Cinnamomum zeylanicum) 50mg, Laung (*Syzygium aromaticum*) 35mg, Kali Maricha (*Piper nigrum*) 10mg and Pudia sat (*Mentha spicata*) 5mg						
128.	Incepta Herbal & Nutricare Ltd.	Tulsi 1gm+Mulethi 1gm+Banaphsa 1gm+Kantkari 1gm+Talispatra 1gm+Sunthi 0.5gm+Pippali 0.5gm+Vasaka 0.5gm+Sathi 0.5gm+Pudina satva 0.06gm+Shudha Each 5ml syrup contain extracts derived from Tulsi (*Ocimum Sanctum*) 50.0mg, Mulethi (*Glycerrhiza glabra*) 50.0mg, Banaphsa (*Viola odorata*) 50.0mg, Kantkari (*Solanum xanthocarpum*) 50.0mg, Talispatra (*Abies webbiana*) 50.0mg, Sunthi	Effective in different kinds of cough and throat imitation.	**Contraindications:** There is no evidence available on contraindication but it may happen in - patients who are hypersensitive to any of its ingredients. **Side Effects:** Herbal medicine is clinically proven as safe & well tolerated. In the recommended doses, side effects are rare.	New	N/A	c‡qvRbxq †idv‡iÝ bvB wea vq Av‡e`b bv gÄy Kiv †h‡Z cv‡i	c‡qvRbxq †idv‡iÝ bvB wea vq Av‡e`b bv gÄy Kiv nj	

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix Kwgwu (Jla wbqšY Kwgwui †UKwbK¨vj mve Kwgwui mvfvi wm×vš	mfvi wm×š
		(*Zingiber officinale*) 25.0mg, Pippali (*Piper longum*) 25.0mg, Vasaka (*Adhatoda vasica*) 25.0mg, Shati (*Zingiber officinale*) 25.0mg, Pudina satva 3.00mg, Shudha madhu (Honey) 1.75gms	Effective in different kinds of cough and throat imitation.	**Contraindications:** There is no evidence available on contraindication but it may happen in - patients who are hypersensitive to any of its ingredients. **Side Effects:** Herbal medicine is clinically proven as safe & well tolerated. In the recommended doses, side effects are rare.	New	N/A	c‡qvRbxq †idv‡iÝ bvB wewfbœ Av‡e`b bv gÄy Kiv †h‡Z cv‡i|	c‡qvRbxq †idv‡iÝ bvB wewfbœ Av‡e`b bv gÄy Kiv nj|
129.	Incepta Herbal & Nutricare Ltd.	Purified Guggulu 0.7gm+Draksha 0.7gm+Vishnu Priya 0.5gm+Jufa 0.5gm+Guduchi 0.4gm+Vasaka 0.3gm+Yashti-madhu 0.3gm+Gojihva 0.2gm+Neelapushpa 0.2gm+Triphala 0.18gm+Trikatu 0.18gm+Vidanga 0.16gm+Kantakari 0.16gm+Taja 0.16gm+Navasagara 0.06gm / 100mL Syrup Each 5ml syrup contains extracts of Madhu (Purified honey) 1.25g, Purified Guggulu (*Balsamodendron mukul*) 35mg, Draksha ext. (*Vitis vinifera*) 35 mg,	• Cough associated with acute and chronic upper and lower respiratory tract infections • Smoker's cough • Cough due to chronic obstructive pulmonary disease (COPD) • Supports healthy bronchial passageway and breathing function. • Supports the body's normal response to occasional irritations of the lung and bronchial passages. • Natural alcohol free support. This non-drowsy is well tolerated in children too.	**Contraindications:** There is no evidence available on contraindication but it may happen in - patients who are hypersensitive to any of its ingredients**.** **Side Effects:** It is not known to have any side effects if taken as per the prescribed dosage**.**	New	N/A	c‡qvRbxq †idv‡iÝ bvB wewfbœ Av‡e`b bv gÄy Kiv †h‡Z cv‡i|	c‡qvRbxq †idv‡iÝ bvB wewfbœ Av‡e`b bv gÄy Kiv nj|

bs	c¯ZKvi‡Ki bvg	JI‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	nvevj GWfvBRix KwgwU (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi mvfvi wm×vš	mfvi wm×š
		Vishnupriya (*Ocimum sanctum*) 25 mg, Jufa (*Hyssopus officinalis*) 25mg, Guduchi (*Tinospora cordifolia*) 20mg, Vasaka (*Adhatoda vasica)* 15mg, Yashti madhu (*Glycyrrhiza glabra*) 15mg, Gojihva (*Onosma bracteatum*) 10mg, Neelapuspha (*Viola odorata*) 10mg, Triphala 9mg, Trikatu 9mg, Vidanga (*Embelia ribes*) 8mg, Kantakari (*Solanum xanthocarpum*) 8mg, Taja (*Cinnamomum cassia*) 8mg, Navasagara 3mg						