Jla wbqšY Kwgwui 22 †de«qvix 2017 Zwi‡L Abyôz 247 Zg mfvi Kvhweeiyx

¯v¯¨ I cwievi Kj¨vY gšYvj‡qi mwPe Rbve †gvt wmivRyj Bmjvg Gwu mfvcwZ‡Z Jla wbqšY Kwgwui 247Zg mfv weMZ 22 †de«qvix 2017 Zwi‡L weKvj 4:00 NwUKvq gšYvj‡qi mfv K‡¶ Abyôz nq|

mfvi কমিটির নিম্নবর্ণিত সদস্যগণ উপস্থিত ছিলেন (জ্যেষ্ঠতার ক্রমানুসারে নয়) ঃ

1| Rbve mykxgvi iÄb †Nvl, cwZwbwa, evsjv‡`k dvgwmDwUK¨vjm B‡¤úvUvm G‡mwm‡qkb, XvKv|
2| Aa¨vcK Wvt Kvgiæj nwmwb Lvb, DcvPvh, e½eÜz‡kL gyRe †gwW‡Kj wekwe`¨vjq, XvKv|
3| ‡gRi †Rbv‡ij †gvt Ave`yj Avjx wgqv, Kbmvj‡U›U wdwRwkqvb ‡Rbv‡ij, evsjv‡`k AvgW †dv‡mm †gwW‡Kj mwf‡mm|
4| cvifxb AvKZvi, hyM¥mwPe, Rb¯v¯¨, ¯v¯¨ I cwievi Kj¨vY gšYvjq, evsjv‡`k mwPevjq, XvKv|
5| Wvt †gvt Ryjvdwdkvi Avjx, cwiPvjK-3, cavbgšxi Kvhvjq, XvKv|
6| Wvt GBPwe Gg †Mvjvg gvngy`, cwiPvjK, wiPvm †Uwbs GÛ Bfvjyqkb, jvBf ÷K wiPvm Bbw÷wUDU, c‡¶ gnvcwiPvjK, cwY m¤ú` Awa`ßi, XvKv|
7| ‡gvt gwRyei ingvb cv‡Uvqvix, AwZwi³ cwiPvjK, gv`K`e¨ wbqšY Awa`ßi, †ZRMvwu, XvKv|
8| Aa¨vcK Wvt †gvt BmgvBj Lvb, Wxb, wPwKrmv Abyl`, XvKv wekwe`¨vjq I cwZwbwa, dvgv‡Kvjvwr wefvM, XvKv †gwW‡Kj K‡jR|
9| Aa¨vcK dwi`v †eMg, wKwbK¨vj dv‡gmx I dvgv‡Kvjvwr wefvM, XvKv wekwe`¨vjq|
10| Aa¨vcK W. Gm Gg Avãyi ingvb, Wxb, dv‡gmx Abyl`, XvKv wekwe`¨vjq|
11| Aa¨vcK Wvt Rwnki †nvmvBb Mwje, Pg I †hŠb †ivM we‡klÁ, m¨vi mwjgyivn †gwW‡Kj K‡jR, XvKv|
12| Wvt gÄyi Avng`, mnKvix cwiPvjK (nvmcvZvj-2), ¯v¯¨ Awa`ßi, gnvLvjx, XvKv|
13| Wvt †gv‡gbyi nK, mv‡eK wmwbqi mn mfvcwZ, evsjv‡`k Jla wkí mvgwZ Ges e¨e¯vcbv cwiPvjK, †Rbv‡ij dvgwmDwUK¨vjm wjt|
14| Rbve Gg †gvQv‡Ïk †nv‡mb, cwZwbwa, evsjv‡`k dv‡gmx KvDwÝj, XvKv|
15| ‡gRi †Rbv‡ij †gvt ‡gv¯wdRyi ingvb, gnvcwiPvjK, Jla ckvmb Awa`ßi, XvKv|

mfvq Av‡jvP¨ welq mgm¨vbgi¦c t

1| Jla wbqšY Kwgwui 246 Zg mfvi Kvhweeiyx wbwðZKiY|

2| Jla wbqšY Kwgwui †UKwbK¨vj mve Kwgwui MZ 05-02-2017 Zwi‡L Abyôz mfvi mycwikmgm¨t
   2.1 Pseudoephedrine- Gi Avg`vbx ‡ivaK‡í Pseudoephedrine I Gi K¤¤‡bkb Jla ewZjKiY cm‡½|
   2.2 ¯vbxqfv‡e Drcv`‡bi Rb¨ Av‡ew`Z 52 wU wnDg¨vb Jl‡ai †iwR‡÷k‡bi Abyôgv`‡bi mycwikmg‡ni wel‡q Av‡jvPbv I wm×vš MnY cm‡½|
   2.3 Avg`vbxi Rb¨ Av‡ew`Z 20 wU wnDg¨vb Jl‡ai †iwR‡÷k‡bi Abyôgv`‡bi mycwikmg‡ni wel‡q Av‡jvPbv I wm×vš MnY cm‡½|
   2.4 Avg`vbxi Rb¨ Av‡ew`Z 20 wU †f‡Uwibvix Jl‡ai †iwR‡÷k‡bi Abyôgv`‡bi mycwikmg‡ni wel‡q Av‡jvPbv I wm×vš MnY cm‡½|

3| Jla wbqšY Kwgwui ‡gwW‡Kj wWfvB‡mi †UKwbK¨vj mve Kwgwui MZ 15-02-2017 Zwi‡L Abyôz mfvi mycwikmgm¨t

   3.1 ¯vbxqfv‡e Drcv`‡bi Rb¨ Av‡ew`Z 3 wU ‡gwW‡Kj wWfvB‡mi †iwR‡÷k‡bi Abyôgv`‡bi mycwikmg‡ni wel‡q Av‡jvPbv I wm×vš MnY cm‡½|

3.2 Avg`vbxi Rb¨ Av‡ew`Z 119 wU ‡gvW‡Kj wWfvB‡mi †iwR‡÷k‡bi Abyôvb mgvwikmg¥ni wel‡q Av‡jvPbv I wm×vš MnY cm‡½|

3.3 Avg`vbxi Rb¨ Av‡ew`Z 8 wU AvBwfwW wi‡q‡RÛU-Gi †iwR‡÷k‡bi Abyôvb mgvwikmg¥ni wel‡q Av‡jvPbv I wm×vš MnY cm‡½|

4| nvevj GWfvBRix KwgwUi (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwU) MZ 09-11-2016 Zwi‡L Abyôv‡Z mfvi mgvwikmgn¥t

4.1 nvevj Jl‡ai Rb¨ c‡e wbawiZ 16(‡lvj)wU eB Gi cvkvcwk †idv‡iÝ eB wnmv‡e USP-DSC, Indonesian Herbal Pharmacopoeia, Vietnam Herbal Pharmacopoeia & North Korean Herbal Pharmacopoeia ‡K AšÍf~³ KiY cm‡½|

4.2 Lmov nvevj Jl‡ai wbeÜb bxwZgvjv ms‡kwaZ AvKv‡i Abyôvb mgvwi‡ki wel‡q wm×vš MnY cm‡½|

4.3 ¯vbxqfv‡e Drcv`‡bi Rb¨ Av‡ew`Z 29 wU nvevj Jl‡ai †iwR‡÷k‡bi Abyôvb mgvwikmg¥ni wel‡q Av‡jvPbv I wm×vš MnY cm‡½|

সভাপতি উপস্থিত সকলকে স্বাগত জানিয়ে সভার কার্যক্রম শুরু করেন। অতঃপর তিনি সদস্য-mwPe ‡gRi †Rbv‡ij †gvt ‡gv¯wdRyi ingvb, gnvcwiPvjK, Jla ckvmb Awa`ßi‡K Av‡jvP¨mPx Abyôvqx welqmgn¥ Dc¯vcb Kiv Rb¨ Abyôviva K‡ib|

1| **Jla wbqšY KwgwUi 246 Zg mfvi Kvhweeiyx wbwðZKiY cm‡½|**

weMZ 18-10-2016 Zwi‡L Abyôv‡Z Jla wbqšY KwgwUi 246 Zg mfvi Kvhweeiyx mfvq Dc¯vcb Kiv nq| Kvhweeiyx mwVKfv‡e wjwce× n‡q‡Q e‡j m`m¨MY gZ ckvk K‡ib|

mfvq সর্বসম্মতিক্রমে ২৪6 Zg mfvi Kvhweeiyx wbwðZ Kiv nq|

2| **Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwUi MZ 05-02-2017 Zwi‡L Abyôv‡Z mfvi mgvwikmgn¥t**

2.1 Pseudoephedrine-Gi Avg`vbx ‡ivaK‡í Pseudoephedrine I Gi K¤¢bkb Jla ewZjKiY cm‡½|

<u>mfvi Av‡jvPbvt</u>

m`m¨-mwPe ‡gRi †Rbv‡ij †gvt ‡gv¯wdRyi ingvb, gnvcwiPvjK, Jla ckvmb Awa`ßi mfv‡K AewnZ K‡ib †h, gvbbxq cavbgšxi Kvhvjq n‡Z 07-08-2016 Zwi‡Li GK c‡Îi gva¨‡g Pseudoephedrine e¨envi K‡i Bqvev ev Ab¨ ‡Kvb †bkv RvZxq `e¨ ˆZwi Kiv m¤e wKbv, †m wel‡q gv`K`e¨ Awa`ß‡ii we‡klÁ, dv‡gmx we‡klÁ I dvgv‡KvjRx we‡klÁ‡`i AskMn‡Y GKwU KwgwU MVb K‡i gZvgZ MnYce~K Zv †ciY Kivi Abyôviva Kiv nq|

GB cwi‡cwÿ‡Z weMZ 10-08-2013 Zwi‡L XvKv †gvW‡Kj K‡j‡Ri Aa¨ÿ Aa¨vcK Wvt †gvt BmgvBj Lvb-‡K Avnevqk K‡i we‡klÁ‡`i mgš‡q 06(Qq) m`m¨ wewkó GKwU we‡klÁ KwgwU MVb Kiv nq| D³ KwgwU we¯wiZ cwZ‡e`b c`vb K‡ib Ges gZvgZ c`vb K‡ib †h,

K| Pseudoephedrine *e¨envi K‡i c¯ZKZ.Jla †_‡K* Pseudoephedrine *†K Avjv`v Kiv hvq*| Pseudoephedrine এর সাথে অন্যান্য সক্রিয় উপাদান সমন্বয়ে উৎপাদিত ঔষধ Fixed Dose Combination (FDC) *n‡ZI* Pseudoephedrine *Avjv`v Kiv hvq*|

L| Pseudoephedrine হতে রাসায়নিক প্রক্রিয়ায় মাধ্যমে Methamphetamine *‰Zix Kiv hvq, hv* ইয়াবার একটি সক্রিয় উপাদান। সুতরাং Pseudoephedrine *e¨envi K‡i Bqvev bvgK †bkv RvZxq c`v_ Drcv`b Kiv m¤e*|

Jla ckvmb Awa`ßi n‡Z D³ KwgwUi wm×vš gvbbxq cavbgšxi Kvhvjq‡K AewnZ Kivi ci gvbbxq cavbgšxi Kvhvj‡qi c‡Îi cwi‡cÿ‡Z Pseudoephedrine *wgwk‡q wewfb Jlamn Bqvev ˆZwii Awf‡hv‡Mi wel‡q mwPe, ¯v¯¨ I cwievi Kj¨vY gšYvj‡qi mfvcwZ‡Z 07-12-2016 Zwi‡L ¯v¯¨ I cwievi Kj¨vY gšYvj‡q GKwU mfv Abyôv nq| D³ mfvq Jla wbqšY KwgwUi cieZx mfvi mycwi‡ki gva¨‡g Ges Jla ckvmb Awa`ßi I gv`K`e¨ wbqšY Awa`ß‡ii †hš_ D‡`¨v‡M* Pseudoephedrine *Avg`wb e‡Üi D‡`¨vM MnY Kivi wm×vš Mp,xZ nq*|

D³ wm×v‡š Av‡jv‡K Jla wbqšY KwgwUi mfvi 247Zg mfvq Dc¯vc‡bi wbwg‡Ë †UKwbK¨vj mve-KwgwUi mfvq Pseudoephedrine *Avg`wb e‡Üi D‡`¨vM MnY Kivi welqwU Dc¯vcb Kiv nq*|

wZwb mfv‡K Av‡iv AewnZ K‡ib †h, evsjv‡`‡k* Pseudoephedrine *Ges Gi wewfb Kw¤‡bkb U¨ve‡jU I wmivc AvKv‡i wb¤i¡c Abyôgw`Z Av‡Q :

1. Pseudoephedrine 60 mg Tablet
2. Pseudoephedrine Hydrochloride Pelletes
3. Pseudoephedrine 60 mg + Triprolidine 2.5 mg Tablet
4. Fexofenadine 60mg + Pseudoephedrine 120 mg ER Tablet
5. Fexofenadine 60mg + Pseudoephedrine 120 mg Tablet
6. Loratadine 5mg + Pseudoephedrine Hydrochloride 120 mg Tablet
7. Loratadine 10mg + Pseudoephedrine Hydrochloride 240 mg Tablet
8. Desloratadine 2.5mg + Pseudoephedrine 120 mg Tablet
9. Desloratadine 5mg + Pseudoephedrine 240 mg Tablet
10. Guaiphenesin 100mg + Pseudoephedrine 30 mg + Triprolidine 1.25 mg/5ml Syrup
11. Dextromethorphan 10mg + Pseudoephedrine Hydrochloride 30 mg + Triprolidine Hydrochloride 1.25mg/5 ml Syrup

Pseudoephedrin-*Gi Avg`vbx eÜ K‡i‡Z n‡j Gme Jl‡ai Drcv`b I †iwR‡÷kb ewZ‡ji welqwU we‡ePbv Kivi c‡qvRb*| *†UKwUK¨vj mve-KwgwUi 05 ‡deªqvix, 2017 Zwi‡L Abyôv mfvq GZ`welqwU*

উপস্থাপন করা হলে, সদস্যগণ সর্বসম্মতিক্রমে Pseudoephedrine-Øviv ^Zwi mKj †Wv‡Rm d‡gi Jl‡ai †iwR‡÷kb ewZj Kivi mgvwik K‡ib|

Jla wbqšY KwgwUi m`m¨-mwPe Pseudoephedrine-Øviv ^Zwi mKj †Wv‡Rm d‡gi Jl‡ai †iwR‡÷kb ewZj Kivi ‡UKwbK¨vj mve-KwgwUi mgvwikwU mfvq Dc¯vcb K‡ib| m`m¨MY G wel‡q we¯wiZ Av‡jvPbv K‡i Pseudoephedrine-Gi Ace¨envi †ivaK‡í Gi Avg`vbx e‡Üi wbwg‡Ë Pseudoephedrine-Øviv ^Zwi mKj †Wv‡Rm d‡gi Jl‡ai †iwR‡÷kb ewZj Kivi c‡ÿ gZ cKvk K‡ib| m`m¨MY Av‡ivI e‡jb †h, BwZg‡a¨ Pseudoephedrine-Øviv c¯ZKZ.Jlamgn- evRvi n‡Z cZ¨vnv‡ii Rb¨ wKQzUv mg‡qi c‡qvRb n‡e| ZvB Pseudoephedrine e¨envi K‡i Jla c¯ZKviKMY‡K 03(wZb) gvm mgq †`qv †h‡Z cv‡i, hv‡Z Zuiv Pseudoephedrine-Øviv ^ZwiKZ. Jla evRvi n‡Z cZ¨vnvice-K aÿsm Ki‡Z cv‡ib|

সভায় সর্ব সম্মতিক্রমে নিম্নলিখিত wm×vš Mp,xZ nq t

K| Pseudoephedrine- Gi Avg`vbx eÜK‡í Pseudoephedrine-Øviv ^Zwi mKj †Wv‡Rm d‡gi Jl‡ai †iwR‡÷kb ewZj Kiv nj|

L| AvMvgx 3(wZb) gv‡mi g‡a¨ evRvi †_‡K Pseudoephedrine-Øviv ^Zwi mKj Jla cZ¨vnvi ce-K aÿsm K‡i Jla ckvmb Awa`ßi‡K AewnZ Ki‡Z n‡e|

M| Jla ckvmb Awa`ßi I gv`K `e¨ wbqšY Awa`ßi n‡Z Kv÷gm KZc.ÿ‡K Pseudoephedrine-Qvo bv Kivi Rb¨ cÎ ‡ciY Ki‡Z n‡e|

N| Pseudoephedrine-Gi Ace¨envi †iva K‡í gwbUwis-Gi Rb¨ (1) cwiPvjK, Jla ckvmb Awa`ßi, (2) AwZwi³ cwiPvjK, gv`K `e¨ wbqšY Awa`ßi, (3) cwZwbwa, evsjv‡`k Jla wkí mwgwZ Ges Wxb, dv‡gmx Abyôv`, XvKv wek¦e`¨vjq mgš‡q GKwU KwgwU MVb Kivi wm×vš Mp,xZ nq| KwgwUi m`m¨e„›` gIRyôKZ. Pseudoephedrine-Gi cwigvb wbbq I Gi mwVK e¨envi gwbUwis Ki‡eb|

2.2 ¯vbxq fv‡e Drcv`‡bi Rb¨ Av‡ew`Z 52(evqvb)wU wnDg¨vb Jl‡ai †iwR‡÷k‡bi Abyôvb`v‡bi mgvwikmg‡ni wel‡q Av‡jvPbv I wm×vš MnY cm‡½|

¯vbxq Drcv`‡bi Rb¨ †iwR‡÷k‡bi wbwg‡Ë `wLjKZ.52wU bZzb Jla mfvq Dc¯vcb Kiv n‡j m`m¨MY KZ„K Jla¸wji Safety, Efficacy and Usefulness we‡ePbv K‡i বিস্তারিত আলোচনাক্রমে wb¤wjwLZ wm×vš Mp,xZ nq t

K) Abyôvb`v‡bi wZb - 19 (Dwbk)wU;
L) ¯wMZ I ‡idv‡iÝ I Av‡iv Z_¨mn cieZx mfvq Av‡e`b `wLj Kivi Rb¨ ejv nq-08(AvU)wU Ges
M) bvgÄyôKZ.-25(cuPk)wU; (Annex-A)|

2.3 Avg`vbxi Rb¨ Av‡ew`Z 20 wU wnDg¨vb Jl‡ai †iwR‡÷k‡bi Abyôgv`‡bi mycwikmg~ni wel‡q Av‡jvPbv I wm×vš MnY cm‡½|

Avg`vbxi Rb¨ †iwR‡÷k‡bi wbwg‡Ë Av‡ew`Z 20wU bZzb wnDg¨vb Jl‡ai wel‡q mfvq Dcw¯Z m`m¨MY KZ„K we¯wrat আলোচনাক্রমে wb¤wjwLZ wm×vš Mp„xZ nq t

K) Abyôgw`Z - 18wU Ges
L) ¯wMZ I ‡idv‡iÝ I Av‡iv Z_¨mn cieZx mfvq Av‡e`b `wLj Kiv Rb¨ ejv nq-02wU; (Annex-B)|

2.4 Avg`vbxi Rb¨ Av‡ew`Z 20 wU †f‡Uwibvix Jl‡ai †iwR‡÷k‡bi Abyôgv`‡bi mycwikmg~ni wel‡q Av‡jvPbv I wm×vš MnY cm‡½|

Avg`vbxi wbwg‡Ë ‡f‡Uwibvix Jl‡ai †iwR‡÷k‡bi Rb¨ Av‡ew`Z 20wU bZzb Jl‡ai wel‡q mfvq m`m¨MY Jla¸wji Jla wbqšY KwgwUi †UKwbK¨vj mve-KwgwUi mycwi‡ki Av‡jv‡K we¯wiZ আলোচনাক্রমে 20wU Jla Avg`vbx †iwR‡÷k‡bi Rb¨ Abyôgv`b K‡ib (Annex-C)|

3| Jla wbqšY KwgwUi ‡gwW‡Kj wWfvB‡mi †UKwbK¨vj mve KwgwUi MZ 15-02-2017 Zwi‡L Abyôq mfvi mycwikmgn~ t

3.1 ¯vbxq fv‡e Drcv`‡bi Rb¨ Av‡ew`Z 3 wU ‡gwW‡Kj wWfvB‡mi †iwR‡÷k‡bi Abyôgv`‡bi mycwikmg~ni wel‡q Av‡jvPbv I wm×vš MnY cm‡½|

¯vbxq Drcv`‡bi wbwg‡Ë bZzb ‡gwW‡Kj wWfvBm †iwR‡÷k‡bi Rb¨ `wLjKZ.03(wZb)wU †gwW‡Kj wWfvBm mfvq Dc¯vcb Kiv n‡j m`m¨MY KZ„K wWfvBm¸wji Safety, Efficacy and Performance gj¨vqb K‡i বিস্তারিত আলোচনাক্রমে wb¤wjwLZ wm×vš Mp„xZ nq t

K) JMI Alcohol prer pads bvgxq ‡gwW‡Kj wWfvBmwU Abyôgv`b Kiv nq Ges
L) JMI Sterile Disposable Scalpel Ges JMI Sononned (Atraumitic Needle for peripheral nerve block) bvgxq ‡gwW‡Kj wWfvBm `ÿwU wb¤wjwLZ kZmv‡c‡ÿ Abyôgv`b Kiv nqt
(1) ‡iwR‡ók‡bi c‡e †mdwU WvUv `wLj Ki‡Z n‡e Ges
(2) †iwR‡÷k‡bi c‡e e‡qU KZ„K mwUwd‡KU MnY Ki‡Z n‡e; (Annex-D)|

3.2 **Avg`vbxi Rb¨ Av‡ew`Z 119 wU ‡gwW‡Kj wWfvB‡mi †iwR‡÷k‡bi Abyôgv`‡bi mycwikmg~ni wel‡q Av‡jvPbv I wm×vš MnY cm‡½|**

Avg`vbxi Rb¨ Av‡ew`Z 119(GKkZ Dwbk)wU †gwW‡Kj wWfvBm mfvq Dc¯vcb Kiv n‡j Dcw¯Z m`m¨MY KZ„K cwZwU ‡gwW‡Kj wWfvB‡mi wbivcËv, KvhKvixZv I e¨envi Dc‡hwMZvi (Safety, Efficacy and Usefulness) Av‡jv‡K we¯wiZ Av‡jvPbvcwK wb¤wjwLZ wm×vš Mp„xZ nq t
K) Abyôgw`Z - 114(GKkZ †PŠÏ)wU,
L) ¯wMZ I ‡mdwU WvU `wLj Kiv‡i Rb¨ ejv nq-01wU Ges

M) ‡iwR‡ókb MvBWjvBb Abyjvqx c‡qvRbxq WK‡gv›U `wLj Ki‡Z ejv nq-4wU; (Annex-E)|

3.3 Avg`vbxi Rb¨ Av‡ewkó `Z 8 wU AvBwfwW wi‡q‡R›U-Gi †iwR‡÷k‡bi Abyjgv`‡bi mgvwikmg~n wel‡q Av‡jvPbv I wm×vš— MnY cm‡½|

Avg`vbxi Rb¨ Av‡ewkó`Z 08(AvU)wU AvB,wf,wW, wi‡q‡R‡›Ui Av‡e`b mfvq Dc¯vcb Kiv n‡j Dcw¯Z m`m¨MY KZ©K cwZwU AvB,wf,wW, wi‡q‡R‡›Ui wbivcËv, KvhKvixZv I e¨envi Dc‡hvMZvi (Safety, Efficacy and usefullness) Av‡jv‡K we¯wiZ Av‡jvPbvcwK wb¤wjwLZ wm×vš— Mp,xZ nq t
K) Abyjgvw`Z-05(cuP) wU Ges
L) ‡iwR‡ókb MvBWjvBb Abyjvqx c‡qvRbxq WK‡gv›U `wLj Ki‡Z ejv nq-03wU; (Annex-F)|

4| nvevj GWfvBRix KwgwUi (Jla wbqšY KwgwUi †UKwbK¨vj mve KwgwU) MZ 09-11-2016 Zwi‡L Abyôv‡Z mfvi mgvwikmg~n t

4.1 nvevj Jl‡ai Rb¨ c‡e wbawiZ 16 (‡lvj) wU eB Gi cvkvcwk †idv‡iÝ eB wnmv‡e USP-DSC, Indonesian Herbal Pharmacopoeia, Vietnam Herbal Pharmacopoeia & North Korean Herbal Pharmacopoeia-‡K Aš—f~wgKiY cm‡½|

<u>mfvi Av‡jvPbv I wm×vš— t</u>

mfvq m`m¨ mwPe e‡jb †h, nvevj Jla GWfvBRix KwgwU Gi 13(‡Zi)Zg mfvq 16(‡lvj)wU Reference eB wnmv‡e wbavi Y Kiv nq| D³ 16(‡lvj)wU Reference eB Gi cvkvcwk USP-DSC, Indonesian Herbal Pharmacopoeia, Vietnam Herbal Pharmacopoeia & North Korean Herbal Pharmacopoeia ‡KI Reference eB wnmv‡e ms‡hvRb Kwievi Rb¨ cwZwbwa, miKvwi BDbvbx I Avqv‡e`K †gvW‡Kj K‡jR I nvmcvZvj Ges cwZwbwa, evsjv‡`k nvevj †cvWv± g¨vbvd¨vKPvivm G‡mvwm‡qkb c¯ve K‡ib| D³ c¯v‡ei Dci m`m¨MY we¯wiZ Av‡jvPbv K‡i nvevj Jl‡ai Rb¨ c‡e wbawiZ 16(‡lvj)wU eB Gi cvkvcwk USP-DSC, Indonesian Herbal Pharmacopoeia, Vietnam Herbal Pharmacopoeia & North Korean Herbal Pharmacopoeia ‡KI Reference eB wnmv‡e wbewPZ Kivi Rb¨ GKgZ †cvlY K‡ib| D³ Reference eBmg~n nvevj Jla Drcv`b I gvb-wbqš‡bi Rb¨ Abyôv‡Z n‡e|

4.2 Lmov nvevj Jl‡ai wbeÜb bwZgvjv ms‡kvwaZ AvKv‡i Abyjgv`‡bi mgvwi‡ki wel‡q wm×vš— MnY cm‡½|

<u>mfvi Av‡jvPbv I wm×vš— t</u>

Lmov nvevj Jl‡ai wbeÜb bwZgvjv Abyjgv`‡bi wbg‡Ë ¯v¯¨ I cwievi Kj¨vY gš¿Yvj‡q †ciY Ki‡Z n‡e|

4.3 ¯vbxqfv‡e Drcv`‡bi Rb¨ Av‡e`Z 29 wU nvevj Jl‡ai †iwR‡÷k‡bi Abyjgv`‡bi mgvwikmg~ni wel‡q Av‡jvPbv I wm×vš MnY cm‡½|

mfvi Av‡jvPbv I wm×vš t

bZzb 29 (Dbœ‡Îk)wU nvevj Jl‡ai †iwR‡ókb Abyjgv`‡b wel‡q mfvq Dc¯vcb Kiv n‡j Dcw¯Z m`m¨MY KZK we¯vরিত আলোচানক্রমে wbœwjwLZ wm×vš Mp,xZ nq t

K) Abyjgvw`Z - 14 (‡PŠÏ) wU Ges
*L) bvgÄyKZ.- 15 (c‡bi) wU; (*Annex-G*)|*

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

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

‡gvt wmivRyi Bmjvg
mwPe
¯v¯¨ I cwievi Kj¨vY gšYvjq
I
mfvcwZ
Jla wbqšY KwgwU|

Annex-A: Proposed Product for locally manufacture (Human)

bs	c¯ZKvi‡K bvg	JI‡ai bvg	‡R‡bwiK bvg	‡_iwcDwUK Kvm	wb‡`kbv	Contra-indication & Side-effect	Status (New Molecule/ Existing)	Av‡e`bKvix KZK USFDA/BNF / MHRA Ref.	‡UKwbK¨vj mve KwgwUi mfvi wm×vš	mfvi wm×vš
1.	Mundipharma (Bangladesh) Private Limited. Mirzapur,JL#08, Gazipur Sadar, Gazipur, Bangladesh	Acbrophylline INN 100 mg Capsules	Acbrophylline INN 100 mg Capsules	Antiasthmatic	Acebrophylline is indicated in the symptomatic treatment of acute and chronic disease of respiratory system characterized by mucus hyper-secretion including acute bronchitis, chronic bronchitis, chronic obstructive pulmonary disease, asthma like bronchitis and pulmonary emphysema. Acebrophylline is administered at a recommended dose of 1 tablet once daily.	**Contraindications:** Acebrophylline is contraindicated in patients with hypersensitivity to xanthine derivatives like Theophylline or to ambroxol. **Side effect:** Commonly reported adverse effects with Acebrophylline include abdominal discomfort, stomach/abdominal distension, vomiting, abdominal pain, diarrhea, constipation, heart burn, loss of appetite, esophageal bleeding, rashes, urticaria, itching, drowsiness, difficulty in breathing, leukocytosis and nasal inflammation. If chills and fever occur the drug should be immediately discontinued.	New		c‡qvRbxq ‡id‡iÝ Ges Therapeutic benefit D‡jLceK Av‡iv Z_¨vw` `wLj Kivi Rb¨ ejv †h‡Z cv‡i	c‡qvRbxq ‡id‡iÝ Ges Therapeutic benefit D‡jLceK Av‡iv Z_¨vw` `wLj Kivi Rb¨ ejv nj	
2.	Mundipharma (Bangladesh) Private Limited. Mirzapur,JL#08, Gazipur Sadar, Gazipur, Bangladesh	Acbrophylline INN 200 mg CR tablets	Acbrophylline INN 200 mg tablets	Antiasthmatic	Acebrophylline is indicated in the symptomatic treatment of acute and chronic disease of respiratory system characterized by mucus hyper-secretion including acute bronchitis, chronic bronchitis, chronic obstructive pulmonary disease, asthma like bronchitis and pulmonary emphysema. Acebrophylline is administered at a recommended dose of 1 tablet once daily.	**Contraindications:** Acebrophylline is contraindicated in patients with hypersensitivity to xanthine derivatives like Theophylline or to ambroxol. **Side effect:** Commonly reported adverse effects with Acebrophylline include abdominal discomfort, stomach/abdominal distension, vomiting, abdominal pain, diarrhea, constipation, heart burn, loss of appetite, esophageal bleeding, rashes, urticaria, itching, drowsiness, difficulty in breathing, leukocytosis and nasal inflammation. If chills and fever occur the drug should be immediately discontinued.	New		c‡qvRbxq ‡id‡iÝ Ges Therapeutic benefit D‡jLceK Av‡iv Z_¨vw` `wLj Kivi Rb¨ ejv †h‡Z cv‡i	c‡qvRbxq ‡id‡iÝ Ges Therapeutic benefit D‡jLceK Av‡iv Z_¨vw` `wLj Kivi Rb¨ ejv nj	

নং	প্রস্তুতকারকের নাম	ঔষধের নাম	জেনেরিক নাম	‡_iwcDwUK Kvm	নির্দেশনা	Contra-indication & Side-effect	Status (New Molecule/ Existing)	আবেদনকারী প্রদত্ত USFDA/BNF/ MHRA Ref.	‡UKwbK¨vj mve KwgwUi mfvi wm×vš	mfvi wm×vš
3.	Mundipharma (Bangladesh) Private Limited. Mirzapur,JL#08, Gazipur Sadar, Gazipur, Bangladesh	Doxophylline 400mg CR Tablet	Doxophylline INN 400 mg	Antiasthmatic	For the treatment of Chronic Obstructive Pulmonary Disease (COPD), bronchial asthma and pulmonary disease with spastic bronchial component. The initial day dosage for adult is 1 tablet, two to three times in a day. For Elderly Patients the daily dosage half of the tablets, two to three times in a day or as prescribed by the Physician	**Contraindications:** Hypersensitivity to any ingredient in the product or to other methylxanthines. Patients with acute myocardial infarction, hypotension, and in breastfeeding women. **Side effects:** After xanthine administration, nausea, vomiting, epigastric pain, cephalalgia, irritability, insomnia, tachycardia, extrasystole, tachypnea, and occasionally hyperglycemia and albuminuria, may occur. If a potential oral overdose is established, the patient may present with severe arrhythmias and seizure; these symptoms could be the first sign of intoxication. Adverse reactions may cause the withdrawal from treatment; a lower dose rechallenge may start only after the advice of physician.	200mg, 400mg Tablet; 100 mg/5 ml Syrup		Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
4.	Navana Pharmaceuticals Ltd.	Seratrodast 40 mg Tablet	Seratrodast INN 40 mg	Antiasthmatic agent	Is indicated in adults (18 years and above) for the prophylactic management of asthma. It also effective in the treatment of allergic rhinitis and COPD.	**Contraindications:** Hypersensitivity to seratrodast or any of the ingredients. Patients with hepatic failure. **Side effects:** Adverse reactions are generally rare, but if present, range from mild to moderate in severity. Generally, rash, inching (hypersensitivity); elevated liver enzymes (hepatic); nausea, loss of appetite, stomach discomfort, abdominal pain, diarrhea, dry mout, anemia, drowsiness, headache,and palpitations, malaise observed	New		c‡qvRbxq ‡id‡i Ý Ges Therapeutic benifit D‡jLce‡K Av‡iv Z_¨vw` `wLj Kiv‡i Rb¨ ejv †h‡Z cv‡i	c‡qvRbxq ‡id‡i Ý Ges Therapeutic benifit D‡jLce‡K Av‡iv Z_¨vw` `wLj Kiv‡i Rb¨ ejv nj

নং	প্রস্তুতকারকের নাম	ঔষধের নাম	জেনেরিক নাম	‡_iwcDwUK Kvm	নির্দেশনা	Contra-indication & Side-effect	Status (New Molecule/ Existing)	আবেদনকারী প্রদত্ত USFDA/BNF/ MHRA Ref.	‡UKwbK¨vj mve Kvgwui mfvi wm×vš	mfvi wm×vš
5.	Navana Pharmaceuticals Ltd.	Seratrodast 80 mg Tablet	Seratrodast INN 80 mg	Antiasthmatic agent	Is indicated in adults (18 years and above) for the prophylactic management of asthma. It also effective in the treatment of allergic rhinitis and COPD.	**Contra-indications:** Hypersensitivity to seratrodast or any of the ingredients. Patients with hepatic failure. **Side effects:** Adverse reactions are generally rare, but if present, range form mild to moderate in severity. Generally, rash, inching (hypersensitivity); elevated liver enzymes (hepatic); nausea, loss of appetite, stomach discomfort, abdominal pain, diarrhea, dry mout, anemia, drowsiness, headache,and palpitations, malaise observed	New		c‡qvRbxq ‡id v‡iÝ Ges Therapeutic benifit D‡jLceK Av‡iv Z_¨vw` `wLj Kivi Rb¨ ejv †h‡Z cv‡i|	c‡qvRbxq ‡id v‡iÝ Ges Therapeutic benifit D‡jLceK Av‡iv Z_¨vw` `wLj Kivi Rb¨ ejv nj|
6.	**Acme Laboratories Ltd., Dhamrai, Dhaka**	Ceftaroline Fosamil 400 mg/Vial Powder for IV Injection	Ceftaroline Fosamil monoacetate monohydrate with L-Arginine (Sterile) INN 742.500 mg eq. to Ceftaroline Fosamil 400mg/Vial	Antibiotic	*Acute Bacterial Skin And Skin Structure Infections* Ceftaroline Fosamil 400 mg IV Injection is indicated for the treatment of acute bacterial skin and skin structure infections (ABSSSI) caused by susceptible isolates of the following Gram-positive and Gram-negative microorganisms: Staphylococcus aureus (including methicillin-susceptible and -resistant isolates), Streptococcus pyogenes, Streptococcus agalactiae, Escherichia coli, Klebsiella pneumoniae, and Klebsiella oxytoca. *Community-Acquired Bacterial Pneumonia* Ceftaroline Fosamil 400 mg IV Injection is indicated for the treatment of community-acquired bacterial pneumonia (CABP) caused by susceptible isolates of the following Gram-positive and Gram-negative microorganisms: Streptococcus pneumoniae (including cases with concurrent bacteremia), Staphylococcus aureus(methicillin-susceptible isolates only), Haemophilus influenzae, Klebsiella pneumoniae, Klebsiella oxytoca, and Escherichia coli.	**Contraindication:** Ceftaroline Fosamil is contraindicated in patients with known serious hypersensitivity to Ceftaroline or other members of the cephalosporin class. Anaphylaxis and anaphylactoid reactions have been reported with Ceftaroline. **Side effects**: Local injection site reactions, rash, pruritus, urticaria, abdominal pain, nausea, vomiting, diarrhoea, pseudomembranous colitis. Rarely Hypersensitivity Reactions, Clostridium difficile-Associated diarrhea, Direct Coombs' Test Seroconversion,	**New**	USFDA	Abyjgv`b Kiv †h‡Z cv‡i|	Abyjgv`b Kiv nj|

নং	প্রস্তুতকারকের নাম	ঔষধের নাম	জেনেরিক নাম	‡_iwcDwUK Kvm	নির্দেশনা	Contra-indication & Side-effect	Status (New Molecule/ Existing)	আবেদনকারী প্রদত্ত USFDA/BNF/ MHRA Ref.	‡UKwbK¨vj mve Kwgwdi mfvi wm×vš	mfvi wm×vš
7.	**Acme Laboratories Ltd., Dhamrai, Dhaka**	Ceftaroline Fosamil 600 mg/Vial Powder for IV Injection	Ceftaroline Fosamil monoacetate monohydrate with L-Arginine (Sterile) INN 1114 mg eq. to Ceftaroline Fosamil 600mg/Vial	Antibiotic	*Acute Bacterial Skin And Skin Structure Infections* Ceftaroline Fosamil 600 mg IV Injection is indicated for the treatment of acute bacterial skin and skin structure infections (ABSSSI) caused by susceptible isolates of the following Gram-positive and Gram-negative microorganisms: Staphylococcus aureus (including methicillin-susceptible and -resistant isolates), Streptococcus pyogenes, Streptococcus agalactiae, Escherichia coli, Klebsiella pneumoniae, and Klebsiella oxytoca. *Community-Acquired Bacterial Pneumonia* Ceftaroline Fosamil 600 mg IV Injection is indicated for the treatment of community-acquired bacterial pneumonia (CABP) caused by susceptible isolates of the following Gram-positive and Gram-negative microorganisms: Streptococcus pneumoniae (including cases with concurrent bacteremia), Staphylococcus aureus(methicillin-susceptible isolates only), Haemophilus influenzae, Klebsiella pneumoniae, Klebsiella oxytoca, and Escherichia coli.	**Contraindication:** Ceftaroline Fosamil is contraindicated in patients with known serious hypersensitivity to Ceftaroline or other members of the cephalosporin class. Anaphylaxis and anaphylactoid reactions have been reported with Ceftaroline. **Side effects**: Local injection site reactions, rash, pruritus, urticaria, abdominal pain, nausea, vomiting, diarrhoea, pseudomembranous colitis. Rarely Hypersensitivity Reactions ,Clostridium difficile-Associated diarrhea, Direct Coombs' Test Seroconversion,	**New**	USFDA	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

নং	প্রস্তুতকারকের নাম	ঔষধের নাম	জেনেরিক নাম	‡_iwcDwUK Kvm	নির্দেশনা	Contra-indication & Side-effect	Status (New Molecule/ Existing)	আবেদনকারী প্রদত্ত USFDA/BNF/ MHRA Ref.	‡UKwbK`vj mve Kwgwdi mfvi wm×vš	mfvi wm×vš
8.	Aristopharma Ltd, Shampur-Kadamtali I/A, Dhaka-1204	Moxifloxacin 0.50gm/100g Sterile Eye Ointment	Moxifloxacin HCl BP 0.5454gm eq. to Moxifloxacin 0.50gm/100gm	Antibiotic	Moxifloxacin is a broad-spectrum antibiotic that is active against both Gram-positive and Gram –negative bacteria. It functions by inhibiting DNA gyrase, a type II topoisomerase, and opoisomerase IV enzymes necessary to separate bacteria DNA, thereby inhibiting cell replication. It is indicated for the treatment of ocular bacteria infection.	**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 effects:** The most frequently reported Ocular adverse events were conjunctivitis, decreased visual acuity, dry eye, keratitis, ocular discomfort, ocular hyperemia, ocular pain, ocular pruritus, subconjunctival hemorrhage, and tearing.	400mg Tablet 0.5g/100 Eye Drop 0.16% IV Infusion	Vietnam, Myanmar, India, Sri Lanka.	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |
9.	Square Pharmaceuticals Ltd., (Dhaka Unit) Kaliakoir, Gazipur	Cefadroxil 1000mg Tablet	Cefadroxil Monohydrate USP 1049.50mg eq. to 1000mg Cefadroxil Tablet	Antibiotic	Indicated for the treatment of patients with infection caused by susceptible strains of the designated organisms in the following diseases: • Urinary tract infections caused by *E. coli*, *P. mirabilis*, and *Klebsiella* species. Skin • skin structure infections caused by staphylococci and/or streptococci. Pharyngitis and/or tonsillitis caused by *Streptococcus pyogenes* (Group A beta-hemolytic streptococci).	**Contraindication**: Contraindicated in patients with known allergy to thecephalosporin group of antibiotics. **Side effect:** Generally well tolerated. The most commonly reported side effects are gastrointestinal disturbances and hypersensitivity. Side effects including nausea, vomiting, 12diarrhea, dyspepsia, abdominal discomfort, fever, dizziness, headache, arthralgia may also occur.	250mg, 500mg Capsule 125 mg/5 ml PFS 250 mg/5 ml PFS	USFDA	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |

নং	প্রস্তুতকারকের নাম	ঔষধের নাম	জেনেরিক নাম	‡_iwcDwUK Kvm	নির্দেশনা	Contra-indication & Side-effect	Status (New Molecule/ Existing)	আবেদনকারী প্রদত্ত USFDA/BNF/ MHRA Ref.	‡UKwbK¨vj mve Kwgwdi mfvi wm×vš	mfvi wm×vš
10.	**Acme Laboratories Ltd., Dhamrai, Dhaka**	Ciprofloxacin 500 mg + Metronidazole 300 mg Tablet	Ciprofloxacin Hydrochloride USP 582.20 mg eq. to 500mg Ciprofloxacin + Metronidazole BP 300 mg	Antibiotic & Antiprotozoal	Ciprofloxacin and Metronidazole combination is indicated for the treatment of mild to moderate Diverticulitis. Ciprofloxacin is an antibiotic that works to kill bacteria and Metronidazole kills protozoans as well as bacterias, especially anaerobic bacteria. In diverticulitis, it is possible to have both bacteria and protozoans in the digestive tract. Diverticulitis can be classified as mild, moderate, or severe. Treatment is based on clinical findings and the results of imaging studies. The mainstay of treatment includes antibiotic therapy, bowel rest, and analgesia. In mild to moderate diverticulitis, localized symptoms are present without evidence of perforation, abscess, or significant comorbidity. Patients can be managed on an outpatient basis with close follow-up. Treatment also includes a clear liquid diet for 3-5 day. Severe diverticulitis may include focal or generalized peritonitis, peridiverticular abscess, and systemic signs of sepsis. Inpatient treatment is recommended; surgical intervention may be required.	**Contraindication:** Contraindicated to patients with known hypersensitivity to Ciprofloxacin and Metronidazole or any other components of the products. **Side effects**: Common side effects may include nausea, vomiting, loss of appetite, a metallic taste, diarrhea, dizziness, headaches, and discolored urine (dark or reddish brown). Another side effect of long-term use is tingling of the hands and feet, which may persist even after the drug is discontinued.	**Ciprofloxacin 250, 500 mg tablet & Metronidazole 200, 400 mg tablet**		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
11.	Aristopharma Ltd, Shampur-Kadamtali I/A, Dhaka-1204	Doxylamine Succinate 20 mg + Pyridoxine Hydrochloride 20mg Extended Release Tablet	Doxylamine Succinate USP 20 mg + Pyridoxine Hydrochloride BP 20 mg	Anticholinergic + Vitamin	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 nausea and vomiting of pregnancy in women who do not respond to conservative management.	**Contraindications:** Known hypersensitivity to doxylamine succinate, other ethanolamine derivative antihistamines, pyridoxine hydrochloride or any inactive ingredient in the formulation Monoamine oxidase (MAO) inhibitors. **Side effects:** The most common adverse reaction with this tablet (≥5 percent and exceeding the rate in placebo) is somnolence.	Doxylamine Succinate 10mg+ Pyridoxine HCl 10mg Delayed Release Tablet	USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

নং	প্রস্তুতকারকের নাম	ঔষধের নাম	জেনেরিক নাম	‡_iwcDwUK Kvm	নির্দেশনা	Contra-indication & Side-effect	Status (New Molecule/ Existing)	আবেদনকারী প্রদত্ত USFDA/BNF/ MHRA Ref.	‡UKwbK`vj mve KwgwUi mfvi wm×vš	mfvi wm×vš
12.	Aristopharma Ltd, Shampur-Kadamtali I/A, Dhaka-1204	Empagliflozin 25 mg Film Coated Tablet	Empagliflozin INN 25 mg	Antidiabetic	It is indicated as an adjunct to diet and exercise to improve glycemic control in adults with type 2 diabetes mellitus. **Limitation of use:** It is not recommended for patients with type 1 diabetes or for the treatment of diabetic ketoacidosis.	**Contraindications:** • History of serious hypersensitivity reaction to empagliflozin. • Severe renal impairment, end-stage renal disease, or dialysis. **Side effects:** The most common adverse reactions associated with empagliflozin were urinary tract infections and female genital mycotic infections.	New	USFDA	c‡qvRb ‡bB weavq Av‡e`b bvgÄyi Kiv †h‡Z cv‡i	c‡qvRb ‡bB weavq Av‡e`b bvgÄyi Kiv nj
13.	Popular Pharmaceuticals Limited	Gliclazide 80mg + Metformin Hydrochloride 500mg Film Coated Tablet	Gliclazide BP 80mg + Metformin Hydrochloride BP 500mg	Antidiabetic	It is indicated as anadjunct to diet and exercise to improve glycemic control in adults with type 2diabetes mellitus.	**Contraindications:** Type-1 diabetes mellitus, renal or hepatic failure, alcoholism, type-2 diabetes 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, known hypersensitivity to any of the ingredients. **Side effects:** **Gastrointestinal disturbances -** Nausea, diarrhea, gastric pain, constipation, vomiting, metallic taste in mouth. **Dermatological effects -** Rash, pruritus, urticaria, erythema and flushing. **Miscellaneous -** Headache and dizziness. Gliclazide appears to be associated with a low incidence of hypoglycaemia. Impaired gastrointestinal absorption of vitamin B_{12} and folic acid has been associated with long term Metformin therapy.			c‡qvRb ‡bB weavq Av‡e`b bvgÄyi Kiv †h‡Z cv‡i	c‡qvRb ‡bB weavq Av‡e`b bvgÄyi Kiv nj

নং	প্রস্তুতকারকের নাম	ঔষধের নাম	জেনেরিক নাম	‡_iwcDwUK Kvm	নির্দেশনা	Contra-indication & Side-effect	Status (New Molecule/ Existing)	আবেদনকারী প্রদত্ত USFDA/BNF/ MHRA Ref.	‡UKwbK¨vj mve KwgwUi mfvi wm×vš	mfvi wm×vš
14.	Square Formulations Ltd., Gorai, Tangail	Metformin HCl 1000mg + Canagliflozin 50mg Extended Release Tablet	Metformin Hydrochloride BP 1000mg + Canagliflozin INN 50mg	Antidiabetic	This 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 when treatment with both canagliflozin and metformin is appropriate	**Contraindications:** Moderate to severe renal impairment (eGFR below 45 mL/min/1.73 m^2), end stage renal disease or dialysis. Metabolic acidosis, including diabetic ketoacidosis History of serious hypersensitivity reaction to canagliflozin or metformin **Adverse Effect:** 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	Metformin Hydrochloride BP 1000mg Tablet Canagliflozin 100mg Tablet	USFDA	c‡qvRb ‡bB ïeavq Av‡e`b bvgÄyi Kiv †h‡Z cv‡i	c‡qvRb ‡bB ïeavq Av‡e`b bvgÄyi Kiv nj
15.	Square Pharmaceuticals Ltd., Pabna Unit, Salgaria, Pabna	Metformin Hydrochloride 1000mg + Canagliflozin 50mg Tablet	Metformin Hydrochloride BP 1000mg + Canagliflozin INN 50mg Tablet	Antidiabetic	This 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 when treatment with both canagliflozin and metformin is appropriate	**Contraindications:** Moderate to severe renal impairment (eGFR below 45 mL/min/1.73 m^2), end stage renal disease or dialysis. Metabolic acidosis, including diabetic ketoacidosis History of serious hypersensitivity reaction to canagliflozin or metformin **Adverse Effect:** 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	Metformin Hydrochloride BP 1000mg Tablet Canagliflozin 100mg Tablet	USFDA	c‡qvRb ‡bB ïeavq Av‡e`b bvgÄyi Kiv †h‡Z cv‡i	c‡qvRb ‡bB ïeavq Av‡e`b bvgÄyi Kiv nj

নং	প্রস্তুতকারকের নাম	ঔষধের নাম	জেনেরিক নাম	‡_iwcDwUK Kvm	নির্দেশনা	Contra-indication & Side-effect	Status (New Molecule/ Existing)	আবেদনকারী প্রদত্ত USFDA/BNF/ MHRA Ref.	‡UKwbK¨vj mve Kwgwdi mfvi wm×vš	mfvi wm×vš
16.	Square Pharmaceuticals Ltd., Pabna Unit, Salgaria, Pabna	Dronabinol 0.50gm/100ml Oral soluton	Dronabinol USP 0.5gm/100ml	Antiemetic	It is indicated in adults for the treatment of • chemotherapy induced nausea and vomiting associated with cancer chemotherapy in patients who have failed to respond adequately to conventional antiemetic treatments and • anorexia associated with weight loss in patients with Acquired Immune Deficiency Syndrome (AIDS).	**Contraindications:** It is contraindicated in patients • with a sensitivity to dronabinol or alcohol • with a history of hypersensitivity of reaction to alcohol • who are receiving disulfiram or metronidazole in past 14 days.	New	USFDA	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 |
17.	Navana Pharmaceuticals Ltd.	Clonazepam 0.25mg Orally Disintegrating Tablet	Clonazepam 0.25 mg	Antiepileptic	Seizure Disorder & Panic Disorder	**Contra-indication** : Liver disease & Narrow angle glaucoma **Side Effects** : Amnesia . bronchial hypersecretion in infants and small children . coordination disturbances. Confusion. dependence .dizziness . drowsiness .fatigue. muscle hypotonia	0.5mg, 1 mg, 2mg Tablet & 1mg/ml, 2.5mg/ml pediatric drops	USFDA (Discontinued)	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 |
18.	Navana Pharmaceuticals Ltd.	Clonazepam 0.5 mg Orally Disintegrating Tablet	Clonazepam 0.5 mg	Antiepileptic	Seizure Disorder & Panic Disorder	Do	0.5mg, 1 mg, 2mg Tablet & 1mg/ml, 2.5mg/ml pediatric drops	USFDA	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.	Navana Pharmaceuticals Ltd.	Clonazepam 1 mg Orally Disintegrating Tablet	Clonazepam 1 mg	Antiepileptic	Seizure Disorder & Panic Disorder	Do	0.5mg, 1 mg, 2mg Tablet & 1mg/ml, 2.5mg/ml pediatric drops	USFDA	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.	Navana Pharmaceuticals Ltd.	Clonazepam 2 mg Orally Disintegrating Tablet	Clonazepam 2 mg	Antiepileptic	Seizure Disorder & Panic Disorder	Do	0.5mg, 1 mg, 2mg Tablet & 1mg/ml, 2.5mg/ml pediatric drops	USFDA	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 |

নং	প্রস্তুতকারকের নাম	ঔষধের নাম	জেনেরিক নাম	‡_iwcDwUK Kvm	নির্দেশনা	Contra-indication & Side-effect	Status (New Molecule/ Existing)	আবেদনকারী প্রদত্ত USFDA/BNF/ MHRA Ref.	‡UKwbK¨vj mve KwgwUi mfvi wm×vš	mfvi wm×vš
21.	Square Pharmaceuticals Ltd., Pabna Unit, Salgaria, Pabna	Perampanel 2mg Tablet	Perampanel INN 2mg	Antiepileptic	This is a non-competitive AMPA glutamate receptor antagonist which is indicated as adjunctive therapy for the treatment of: • Partial-Onset Seizures with or without secondarily generalized seizures in patients with epilepsy 12 years of age and older. • Primary Generalized Tonic-Clonic Seizures in patients with epilepsy 12 years of age and older.	**Contraindications:** None **Advarse Effect** • Neurologic Effects: Monitor for dizziness, gait disturbance, somnolence, and fatigue Patients should use caution when driving or operating machinery. • Falls: Monitor for falls and injuries • Withdrawal of Antiepileptic Drugs: In patients with epilepsy, there may be an increase in seizure frequency Most common adverse reactions (≥5% and ≥1% higher than placebo) include dizziness, somnolence, fatigue, irritability, falls, nausea, weight gain, vertigo, ataxia, headache, vomiting, contusion, abdominal pain, and anxiety.	New	USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
22.	Square Pharmaceuticals Ltd., Pabna Unit, Salgaria, Pabna	Perampanel 4mg Tablet	Perampanel INN 4mg	Antiepileptic	-Do-	-Do-	New	USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
23.	Navana Pharmaceuticals Ltd.	Naftifine HCl 2% Cream	Naftifine HCl 2%	Antifungal	It is an allylamine antifungal indicated for the treatment of interdigital tinea pedis, tinea cruris, and tinea corporis caused by the organism Trichophyton rubrum.	**Contraindications:** None **Side Effects:** The most common adverse reaction (≥1%) is pruritus.	New	USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
24.	Square Pharmaceuticals Ltd., (Dhaka Unit) Kaliakoir, Gazipur	Bimatoprost 0.3mg + Timolol 5mg/ml Opthalmic Solution	Bimatoprost INN 0.3mg + Timolol maleate BP 6.834mg eq. to 5.0mg Timolol/ml	Antiglucoma	It is indicated for the raised intra-ocular pressure in patient with open angle glaucoma or ocular hypertension when beta-blocker or prostaglandin analogue alone not adequate.	**Contraindication**: none **Side effect:** Blepharitis, blood pressure changes, brown pigmentation, conjuctival disorder, corneal erosion, headache and ocular discomfort	Bimatoprost 0.03% Eye Drops	BNF-72 (Page-1029)	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

নং	প্রস্তুতকারকের নাম	ঔষধের নাম	জেনেরিক নাম	‡_iwcDwUK Kvm	নির্দেশনা	Contra-indication & Side-effect	Status (New Molecule/ Existing)	আবেদনকারী প্রদত্ত USFDA/BNF/ MHRA Ref.	‡UKwbK¨vj mve Kwgwdi mfvi wm×vš	mfvi wm×vš
25.	**Acme Laboratories Ltd., Dhamrai, Dhaka**	Levocetirizine Dihydrochloride 5mg + Montelukast 10 mg Tablet	Levocetirizine Dihydrochloride USP 5 mg + Montelukast Sodium USP 10.38 mg eq. to Montelukast 10 mg	Antihistamine & Anti-Asthmatic	The combination is indicated for chronic allergic conditions like seasonal allergic rhinitis, perennial allergic rhinitis, Rhinitis associated with Asthma.	**Contraindication:** The combination is contraindicated in patients known to have hypersensitivity to the drug or any of its components. Patients with completely impaired renal function (anuria). **Side effects:** Possible side effects are asthenia/ fatigue, fever, abdominal pains, trauma, dyspepsia, infectious gastroenteritis, dental pain, dizziness, headache, cough, nasal congestion, influenza, rash. Somnolence, dry mouth, hypersensitivity, convulsion, visual disturbances, palpitation, dyspnea, nausea, hepatitis, angioneurotic edema, pruritus, urticaria, myalgia, weight gain, abnormal liver function test.	Levocetirizine Tablet 5 mg & Montelukast Tablet 10 mg		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 |
26.	Healthcare Pharmaceutical Ltd., Rajendrapur,Gazipur	Sacubitril 24 mg + Valsartan 26mg Tablet	Sacubitril Sodium INN 25.28 mg eq. to Sacubitril INN 24 mg) + Valsartan USP 26 mg	Antihypertensive	It is indicated for the treatment of heart failure with reduced ejection fraction in patients with NYHA class II or III, to reduce the incidence of cardiovascular death & heart failure hospitalization	**Contraindications:** Recent symptomatic hypotension prior to initiation of treatment with Sacubitril+ Valsartan. **Side-Effects:** Low blood pressure, Excessively high level of potassium in the blood, as shown in a blood test, Dizziness, Tiredness, sudden loss of consciousness ,weakness & rash	Sacubitril 49mg + Valsartan 51mg Tablet Sacubitril 97mg + Valsartan 103mg Tablet	USFDA	Abyjvg`b Kiv †h‡Z cv‡i |	Abyjvg`b Kiv nj |
27.	Nuvista Pharma Ltd.	Selexipag 0.2mg Tablet.	Selexipag INN 0.20 mg	Antihypertensive	Pulmonary Arterial Hypertension (PAH): For the treatment of pulmonary arterial hypertension (PAH, WHO Group I) to delay disease progression and reduce the risk of hospitalization for PAH. Effectiveness was established in a long-term study in PAH patients with WHO Functional Class II-III symptoms. Patients had idiopathic and heritable PAH (58%), PAH associated with connective tissue disease (29%), PAH associated with congenital heart disease with repaired shunts (10%).	**Contraindications:**None; **Side-effects:** Headache, Diarrhea, jaw pain, nausea, muscle pain, vomiting, pain in arms or legs, pain in joints, decreased appetite, flushing, low red blood cell count, rash, hypohemoglobinemia, a reduction in median thyroid-stimulating hormone (TSH)	New	USFDA	Abyjvg`b Kiv †h‡Z cv‡i |	Abyjvg`b Kiv nj |

নং	প্রস্তুতকারকের নাম	ঔষধের নাম	জেনেরিক নাম	‡_iwcDwUK Kvm	নির্দেশনা	Contra-indication & Side-effect	Status (New Molecule/ Existing)	আবেদনকারী প্রদত্ত USFDA/BNF/ MHRA Ref.	‡UKwbK`vj mve KwgwUi mfvi wm×vš	mfvi wm×vš
28.	Nuvista Pharma Ltd.	Selexipag 0.4 mg Tablet.	Selexipag INN 0.40 mg	Antihypertensive	Pulmonary Arterial Hypertension (PAH): For the treatment of pulmonary arterial hypertension (PAH, WHO Group I) to delay disease progression and reduce the risk of hospitalization for PAH. Effectiveness was established in a long-term study in PAH patients with WHO Functional Class II-III symptoms. Patients had idiopathic and heritable PAH (58%), PAH associated with connective tissue disease (29%), PAH associated with congenital heart disease with repaired shunts (10%).	**Contraindications:**None; **Side-effects:** Headache, Diarrhea, jaw pain, nausea, muscle pain, vomiting, pain in arms or legs, pain in joints, decreased appetite, flushing, low red blood cell count, rash, hypohemoglobinemia, a reduction in median thyroid-stimulating hormone (TSH)	New Molecule	USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
29.	NIPRO JMI Pharma Ltd.	Diacerein 50 mg Tablet	Diacerein INN 50 mg	Antinflammatory	Symptomatic relief in long-term treatment of osteoarthritis, anti-inflammatory agent, prescribed for chronic inflammatory arthritis.	**Contraindications:** Contraindicated in those patients with known hypersensitivity to the drug itself or to those with previous episodes of hypersensitivity to anthraquinone derivatives. **Side effects:** Diarrhea, Stomach pain, Nausea, Vomiting, Intense yellow coloring of urine, Epigastric pain	New		c‡qvRbxq ‡idv‡iÝ Ges Therapeutic benefit D‡jLceK Av‡iv Z_¨vw` `wLj Kivi Rb¨ ejv †h‡Z cv‡i	c‡qvRbxq ‡idv‡iÝ Ges Therapeutic benifit D‡jLceK Av‡iv Z_¨vw` `wLj Kivi Rb¨ ejv nBj
30.	Navana Pharmaceuticals Ltd.	Aspirin 325mg + Omeprazole 40mg Delayed Release Tablet	Aspirin 325 mg + Omeprazole 40 mg	Antiplatelet agent + PPI	Do	Do	Aspirin 75mg Tablet Omeprazole 40mg Capsule	USFDA	c‡qvRb ‡bB wewq Av‡e`b bvgÄyi Kiv †h‡Z cv‡i	c‡qvRb ‡bB wewq Av‡e`b bvgÄyi Kiv nj

নং	প্রস্তুতকারকের নাম	ঔষধের নাম	জেনেরিক নাম	‡_iwcDwUK Kvm	নির্দেশনা	Contra-indication & Side-effect	Status (New Molecule/ Existing)	আবেদনকারী প্রদত্ত USFDA/BNF/ MHRA Ref.	‡UKwbK`vj mve KwgwUi mfvi wm×vš	mfvi wm×vš
31.	a) Navana Pharmaceuticals Ltd. b) Delta Pharma Ltd.	Aspirin 81 mg + Omeprazole 40 mg Delayed Release Ta blet	Aspirin 81 mg + Omeprazole 40 mg	Antiplatelet agent + PPI	is a combination of aspirin, an anti-platelet agent, and omeprazole, a proton pump inhibitor (PPI), indicated for patients who require aspirin for secondary prevention of cardiovascular and cerebrovascular events and who are at risk of developing aspirin associated gastric ulcers. The aspirin component of YOSPRALA is indicated for reducing the combined risk of death and nonfatal stroke in patients who have had ischemic stroke or transient ischemia of the brain due to fibrin platelet emboli, reducing the combined risk of death and nonfatal MI in patients with a previous MI or unstable angina pectoris, reducing the combined risk of MI and sudden death in patients with chronic stable angina pectoris, use in patients who have undergone revascularization procedures (Coronary Artery Bypass Graft [CABG] or Percutaneous Transluminal Coronary Angioplasty [PTCA]) when there is a pre-existing condition for which aspirin is already indicated The omeprazole component of YOSPRALA is indicated for decreasing the risk of developing aspirin associated gastric ulcers in patients at risk for developing aspirin associated gastric ulcers due to age (≥55) or documented history of gastric ulcers **Limitations of Use**: Not for use as the initial dose of aspirin therapy during onset of acute coronary syndrome, acute myocardial infarction or before percutaneous coronary intervention.Has not been shown to reduce the risk of gastrointestinal bleeding due to aspirint is not interchangeable with the individual components of aspirin and	**Contraindications:** Patients with known allergy to aspirin and other nonsteroidal anti-inflammatory drug Products (NSAIDs) and in patients with the syndrome of asthma, rhinitis, and nasal polyps. Aspirin may cause severe urticaria, angioedema, or bronchospasm (asthma), Pediatric patients with suspected viral infections, with or without fever, because of the risk of Reye's syndrome with concomitant use of aspirin in certain viral illnesses. It contraindicated in patients with known hypersensitivity to aspirin, omeprazole, substituted benzimidazoles, or to any of the excipients in the formulation Proton pump inhibitor (PPI)–containing products , including (Aspirin + Omeprazole) contraindicated in patients receiving rilpivirine - containing products. **Side effects:** Most common adverse reactions in adults (≥2%) are: gastritis, nausea, diarrhea, gastric polyps, and non-cardiac chest pain.	Aspirin 75mg Tablet Omeprazole 40mg Capsule	USFDA	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 |

নং	প্রস্তুতকারকের নাম	ঔষধের নাম	জেনেরিক নাম	‡_iwcDwUK Kvm	নির্দেশনা	Contra-indication & Side-effect	Status (New Molecule/ Existing)	আবেদনকারী প্রদত্ত USFDA/BNF/ MHRA Ref.	‡UKwbK¨vj mve Kwgwdi mfvi wm×vš	mfvi wm×vš
32.	Square Pharmaceuticals Ltd., Pabna Unit, Salgaria, Pabna	Eflornithine 11.5gm/100gm Cream	Eflornithine Hydrochloride INN 13.9gm eq. to Eflornithine 11.5gm/100gm	Antiprotozoal	It's inhibits the enzyme ornithine decarboxylase in hair follicles and is use to reduce unwanted facial hair growth in women	**Contraindication**: Contraindicated in patients with a history of sensitivity to any components of the preparation. **Adverse Reactions:** The most frequent adverse events related to treatment with VANIQA® were skin-related. The following table notes the percentage of adverse events associated with the use of VANIQA® or its vehicle that occurred at greater than 1% in both the vehicle-controlled studies and the open-label safety studies up to 1 year of continuous use.	New	BNF-72 (Page 1118)	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
33.	Aristopharma Ltd, Shampur-Kadamtali I/A, Dhaka-1204	Levosulpiride 25mg Tablet	Levosulpiride INN 25mg	Antipsychotic	It is a gastroprokinetic agent.. For the treatment of gastroesophageal reflux disease, various forms of dyspepsia, diabetic gastroparesis, vomiting and nausea.	**Contraindications:** Levosulpiride is contraindicated in conditions like epilepsy, hyperprolactinaemia, breast feeding, and hypersensitivity to any component of product,gastrointestinal Hemorrhage and pheochromocytoma. **Side effects**: The symptomatic adverse reactions produced by Levosulpiride are more or less tolerable and if they become severe, they can be treated symptomatically, these include sedation, hypotension, and dyskinesia hyperprolactinemia .	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

নং	প্রস্তুতকারকের নাম	ঔষধের নাম	জেনেরিক নাম	‡_iwcDwUK Kvm	নির্দেশনা	Contra-indication & Side-effect	Status (New Molecule/ Existing)	আবেদনকারী প্রদত্ত USFDA/BNF/ MHRA Ref.	‡UKwbK¨vj mve Kwgwdi mfvi wm×vš	mfvi wm×vš
34.	Popular Pharmaceuticals Limited	Esomeprazole 2.50mg/Sachet Dealyed Release Granules	Esomeprazole Magnesium Trihydrate BP 4.17mg eq. t Esomeprazole 2.50mg	Antiulcerant	**Treatment of Gastroesophageal Reflux Disease (GERD)** ***Healing of Erosive Esophagitis*** Esomeprazole is indicated for the short-term treatment (4 to 8 weeks) in the healing and symptomatic resolution of diagnostically confirmed erosive esophagitis. For those patients who have not healed after 4 to 8 weeks of treatment, an additional 4 to 8 week course of Esomeprazole may be considered. In infants 1 month to less than 1 year, Esomeprazole is indicated for short-term treatment (up to 6 weeks) of erosive esophagitis due to acid-mediated GERD.	**Contraindications:** Esomeprazole is contraindicated in patients with known hypersensitivity to substituted benzimidazoles or to any component of the formulation. Hypersensitivity reactions may include anaphylaxis, anaphylactic shock, angioedema, bronchospasm, acute interstitial nephritis, and urticaria. For information about contraindications of antiba agents (clarithromycin and amoxicillin) indicated i Combination with Esomeprazole, **Side effects:** In general, Esomeprazole is well tolerated in both short and long-term use. Common side effects are headache, diarrhea. Other effects include nausea, flatulence, abdominal pain, constipation and dry mouth.	20mg & 40mg Capsule 20mg/Sachet	USFDA	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
35.	Navana Pharmaceuticals Ltd.	Doxoyphylline 400mg Sustained Release Tablet	Doxophylline INN 400 mg	Bronchodilator	Treatment of bronchial asthma and pulmonary disease with spastic bronchial component	**Contraindications:** Individuals who have shown hypersensitivity to Doxofylline and its components. Patients with acute myocardial infarction and hypotension. **Side Effects:** After xanthine administration, nausea, vomiting, epigastric pain, cephalalgia, irritability, insomnia, tachycardia, extrasystole, tachypnea and occasionally, hyperglycemia and albuminuria, may occur. If a potential oral overdose is established, the patient may present with severe arrhythmias and seizure; these symptoms could be the 1st sign of intoxication. Adverse reactions may cause the withdrawal from treatment; a lower dose rechallenge may start only after the advice of a physician	200 mg, 400mg Tablet and 100mg/5ml Syrup		Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

নং	প্রস্তুতকারকের নাম	ঔষধের নাম	জেনেরিক নাম	‡_iwcDwUK Kvm	নির্দেশনা	Contra-indication & Side-effect	Status (New Molecule/ Existing)	আবেদনকারী প্রদত্ত USFDA/BNF/ MHRA Ref.	‡UKwbK¨vj mve Kvgv‡Ui mfvi wm×vš	mfvi wm×vš
36.	Square Pharmaceuticals Ltd., Pabna Unit, Salgaria, Pabna	Lisdexamfetamine dimesylate 10mg Capsule	Lisdexamfetamine dimesylate INN 10mg	CNS Stimulant	Indicated for the treatment of – - Attention Deficit Hyperactivity Disorder (ADHD) - Moderate to Severe Binge Eating Disorder (BED)	**Contraindications:** Known hypersensitivity to amphetamine products and is also contraindicated with monoamine oxidase (MAO) inhibitor, or within 14 days of the last MAO inhibitor dose. **Side effects:** The most frequent adverse reactions leading to discontinuation were insomnia, tachycardia, irritability, hypertension, headache, anxiety and dyspnea. The most common adverse reactions (incidence ≥5% and at a rate at least twice placebo) reported in children, adolescents, and/or adults were anorexia, anxiety, decreased appetite, decreased weight, diarrhea, dizziness, dry mouth, irritability, insomnia, nausea, upper abdominal pain, and vomiting.	New	USFDA	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 |
37.	Square Pharmaceuticals Ltd., Pabna Unit, Salgaria, Pabna	Lisdexamfetamine Dimesylate 30mg Capsule	Lisdexamfetamine Dimesylate INN 30mg	CNS Stimulant	Indicated for the treatment of – - Attention Deficit Hyperactivity Disorder (ADHD) - Moderate to Severe Binge Eating Disorder (BED)	**Contraindications:** Known hypersensitivity to amphetamine products and is also contraindicated with monoamine oxidase (MAO) inhibitor, or within 14 days of the last MAO inhibitor dose. **Side effects:** The most frequent adverse reactions leading to discontinuation were insomnia, tachycardia, irritability, hypertension, headache, anxiety and dyspnea. The most common adverse reactions (incidence ≥5% and at a rate at least twice placebo) reported in children, adolescents, and/or adults were anorexia, anxiety, decreased appetite, decreased weight, diarrhea, dizziness, dry mouth, irritability, insomnia, nausea, upper abdominal pain, and vomiting.	New	USFDA	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 |

নং	প্রস্তুতকারকের নাম	ঔষধের নাম	জেনেরিক নাম	‡_iwcDwUK Kvm	নির্দেশনা	Contra-indication & Side-effect	Status (New Molecule/ Existing)	আবেদনকারী প্রদত্ত USFDA/BNF/ MHRA Ref.	‡UKwbK`vj mve Kwgwdi mfvi wm×vš	mfvi wm×vš
38.	Aristopharma Ltd, Shampur-Kadamtali I/A, Dhaka-1204	Mesalazine (Mesalamine) 1200 mg DR Tablet	Mesalazine (Mesalamine) BP 1200 mg	Gastrointestinal Agent	It is an aminosalicylate indicated for: 1. Treatment of mildly to moderately active ulcerative colitis in patients 5 years of age and older. 2. Maintenance of remission of ulcerative colitis in adults.	**Contraindication**: Patients with hypersensitivity to salicylates or to any of the components of the product. **Side effects:** The most common adverse reactions (observed in greater than or equal to 5 percent of adults in controlled clinical studies) were abdominal pain, eructation, pain, back pain, rash, dyspepsia, rhinitis, flu syndrome, asthenia, flatulence, vomiting, fever, arthralgia, constipation, and gastrointestinal bleeding.	400mg DR Tablet	USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
39.	Square Pharmaceuticals Ltd., Pabna Unit, Salgaria, Pabna	Mesalazine 800mg Delayed Release Tablet	Mesalazine USP 800mg	Gastrointestinal Agent	Indicated for the treatment of • mild to moderate ulcerative colitis and acute attack • maintenance of remission of ulcerative colitis • maintenance of remission of crohn's ileo-colitis	**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%). Hypersensitivity reactions and drug fever may occasionally occur.	250mg, 500mg ER Capsule, 1gm/Sachet, 2.0g/Sachet	BNF-72 (page 35)	c‡qvRb ‡bB weavq Av‡e`b bvgÄyi Kiv †h‡Z cv‡i	c‡qvRb ‡bB weavq Av‡e`b bvgÄyi Kiv nj

নং	প্রস্তুতকারকের নাম	ঔষধের নাম	জেনেরিক নাম	‡_iwcDwUK Kvm	নির্দেশনা	Contra-indication & Side-effect	Status (New Molecule/ Existing)	আবেদনকারী প্রদত্ত USFDA/BNF/ MHRA Ref.	‡UKwbK¨vj mve KwgwUi mfvi wm×vš	mfvi wm×vš
40.	Popular Pharmaceuticals Limited	Human Menopausal Gonadotrophin (HMG) (Menotrophi) 100 IU/Vial Lyophilized Injection	Human Menopausal Gonadotrophin (HMG) (Menotrophi) BP 100IU/Vial	Hormone	Sterility in females with hypo- or normogonadotropic ovarian insufficiency: Stimulation of follicle growth. Sterility in males with hypo- or normogonadotropic hypogonadism: in combination with HCG to stimulate spermatogenesis.	**Contraindications:** **In females:** - Pregnancy, - Enlargement of the ovaries or cysts that is not caused by polycystic ovarian syndrome, - Gynecological bleeding whose cause is unknown, - Tumors in the uterus, ovaries and breasts, - Prior hypersensitivity to Menotrophins or to any of the excipients, - A high FSH level indicating primary ovarian failure, - The presence of uncontrolled thyroid and adrenal dysfunction, - The presence of any cause of infertility other than anovulation. **In males:** - Carcinoma of the prostate. - Tumors in the tests, - Normal gonadotrophin levels indicating normal pituitary function, - Elevated gonadotrophin levels indicating primary testicular failure, - Infertility disorders other than hypogonadotropic hypogonadism. The following conditions should be properly treated before HMG-therapy is begun: - Dysfunctions of the thyroid gland and cortex of the suprarenal gland, - Hyperprolactinemia, - Tumors in the pituitary or in the hypothalamic glands.	New		c‡qvRbxq ‡idv‡iÝ Ges Therapeutic benefit D‡jLceK Av‡iv Z_¨vw` `wLj Kivi Rb¨ ejv †h‡Z cv‡i	c‡qvRbxq ‡idv‡iÝ Ges Therapeutic benifit D‡jLceK Av‡iv Z_¨vw` `wLj Kivi Rb¨ ejv nj

নং	প্রস্তুতকারকের নাম	ঔষধের নাম	জেনেরিক নাম	‡_iwcDwUK Kvm	নির্দেশনা	Contra-indication & Side-effect	Status (New Molecule/ Existing)	আবেদনকারী প্রদত্ত USFDA/BNF/ MHRA Ref.	‡UKwbK¨vj mve Kwgwdi mfvi wm×vš	mfvi wm×vš
41.	Popular Pharmaceuticals Limited	Vitamin A 500IU+Ascorbic Acid (DC) BP 75mg+ Vitamin D3 200IU+ Vitamin E 25IU+ Vitamin K 15mcg+ Thiamine 0.75mg + Riboflavin BP 0.85mg+Niacin USP 10mg+Pyridoxine Hydrochloride BP 1mg+Folic Acid BP 0.20mg+ Cyanocobalamin 3mcg+ Biotin BP 0.015mg+ Pantothenic Acid 5mg and Calcium 0.39mg+ Calcium 90.57mg+ Phosphorous 7mg and Calcium 9.04mg+ Iodine 75mcg+ Magnesium 50mg+ Zinc 12.50mg+ Selenium 35mcg+ Copper 1mg+ Manganese 1mg+ Chromium 60mcg+ Molybdenum 37.5mcg+ Lutein	Dry Vitamin A Acetate 500 BP 1.00mg eq. to Vitamin A 500IU+Ascorbic Acid (DC) BP 75mg+Dry Vitamin D3 100 BP 2mg eq. to Vitamin D3 200IU+Dry Vitamin E Acetate 50% DC BP 50mg eq. to Vitamin E 25IU+Dry Vitamin K1 5% BP 0.30mg eq. to Vitamin K 15mcg+Thiamine Nitrate BP 0.93mg eq. to Thiamine 0.75mg+Riboflavin BP 0.85mg+Niacin USP 10mg+Pyridoxine Hydrochloride BP 1mg+Folic Acid BP 0.20mg+Cyanocobalamin (1%) BP 0.30mg eq. to Cyanocobalamin 3mcg+Biotin BP 0.015mg+Calcium Pantothenate BP 4.663mg eq. to Pantothenic Acid 5mg and Calcium 0.39mg+Calcium Carbonate (Heavy) BP 226.425mg eq. to Calcium 90.57mg+Anhydroous Calcium Hydrogen Phosphate BP 30.742mg	Multivitamins and minerals	Multivitamins and minerals are used to provide substances that are not taken in through the diet. Multivitamins and minerals are also used to treat vitamin or mineral deficiencies caused by illness, pregnancy, poor nutrition, digestive disorders, certain medications, and many other conditions. An all-in-one eye vitamin and multivitamin/ multimineral supplement Helps replenish vital nutrients that can protect eye health Provides the benefits of a complete multivitamin	**Contraindications:** Eye Multivitamin & Mineral Supplement is contraindicated in patients who are hypersensitivity to or any ingredient in the formulation. **Side effects:** When taken as directed, multivitamins and minerals are not expected to cause serious side effects. Common side effects may include: Upset stomach, Headache, Unusual or unpleasant taste in your mouth.			c‡qvRb ‡bB weavq Av‡e`b bvgÄyi Kiv †h‡Z cv‡i	c‡qvRb ‡bB weavq Av‡e`b bvgÄyi Kiv nj

		5mg+ Zeaxanthin 1mg+ Lycopene 150mcg Tablet	eq. to Phosphorous 7mg and Calcium 9.04mg+Potassium Iodide BP 0.098mg eq. to Iodine 75mcg+Magnesium Oxide (Heavy) BP 82.50mg eq. to Magnesium 50mg+Zinc Oxide BP 15.55mg eq. to Zinc 12.50mg+Sodium Selenite (Se 45%) Ph. Grade 0.084mg eq. to Selenium 35mcg+Copper Gluconate USP 27.00mg eq. to Copper 1mg+Mangannese Sulphate Monohydrate BP 3.075mg eq. to Manganese 1mg+Chromic Chloride Hexahydrate USP 0.246mg eq. to Chromium 60mcg + Sodium Molybdate Dihydrate 1% Ph. Grade 9.46mg eq. to Molybdenum 37.5mcg+Lutein (5%) Ph. Grade 100.00mg eq. to Lutein 5mg+Zeaxanthin (5%) Ph. Grade 20mg eq. to Zeaxanthin 1mg+Lycopene (5%) Ph. Grade 3mg eq. to Lycopene 150mcg							

নং	প্রস্তুতকারকের নাম	ঔষধের নাম	জেনেরিক নাম	‡_iwcDwUK Kvm	নির্দেশনা	Contra-indication & Side-effect	Status (New Molecule/ Existing)	আবেদনকারী প্রদত্ত USFDA/BNF/ MHRA Ref.	‡UKwbK¨vj mve Kwgwdi mfvi wm×vš	mfvi wm×vš
42.	Mundipharma (Bangladesh) Private Limited. Mirzapur,JL#08, Gazipur Sadar, Gazipur, Bangladesh	Chlorzoxazone 250mg + Paracetamol 500mg Tablet	Chlorzoxazone USP 250mg + Paracetamol BP 500mg	Muscle relaxant + Analgesic	Indicated For the relief of pain and muscle spasm associated with inflammatory and degenerative processes; fibrositis. myositis, bursitis, tenosynovitis, torticollis, osteoarthritis, trauma, intervertebral disc syndrome, lumbago, sacroiliac pain, muscular and tendinous sprains, contusions, postoperative myalgia, post tooth extraction.	**Contraindications:** Sensitivity to Paracetamol or Chlorzoxazone **Adverse reactions:** Adverse effects reported to occur with Chlorzoxazone include : Gl: nausea, vomiting, epigastric distress CNS : drowsiness, dizziness, lightheadedness, malaise; Skin : Allergic skin rashes (rarely) Hepatic : hepatitis; Miscellaneous : urine	Paracetamol BP 500mg Tablet		c‡qvRbxq ‡idv‡iÝ Ges Therapeutic benefit D‡jLceK Av‡iv Z_¨vw` `wLj Kiv‡i Rb¨ ejv †h‡Z cv‡i	c‡qvRbxq ‡idv‡iÝ Ges Therapeutic benifit D‡jLceK Av‡iv Z_¨vw` `wLj Kiv‡i Rb¨ ejv nj	
43.	Mundipharma (Bangladesh) Private Limited. Mirzapur,JL#08, Gazipur Sadar, Gazipur, Bangladesh	Diclofenac Sodium 50mg + Paracetamol 500mg Tablet.	Diclofenac Sodium BP 50mg + Paracetamol BP 500 mg	NSAID+ Analgesic	Rheumatoid arthritis, Osteoarthritis, ankylosing spondylitis,cervical spondylitis, intervertebral disc syndrome and sciatica. Non articular rheumatic conditions such as fibrositis, myositis, bursitis, low back pain, Soft tissue injuries such as sprains, strains and sports injury. Painful inflammatory condition in gynaecology. Post-operative and Post traumatic inflammation and swelling, The initial daily dosage for adult is one tablet, two or three times in a day. The drug should be taken with or after meals. For long term therapy, one tablet, two times a day is sufficient.	**Contraindications:** Known hypersensitivity to the Diclofenac sodium or Paracetamol Peptic ulcer, In patients with asthma, seasonal allergic rhinitis, swelling of the nasal mucosa (i.e. nasal polyps), chronic obstructive pulmonary diseases or chronic infections of the respiratory tract (especially if linked to allergic rhinitis-like symptoms), reactions on NSAIDs like asthma exacerbations,. **Side effects:** At recommended dosages Diclomol Plus is generally well tolerated. At the start of treatment, however, patients may sometimes complain of epigastric pain, nausea, diarrhea, dizziness or headache. These unwanted effects are usually of a mild nature. Peripheral oedema and skin reactions such as drug rash, urticarial and eczema have also been observed.	Diclofenac Sodium BP 50mg Tablet Paracetamol 500mg Tablet		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

নং	প্রস্তুতকারকের নাম	ঔষধের নাম	জেনেরিক নাম	‡_iwcDwUK Kvm	নির্দেশনা	Contra-indication & Side-effect	Status (New Molecule/ Existing)	আবেদনকারী প্রদত্ত USFDA/BNF/ MHRA Ref.	‡UKwbK¨vj mve KwgwUi mfvi wm×vš	mfvi wm×vš
44.	Aristopharma Ltd, Shampur-Kadamtali I/A, Dhaka-1204	Vonoprazan 10mg Film Coated Tablet	Vonoprazan Fumarate INN 13.36mg eq. to Vonoprazan 10 mg	Potassium-Competitive Acid Blocker	Vonoprazan Fumarate is a first-in-class potassium-competitive acid blocker. It inhibits H^+, K^+-ATPase activities in a reversible and potassium-competitive manner with a potency of inhibition approximately 350 times higher than the proton pump inhibitor, Lansoprazole. Vonoprazan Fumarate tablet is indicated in the treatment of Gastroduodenal Ulcer, Secondary prevention of Reflux Esophagitis, Secondary prevention of low-dose aspirin or non-steroidal anti-inflammatory drug induced Peptic Ulcer, in combination with Clarithromycin and Amoxicillin for the eradication of Hlicobacter pylori.	**Contra-indication:-** Caution is necessary when oral vonoprazan Fumarate is used in patients with the following conditions and frequent monitoring is necessary: Liver/ Renal disorders, allergic rections (itch, rash, etc.) or in those who are taking any other medicinal products, over-the-counter medicines & dietary supplements as well as other prescription medicines. Caution is also necessary for women who are pregnant & breastfeeding. **Side-effect:-** The most commonly reported adverse reactions include constipation, diarrhea, enlarged feeling of abdomen, nausea, rash and edema.	New		c‡qvRbxq ‡idv‡iÝ Ges Therapeutic benefit D‡jLceK Av‡iv Z_¨vw` `wLj Kivi Rb¨ ejv †h‡Z cv‡i	c‡qvRbxq ‡idv‡iÝ Ges Therapeutic benifit D‡jLceK Av‡iv Z_¨vw` `wLj Kivi Rb¨ ejv nj	
45.	Aristopharma Ltd, Shampur-Kadamtali I/A, Dhaka-1204	Vonoprazan 20 mg Film Coated Tablet	Vonoprazan Fumarate 26.72mg eq. to Vonoprazan 20mg	Potassium-Competitive Acid Blocker	Vonoprazan Fumarate is a first-in-class potassium-competitive acid blocker. It inhibits H^+, K^+-ATPase activities in a reversible and potassium-competitive manner with a potency of inhibition approximately 350 times higher than the proton pump inhibitor, Lansoprazole. Vonoprazan Fumarate tablet is indicated in the treatment of Gastroduodenal Ulcer, Secondary prevention of Reflux Esophagitis, Secondary prevention of low-dose aspirin or non-steroidal anti-inflammatory drug induced Peptic Ulcer, in combination with Clarithromycin and Amoxicillin for the eradication of Hlicobacter pylori.	**Contra-indication:-** Caution is necessary when oral vonoprazan Fumarate is used in patients with the following conditions and frequent monitoring is necessary: Liver/ Renal disorders, allergic rections (itch, rash, etc.) or in those who are taking any other medicinal products, over-the-counter medicines & dietary supplements as well as other prescription medicines. Caution is also necessary for women who are pregnant & breastfeeding. **Side-effect:-** The most commonly reported adverse reactions include constipation, diarrhea, enlarged feeling of abdomen, nausea, rash and edema.	New		c‡qvRbxq ‡idv‡iÝ Ges Therapeutic benefit D‡jLceK Av‡iv Z_¨vw` `wLj Kivi Rb¨ ejv †h‡Z cv‡i	c‡qvRbxq ‡idv‡iÝ Ges Therapeutic benifit D‡jLceK Av‡iv Z_¨vw` `wLj Kivi Rb¨ ejv nj	

নং	প্রস্তুতকারকের নাম	ঔষধের নাম	জেনেরিক নাম	‡_iwcDwUK Kvm	নির্দেশনা	Contra-indication & Side-effect	Status (New Molecule/ Existing)	আবেদনকারী প্রদত্ত USFDA/BNF/ MHRA Ref.	‡UKwbK¨vj mve KwgwUi mfvi wm×vš	mfvi wm×vš
46.	Aristopharma Ltd, Shampur-Kadamtali I/A, Dhaka-1204	Rabeprazole Sodium 20 mg/Vial Lyophilized Injection	Rabeprazole Sodium (Injectable Grade) INN 20 mg/Vial	Proton Pump Inhibitor	It is an alternative in patients for whom oral administration of Rabeprazole is not indicated. Rabeprazole Injection is indicated in the treatment of: * Sequential-therapy (step-up) from oral Rabeprazole, e.g. a patient previously on oral Rabeprazole who is temporarily unable to take oral medication for any reason. * Active duodenal ulcer with bleeding or severe erosions. * Active gastric ulcer with bleeding or severe erosions. *Short-term treatment of erosive or ulcerative gastroesophageal reflux disease (GERD) * Prevention of acid-aspiration. * Stress-induced mucosal injury in critical care. *Pathological hypersecretory conditions, including Zollinger-Ellison syndrome.	**Contraindication:** Rabeprazole is contraindicated in patients with known hypersensitivity to Rabeprazole, substituted Benzimidazoles or to any component of the formulation. **Side-effect:** Headache, abdominal pain, diarrhea, dry mouth, dizziness, peripheral edema, hepatic enzyme increase, hepatitis, hepatic encephalopathy, myalgia, and and arthralgia.	20mg Tablet & Capsule		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
47.	**Acme Laboratories Ltd., Dhamrai, Dhaka**	Sumatriptan 0.50gm/1000 Actuations (100 ml) Nasal Spray	Sumatriptan USP 0.50gm/1000 Actuations (100 ml)	Serotonin 5-HT1 Receptor Agonists	Acute treatment of migraine.	**Contraindication:** Sumatriptan Nasal Spray is contraindicated in patients with: • Ischemic coronary artery disease (CAD) (angina pectoris, history of myocardial infarction, documented silent ischemia) or coronary artery vasospasm, including Prinzmetal's angina. • Wolff-Parkinson-White syndrome or arrhythmias associated with other cardiac accessory conduction pathway disorders. • History of stroke, transient ischemic attack (TIA), or history of hemiplegic or basilar migraine because these patients are at a higher risk of stroke. • Peripheral vascular disease.	**100mg, 50 mg Tablet**	USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

						• Ischemic bowel disease. • Uncontrolled hypertension. • Recent use (i.e., within 24 hours) of ergotamine-containing medication, ergot-type medication (such as dihydroergotamine or methysergide), or another 5hydroxytryptamine1 (5-HT1) agonist. • Concurrent administration of a monoamine oxidase (MAO)-A inhibitor or recent (within 2 weeks) use of an MAO-A inhibitor. • Hypersensitivity to Sumatriptan (angioedema and anaphylaxis seen). • Severe hepatic impairment. **Side effects**: The following side effects are discussed in more detail in other sections of the prescribing information: Myocardial ischemia, myocardial infarction, and prinzmetal's angina, arrhythmias, chest, throat, neck, and/or jaw pain/tightness/pressure, cerebrovascular events, other vasospasm reactions, medication overuse headache, serotonin syndrome , increase in blood pressure , local irritation, hypersensitivity reactions, seizures.				

নং	প্রস্তুতকারকের নাম	ঔষধের নাম	জেনেরিক নাম	‡_iwcDwUK Kvm	নির্দেশনা	Contra-indication & Side-effect	Status (New Molecule/ Existing)	আবেদনকারী প্রদত্ত USFDA/BNF/ MHRA Ref.	‡UKwbK¨vj mve Kwgwdi mfvi wm×vš	mfvi wm×vš
48.	**Acme Laboratories Ltd., Dhamrai, Dhaka**	Sumatriptan 2gm/1000 Actuations (100 ml) Nasal Spray	Sumatriptan USP 2gm/1000 Actuations (100 ml)	Serotonin 5-HT1 Receptor Agonists	Acute treatment of migraine.	**Contraindication:** Sumatriptan Nasal Spray is contraindicated in patients with: • Ischemic coronary artery disease (CAD) (angina pectoris, history of myocardial infarction, documented silent ischemia) or coronary artery vasospasm, including Prinzmetal's angina. • Wolff-Parkinson-White syndrome or arrhythmias associated with other cardiac accessory conduction pathway disorders. • History of stroke, transient ischemic attack (TIA), or history of hemiplegic or basilar migraine because these patients are at a higher risk of stroke. • Peripheral vascular disease. • Ischemic bowel disease. • Uncontrolled hypertension. • Recent use (i.e., within 24 hours) of ergotamine-containing medication, ergot-type medication (such as dihydroergotamine or methysergide), or another 5hydroxytryptamine1 (5-HT1) agonist. • Concurrent administration of a monoamine oxidase (MAO)-A inhibitor or recent (within 2 weeks) use of an MAO-A inhibitor. • Hypersensitivity to Sumatriptan (angioedema and anaphylaxis seen). • Severe hepatic impairment. **Side effects**: The following side effects are discussed in more detail in other sections of the prescribing information: Myocardial ischemia, myocardial infarction, and prinzmetal's angina, arrhythmias, chest, throat, neck, and/or jaw pain/tightness/pressure, cerebrovascular events, other vasospasm reactions, medication overuse headache, serotonin syndrome , increase in blood pressure , local irritation, hypersensitivity reactions, seizures.	**100 mg, 50 mg Tablet**	USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

নং	প্রস্তুতকারকের নাম	ঔষধের নাম	জেনেরিক নাম	‡_iwcDwUK Kvm	নির্দেশনা	Contra-indication & Side-effect	Status (New Molecule/ Existing)	আবেদনকারী প্রদত্ত USFDA/BNF/ MHRA Ref.	‡UKwbK¨vj mve Kwgwdi mfvi wm×vš	mfvi wm×vš
49.	Square Pharmaceuticals Ltd., Pabna Unit, Salgaria, Pabna	Prucalopride 1mg Tablet	Prucalopride INN 1mg	Serotonin 5-HT_4 receptor agonist	This is a selective serotonin (5-HT_4) receptor agonist which is indicated for symptomatic treatment of chronic constipation in adults especially in women when other laxatives fail to provide adequate relief.	**Contraindications:** Crohn's disease intestinal obstruction, intestinal perforation, severe inflammatory condition of intestinal tract, toxic megacolon and ulcerative colitis. Hypersensitivity to the active substance or to any of the excipients. Renal impairment requiring dialysis	New	BNF-72 Page: 53	c‡qvRb ‡bB ïeavq Avi‡e`b bv gÄwy Kiv †h‡Z cv‡i	c‡qvRb ‡bB ïeavq Avi‡e`b bv gÄwy Kiv nj
50.	Square Pharmaceuticals Ltd., Pabna Unit, Salgaria, Pabna	Prucalopride 2mg Tablet	Prucalopride INN 2mg	Serotonin 5-HT_4 receptor agonist	-Do-	-Do-	New	BNF-72 Page: 53	c‡qvRb ‡bB ïeavq Avi‡e`b bvgÄwy Kiv †h‡Z cv‡i	c‡qvRb ‡bB ïeavq Avi‡e`b bvgÄwy Kiv nj
51.	**Acme Laboratories Ltd., Dhamrai, Dhaka**	Cholecalciferol (Vitamin D_3), Oily (Potency: 1 MIU/g) 14,40,000 IU Vitamin D_3/100 ml Syrup	Cholecalciferol (Vitamin D_3), Oily (Potency: 1 MIU/g) BP 1.440 gm eq. to 14,40,000 IU Vitamin D_3/100 ml	Vitamin	Vitamin D helps the body to utilize calcium necessary for the normal development of healthy bones and teeth. Vitamin D3 Pediatric Drops is indicated for use in patients deficient in Vitamin D. Vitamin D is generally produced by the body through sunlight and is necessary for the absorption of calcium. Patients who are not exposed to sufficient sunlight are unable to absorb Vitamin D require additional Vitamin D supplementation. Vitamin D supplementation is required in persons with rickets, inadequate exposure to sunlight infantile tetany and osteomalacia in adults.	**Contraindication:** Hypersensitivity to Vitamin D. Vitamin D should not be given to patients with hypercalcemia, renal impairment, calculi or heart disease. **Side effects**: Vitamin D in excessive amounts may cause hypercalcaemia.	**400 IU, 1000 IU Tablet, 25 mcg/ 5ml syrup, 5mg/ml Injection**		c‡qvRb ‡bB ïeavq Avi‡e`b bvgÄwy Kiv †h‡Z cv‡i	c‡qvRb ‡bB ïeavq Avi‡e`b bvgÄwy Kiv nj
52.	**Ziska Pharmaceuticals Ltd. Gazipure**	Dantrolene Sodium 20mg/70ml Vial Sterile lyophilized Powder for injection	Dantrolene Sodium USP 20mg/70 ml vial	Skeletal Muscle Relaxant	It is indicated, along with appropriate supportive measures, for the management of the fulminant hypermetabolism of skeletal muscle characteristic of malignant hyperthermia crises in patients of all ages. Dantrium Intravenous should be administered by continuous rapid intravenous push as soon as the malignant hyperthermia reaction is recognized (i.e., tachycardia, tachypnea, central venous desaturation, hypercarbia, metabolic acidosis, skeletal muscle rigidity, increased utilization of anesthesia circuit carbon dioxide absorber, cyanosis and mottling of the skin, and, in many cases, fever). Dantrium Intravenous is also indicated preoperatively, and sometimes postoperatively, to prevent or attenuate the development of clinical and laboratory signs of malignant hyperthermia in individuals judged to be malignant hyperthermia susceptible.	Contraindications: None Side effects: There have been occasional reports of death following malignant hyperthermia crisis even when treated with intravenous dantrolene; incidence figures are not available (the pre-dantrolene mortality of malignant hyperthermia crisis was approximately 50%). Most of these deaths can be accounted for by late recognition, delayed treatment, inadequate dosage, lack of supportive therapy, intercurrent disease and/or the development of delayed complications such as renal failure or disseminated intravascular coagulopathy. In some cases there are insufficient data to completely rule out therapeutic failure of dantrolene.	25mg & 50 mg capsule	USFDA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

Annex-B: Proposed Product for Import (Human)

নং	c¯ZKvi‡Ki bvg	JI‡ai bvg	†RvbvwiK bvg	‡_iwcDwUK Kvm	wb‡`kbv	Contraindication & Side-effect	Status (New Molecule/ Existing)	FSC/CPP	‡UKwbK¨vj mve-KvgwUi mfvi wm×vš	mfvi wm×vš
1.	ALMAC Pharma Services Ltd., UK for **GALEN LTD., SEAGOE INDUSTRIAL ESTATE, Craigavon, BT63 5UA, UK** **Local agent:** Radiant Export Import Enterprise. Uttara, Dhaka.	**DaunoXome** Injection 2mg/ml concentrate for solution for Infusion (50mg/25ml vial)	Daunorubicin 50mg/Dose	Anticancer	It is indicated for the treatment of Advanced AIDS-related Kaposi's sarcoma.	**Contraindication**: Myocardial insufficiency, previous treatment with maximum cumulative doses of daunorubicin or other anthracycline, recent myocardial infarction, severe arrhythmia. **CAUTIONS** Caution in handling—irritant to tissues **SIDE-EFFECTS** Alopecia, extravasation, bone-marrow suppression, hyperuricaemia nausea oral mucositis thromboembolism tumour lysis syndrome vomiting **Side-Effects, Further Information** **Cardiotoxicity** All anthracycline antibiotics have been associated with varying degrees of cardiac toxicity—this may be idiosyncratic and reversible, but is commonly related to total cumulative dose and is irreversible**.** **HEPATIC IMPAIRMENT** Reduce dose according to serum bilirubin concentration—consult local protocol for details. Avoid in severe impairment. **RENAL IMPAIRMENT** Reduce dose by 25% if serum creatinine 105–265 micromol/litre. Reduce dose by 50% if serum creatinine greater than 265 micromol/litre. Avoid in severe impairment. **MONITORING REQUIREMENTS** Cardiac monitoring essential.	20 mg/10 ml Injection	MHRA-UK	Abyjgv`b Kiv †h‡Z cv‡i|	Abyjgv`b Kiv nj|

নং	c¯ZKvi‡Ki bvg	Jl‡ai bvg	†Rvbv‡iK bvg	‡_‡iwcDwUK Kvm	wb‡`kbv	Contraindication & Side-effect	Status (New Molecule/ Existing)	FSC/CPP	‡UKwbK¨vj mve-Kvgv‡Ui mfvi wm×vš	mfvi wm×vš
2.	**Chiesi Farmaceuitici S.p.A, Via** Palermo, 26/A-43100 Parma-Italy **Local agent:** Radiant Export Import Enterprise. Uttara, Dhaka.	**CUROSURF** Intratracheal Suspension (Lung Surfactanat) 80mg/ml vial	Poractant alfa INN (procine lung phospholipid fraction) Intratracheal Suspension (1.5ml vial contains 120 mg ; 3ml vial contains 240 mg)	Lung Surfactant	treatment of respiratory distress syndrome in neonates over 700 g; prophylaxis of respiratory distress syndrome in preterm neonates 24–31 weeks post-menstrual age	**Contraindication**: None **Side effects**: Pulmonary surfactants have been associated with intracranial haemorrhage. Bradycardia, pulmonary haemorrhage, and decreased oxygen saturation have been reported rarely; hyperoxia and obstruction of the endotracheal tube by mucous secretions have also been reported.	New	FSC Not submitted	c`wU ¯wMZ Ges ‡idv‡iÝ I Avi I Z_¨vmn cieZx mfvq Av‡e`b `wLj Kivi Rb¨ ejv †h‡Z cv‡i	c`wU ¯wMZ Ges ‡idv‡iÝ I Avi I Z_¨vmn cieZx mfvq Av‡e`b `wLj Kivi Rb¨ ejv nj	
3.	**Fresenius Kabi Austria GmbH,** Hafnerstrasse 36, A-8055 Graz, Austria for **Fresenius Kabi** Deutchland GmbH, Germany Local agent: Radiant Export Import Enterprise. Uttara, Dhaka.	Aminoven Infant 10% 100 ml/glass bottle For intravenous infusion (cava catheter recommended)	Arginine Ph. Eur 7.50gm +Isoleucine Ph. Eur 8.0gm + L- Leucine Ph. Eur 13.0gm + L-Lysine Acetate 12.0gm + Methionine Ph. Eur 3.12gm + Phenylalanine Ph. Eur 3.75gm + Threonine Ph. Eur 4.40gm + Tryptophan Ph. Eur 2.01gm + Valine Ph. Eur 9.0gm + Glycine Ph. Eur 4.15gm + Alanine Ph. Eur 9.30gm + Proline Ph. Eur 9.71gm + Histidine Ph. Eur 4.76gm + Serine Ph. Eur 7.67gm + N-acetyl-L-tyrosine 5.176gm + Taurine 0.40gm + L-malic acid 2.62gm + Acetyl-L-cysteine 0.70gm/1000ml	Amino Acid	10% amino acid solution for partial parenteral nutrition of infants (preterm, underterm newborns, babies) and young children. Together with corresponding amounts of carbohydrates and fat as energy donors, and vitamins, electrolytes and trace elements, the solution may serve for total parenteral nutrition.	**Contraindications:** As for all amino acid solutions Aminoven infant 10% should not be administered in the following conditions: Disturbances in amino acid metabolism, metabolic acidosis, hyperhydratation, hyperkalemia. Patient with insufficient renal or hepatic function require an individual dosage. Attention in case of hyponatremia. **Special Warnings And Precautions For Use** Frequent evaluation and determination of the following laboratory values should be recommended for monitoring parenteral nutrition in infants: urea-nitrogen, ammonia, electrolytes, glucose and triglycerides (when a fat emulsion is administered), acid-base and fluid balance, liver enzymes and serum osmolality. Infusion via peripheral veins in general can cause irritation of the vein intima and	New	Germany	c`wU ¯wMZ Ges Therapeutic benefit D‡jLceK Av‡iv Z_¨vw` `wLj Kivi Rb¨ ejv †h‡Z cv‡i	c`wU ¯wMZ Ges Therapeutic benefit D‡jLceK Av‡iv Z_¨vw` `wLj Kivi Rb¨ ejv nj	

						thrombophlebitis. To minimize the risk of vein irritation, daily controls of the punction site are recommended. Aminoven infant 10% is applicable as part of total parenteral nutrition regimen in combination with adequate amounts of enery donors (carbohydrate solutions, fat emulsions), electrolytes, vitamins and trace elements. **Side effects**: None known when correctly administered				
4.	**Fresenius Kabi Deutschland GmbH,** Freseniusstraβe 61169 Friedberg, Germany **Local agent:** Radiant Export Import Enterprise. Uttara, Dhaka.	Volulyte 6% solution for infusion 500 ml in a free flex bag	Each 1000 ml solution for infusion contain: Poly(O-2-hydroxyethyl)starch (Ph.Eur.) 60.00 gm + - Molar substitution: 0.38 – 0.45 - Mean molecular weight: 130,000 Da (manufactured from waxy maize starch) Sodium acetate trihydrate Ph.Eur 4.63 gm Sodium chloride Ph.Eur 6.02 gm Potassium chloride Ph.Eur 0.30 gm Magnesium chloride hexahydrate Ph.Eur 0.30 gm	Isotonic Solution	Treatment of hypovolaemia due to acute blood loss when crystalloids alone are not sufficient	**Contraindication**: Burns, cerebral haemorrhage, critically ill patients, dehydration, hyperhydration, intracranial haemorrhage, pulmonary oedema, sepsis, severe coagulopathy **CAUTIONS** Cardiac disease, care should be taken to avoid haematocrit concentration from falling below 25-30%; children, renal impairment, severe liver disease, surgery, trauma **SIDE-EFFECTS:** Rare Severe anaphylactic reactions Frequency not known Hypersensitivity reactions, pruritus, raised serum amylase, transient increase in bleeding time **HEPATIC IMPAIRMENT** Avoid in severe impairment. **RENAL IMPAIRMENT** Avoid	New	Germany	*Abyjgv`b Kivth‡Z cv‡i|*	*Abyjgv`b Kivnj|*

নং	c¯ZKvi‡Ki bvg	JI‡ai bvg	†RvbviK bvg	‡_iwcDwUK Kvm	wb‡`kbv	Contraindication & Side-effect	Status (New Molecule/ Existing)	FSC/CPP	‡UKwbK¨vj mve-KwgwUi mfvi wm×vš	mfvi wm×vš
5.	Made for F. Hoffmann-La Roche Ltd, Basel, Switzerland. By Vetter Pharma-Fertigung GmbH & Co. KG, Ravensburg, Germany **Local agent:** Roche Bangladesh Limited	Actemra® 162mg/0.9 ml Pre-filled syringes	Tocilizumab INN 162mg/0.9ml	Immunological Agent	Tocilizumab is indicated for the treatment of moderate to severe active rheumatoid arthritis (RA) in adult patients. Tocilizumab can be used alone or in combination with methotrexate (MTX) and/or other disease-modifying anti-rheumatic drugs (DMARDs). Tocilizumab has been shown to inhibit progression of joint damage as measured by X-ray and to improve physical function.	**Contraindication**: Known hypersensitivity to tocilizumab or to any of the excipients. **Side effects**: Abdominal pain, antibody formation, dizziness, gastritis, headache, hypercholesterolemia, hypersensitivity, hypertension, infection, leucopenia, mouth ulceration, neutropenia, peripheral oedema, pruritis, raised hepatic Transaminases. rash, upper respiratory-tract infection	New	EMA	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj
6.	Made for F. Hoffmann-La Roche, Ltd. Basel, Switzerland. by Roche Diagnostics GmbH, Mannheim, Germany **Local agent:** Roche Bangladesh Limited	Gazyva® 1000mg/40 ml vial concentrate for solution for infusion	Obinutuzumab INN 1000mg/40ml Vial	Anticancer	Gazyva in combination with chlorambucil is indicated for the treatment of patients with Previously untreated chronic lymphocytic leukemia (CLL).	**Contraindication**: Gazyva is contraindicated in patients with a known hypersensitivity (IgE-mediated) to obinutuzumab or to any of the excipients. **Side effects**: Infusion-related reaction, Neutropenia, Thrombocytopenia, Pyrexia, Cough, Diarrhea, Leukopenia.	New	EMA	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj
7.	**Manufacturer :** Cilag AG, Hochstrasse 201, 8200 sCHAFFHAUSEN, Switzerland **Local agent:** Unihealth Limited, House : 46, Sheikh Kalam saroni, Road no. 16, Rangs Nasim Square (6th Floor), Dhanmondi	Imbruvica 140mg Capsule	Ibrutinib INN 140mg	Anticancer	It is indicated for the the following indication: a) Mantle cell lymphoma b) Chronic lymphocytic leukemia/Small lymphocytic lymphoma c) Chronic lymphocytic leukemia/Small lymphocytic lymphoma with deletion 17 p.	**Side Effect** : Adverse reaction are adverse events that have been considered to be reasonably casually associaated with the use of ibrutinib based on the comprehensive assessment of the avilable adverse event information. A causal relationship with ibrutinib cannot be reliably established in individual case. Further, beacuse 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 clinical practice.	New	Switzerland	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj

নং	c¯ZKvi‡Ki bvg	JI‡ai bvg	†Rvbv‡iK bvg	‡_‡iwcDwUK Kvm	wb‡`kbv	Contraindication & Side-effect	Status (New Molecule/ Existing)	FSC/CPP	‡UKwbK¨vj mve-KvgwUi mfvi wm×vš	mfvi wm×vš
8.	**Manufacturer :** Eli lilly and Company Indianapolis, IN 46285, USA **Local agent:** International Agencies (Bd.) Ltd.	Trulicity 1.5mg/0.5ml Solution for Injection in pre-filled pen	Dulaglutide 1.5mg/ 0.5ml	Antidiabetic	Trulicity is indicated in adult with type 2 diabetes mellitus to inmrove glycaemic control as : **Monotherapy** when diet and exercise alone do not provide adequate glycaemic control in patients for whom the use of metformin is considered inappropriate due to intoolerance or Contraindications **add-on therapy** In combination with other glucose-lowering medicinal products including insulin, When these, together with diet and exercis, do not provide adequate glycaemic control (See section 5.1 for data with respect to different combinations)	**Contraindications:** Hypersensitivity to the active substance or to any of the excipients. **Side Effect**: Dulaglutide should not be used in patients with type 1 diabetes mellitus or for the treatment of diabetic ketoacidosis.	New	USA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
9.	**Manufacturer :** Eli lilly and Company, Indianapolis, IN 46285, USA. **Local agent:** International Agencies (Bd.) Ltd.	Trulicity 0.75mg/0.5ml Solution for Injection in pre-filled pen	Dulaglutide 0.75mg/0.5ml	Antidiabetic	Trulicity is indicated in adult with type 2 diabetes mellitus to inmrove glycaemic control as : **Monotherapy** when diet and exercise alone do not provide adequate glycaemic control in patients for whom the use of metformin is considered inappropriate due to intoolerance or Contraindications **add-on therapy** In combination with other glucose-lowering medicinal products including insulin, When these, together with diet and exercis, do not provide adequate glycaemic control (See section 5.1 for data with respect to different combinations)	**Contraindications:** Hypersensitivity to the active substance or to any of the excipients. **Side Effect**: Dulaglutide should not be used in patients with type 1 diabetes mellitus or for the treatment of diabetic ketoacidosis.	New	USA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

নং	c¯ZKvi‡Ki bvg	Jl‡ai bvg	†Rvbv‡iK bvg	‡_‡ivwcDwUK Kvm	wb‡`kbv	Contraindication & Side-effect	Status (New Molecule/ Existing)	FSC/CPP	‡UKwbK¨vj mve-Kvgvi mfvi wm×vš	mfvi wm×vš
10.	**Manufacturer:** Sandoz GmbH, Austria **Local Agent:** (Novartis Bangladesh Ltd) AHN Tower (7th) floor, 13, bir uttam c.r. dutta road Biponan, C/A, Dhaka-1000.	Sandostatin LAR Powder and solvent for suspension for injection 10 mg	Octreotide Acetate Ph. Gr. 11.20mg eq. to Octreotide INN 10 mg	Anticancer	Treatment of patients with ***acromegaly:*** • in whom surgery or radiotherapy is inappropriate or ineffective, or in the interim period until radiotherapy becomes fully effective. ***Gastro-entero-pancreatic (GEP) endocrine tumors:*** Treatment of patients with symptoms associated with functional gastro-entero-pancreatic (GEP) endocrine tumors: • Carcinoid tumors with features of the carcinoid syndrome. • VIPomas. • Glucagonomas. • Gastrinomas/Zollinger-Ellison syndrome. • Insulinomas, for pre-operative control of hypoglycemia and for maintenance therapy. • GRFomas. Treatment of patients with advanced, well-differentiated (G1, G2) Neuroendocrine Tumors of the midgut.	**Contraindication:** Known hypersensitivity to octreotide or to any of the excipients. **Side Effects:** Most commonly reported side effects are diarrhea, abdominal pain, nausea, flatulence, headache, hyperglycaemia and constipation. Commonly reported effects are dizziness, localized pain, thyroid dysfunction, vomiting, loose stools, impaired glucose tolerance and hypoglycaemia.	New	Austria and Switzerland	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
11.	**Manufacturer:** Sandoz GmbH, Austria **Local Agent:** (Novartis Bangladesh Ltd) AHN Tower (7th) floor, 13, bir uttam c.r. dutta road Biponan, C/A, Dhaka-1000.	Sandostatin LAR Powder and solvent for suspension for injection 20 mg	Octreotide Acetate Ph. Gr. 22.40mg eq. to Octreotide INN 20 mg	Anticancer	DO	DO	New	Austria and Switzerland	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

ক্রং	c¯ZKvi‡Ki bvg	JI‡ai bvg	†Rvbv‡iK bvg	‡_‡ivwcDwUK Kvm	wb‡`kbv	Contraindication & Side-effect	Status (New Molecule/ Existing)	FSC/CPP	‡UKwbK¨vj mve-KvgvUi mfvi wm×vš	mfvi wm×vš
12.	**Manufacturer:**Sandoz GmbH, Austria **Local Agent:** (Novartis Bangladesh Ltd) AHN Tower (7th) floor, 13, bir uttam c.r. dutta road Biponan, C/A, Dhaka-1000.	Sandostatin LAR 30 mg Powder and solvent for suspension for injection	Octreotide Acetate Ph. Gr. 33.60mg eq. to Octreotide INN 30 mg	Anticancer	Treatment of patients with ***acromegaly:*** • in whom surgery or radiotherapy is inappropriate or ineffective, or in the interim period until radiotherapy becomes fully effective. ***Gastro-entero-pancreatic (GEP) endocrine tumors:*** Treatment of patients with symptoms associated with functional gastro-entero-pancreatic (GEP) endocrine tumors: • Carcinoid tumors with features of the carcinoid syndrome. • VIPomas. • Glucagonomas. • Gastrinomas/Zollinger-Ellison syndrome. • Insulinomas, for pre-operative control of hypoglycemia and for maintenance therapy. • GRFomas. Treatment of patients with advanced, well-differentiated (G1, G2) Neuroendocrine Tumors of the midgut.	**Contraindication:** Known hypersensitivity to octreotide or to any of the excipients. **Side Effects:** Most commonly reported side effects are diarrhea, abdominal pain, nausea, flatulence, headache, hyperglycaemia and constipation. Commonly reported effects are dizziness, localized pain, thyroid dysfunction, vomiting, loose stools, impaired glucose tolerance and hypoglycaemia.	New	Austria and Switzerland	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
13.	Merck Serono S.p.A. Italy (Responsible for batch release), Merck Serono S.A. Switzerland (Responsible for primary & secondary packaging) **Local agent :** Janata Traders TCB Bhabon , 1 kawran Bazar, Dhaka	Gonal- f solution for Injection in a prefilled pen 300 IU/0.5ml	Follitropin alfa 300IU/0.5ml (22 microgram /0.5ml)	Human Follicle Stimulating Hormone	- Anovulation including polycystic ovarian syndrome in women who have been unresponsive to treatment with clomiphene citrate. - Stimulation of multifollicular development in women undergoing superovulation for assisted reproductive technologies (ART) such as in vitro fertilisation (IVF), gamete intra-fallopian transfer (GIFT) and zygote intra-	**Contraindications**: Allergy (hypersensitivity) to the active ingredient follitropin alfa, FSH or to any of the excipients (see section 'Composition' above) s Tumours of the hypothalamus or pituitary gland s Large ovaries or sacs of fluids within the ovaries (ovarian cyst) not due to polycystic ovarian syndrome s Unexplained vaginal bleeding	New	EMA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

					fallopian transfer (ZIFT).	(gynaecological haemorrhages of unknown aetiology) s Cancer in the ovaries, uterus or breasts GONAL-f® must not be used when an effective response cannot be obtained, such as: s primary ovarian failure s malformations of sexual organs incompatible with pregnancy s fibroid tumours of the uterus incompatible with pregnancy s primary testicular insufficiency Do not use GONAL-f ® if any of the above applies to you. If you are not sure, talk to your doctor or pharmacist before using this medicine. **Side Effect** : The following medical events have been reported subsequent to pregnancies resulting from Gonal-F® (follitropin alfa) therapy in controlled clinical studies: 1. Spontaneous Abortion 2. Ectopic Pregnancy 3. Premature Labor 4. Postpartum Fever 5. Congenital abnormalities				

বং	c¯ZKvi‡Ki bvg	JI‡ai bvg	†RvbviK bvg	‡_iwcDwUK Kvm	wb‡`kbv	Contraindication & Side-effect	Status (New Molecule/ Existing)	FSC/CPP	‡UKwbK¨vj mve-KvgwUi mfvi wm×vš	mfvi wm×vš
14.	Merck Serono S.p.A. Italy (Responsible for batch release), Merck Serono S.A. Switzerland (Responsible for primary & secondary packaging) **Local agent :** Janata Traders TCB Bhabon , 1 kawran Bazar, Dhaka	Gonal- f solution for Injection in a prefilled pen 900IU/1.5ml	Follitropin alfa 900IU/1.5ml (66.0 microgram /1.5ml)	Human Follicle Stimulating Hormone	- Anovulation including polycystic ovarian syndrome in women who have been unresponsive to treatment with clomiphene citrate. - Stimulation of multifollicular development in women undergoing superovulation for assisted reproductive technologies (ART) such as in vitro fertilisation (IVF), gamete intra-fallopian transfer (GIFT) and zygote intra-fallopian transfer (ZIFT).	**Contraindications**: Allergy (hypersensitivity) to the active ingredient follitropin alfa, FSH or to any of the excipients (see section 'Composition' above) s Tumours of the hypothalamus or pituitary gland s Large ovaries or sacs of fluids within the ovaries (ovarian cyst) not due to polycystic ovarian syndrome s Unexplained vaginal bleeding (gynaecological haemorrhages of unknown aetiology) s Cancer in the ovaries, uterus or breasts GONAL-f® must not be used when an effective response cannot be obtained, such as: s primary ovarian failure s malformations of sexual organs incompatible with pregnancy s fibroid tumours of the uterus incompatible with pregnancy s primary testicular insufficiency Do not use GONAL-f ® if any of the above applies to you. If you are not sure, talk to your doctor or pharmacist before using this medicine. **Side Effect** : The following medical events have been reported subsequent to pregnancies resulting from Gonal-F® (follitropin alfa) therapy in controlled clinical studies: 1. Spontaneous Abortion 2. Ectopic Pregnancy 3. Premature Labor 4. Postpartum Fever 5. Congenital abnormalities	New	EMA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

বং	c¯ZKvi‡Ki bvg	JI‡ai bvg	†RvbviK bvg	‡_iwcDwUK Kvm	wb‡`kbv	Contraindication & Side-effect	Status (New Molecule/ Existing)	FSC/CPP	‡UKwbK¨vj mve-KvgwUi mfvi wm×vš	mfvi wm×vš
15.	Trommsdorff GmbH & Co. Kg , TrommsdoffsreBe 2-6 52477 Alsdorf, Germany . **Local agent:** Zas Corporation 80/22, Mymensingh Road , Banglamotor , Dhaka	Zalain Vaginal Suppository	Sertaconazol Nitrate EU. Ph. 0.30g	Antifungal	Local Treatment of infections by Candida of the Vaginal mucosa.	**Contraindication:** Known hypersensitivity to an antimycotic of the imidazoles group or to one of the Excipients. **Side Effect:** The absence of systemic absorption of Sertaconazole after the application of a vaginal suppository by vaginal route makes it hardly probable that side-effects appear.	New	Germany	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
16.	**Manufacturer:** Haupt Pharma Wolfrastshausen GmbH, pfaffenrieder Str.5,82515, Germany **Local agent:** Zas Corporation 80/22, Mymensingh Road, Banglamotor , Dhaka-1000	**Methotrexate PhaRes** 5mg/2ml (Solution for Injection)	Methotrexate 5mg/2ml	Anticancer	Antineoplastic Chemotherapy; Breast Carcinoma, Leukaemia, Meningeal Leukaemia & Psoriasis Chemotherapy.	**Contraindication:** Patients with significantly impaired renal function, hepatic function, Pre-existing blood dyscrasias, such as significant marrow hypoplasia, leucopenia, thrombocytopenia or anaemia, laboratory evidence of immunodeficiency syndrome methotrexate is contraindicated in pregnancy. Methotrexate in breast fed infants allergic hypersensitivity methotrexate. **Side Effect:** Skin: severe, occasionally fatal, dermatologic reactions, Hepatic toxicity resulting in significant elevations of liver enzymes, Urogenital systems renal failure, azotaemia, cystitis, haematuria, blood eosinophilia, headaches, drowsiness, blurred vision, aphasia cognitive disorder Nervous systems. Ear disorders tinnitus, eye disorders Conjunctivitis.Vasculitis, hypotension, thromboembolic events vascular disorder.	New	Germany	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

ক্রং	c¯ZKvi‡Ki bvg	JI‡ai bvg	†RvbviK bvg	‡_ivwcDwUK Kvm	wb‡`kbv	Contraindication & Side-effect	Status (New Molecule/ Existing)	FSC/CPP	‡UKwbK¨vj mve-KvgwUi mfvi wm×vš	mfvi wm×vš
17.	**Manufacturer:** Haupt Pharma Wolfrastshausen GmbH, pfaffenrieder Str.5,82515, Germany **Local agent:** Zas Corporation 80/22, Mymensingh Road, Banglamotor , Dhaka-1000	Methotrexate PhaRes 50mg/2ml (Solution for Injection)	Methotrexate 50mg/2ml	Anticancer	Antineoplastic Chemotherapy; Breast Carcinoma, Leukaemia, Meningeal Leukaemia & Psoriasis Chemotherapy.	**Contraindication:** Patients with significantly impaired renal function, heparic function, Pre-existing blood dyscrasias, such as significant marrow hypoplasia, leucopenia, thrombocytopenia or anaemia, laboratory evidence of immunodeficiency syndrome methotrexate is contraindicated in pregnancy. Methotrexate in breast fed infants allergic hypersensitivity methotrexate. **Side Effect:** Skin: severe, occasionally fatal, dermatologic reactions, hepatic toxicity resulting in significant elevations of liver enzymes, Urogenital systems renal failure, azotaemia, cystitis, haematuria, blood eosinophilia, headaches, drowsiness, blurred vision, aphasia cognitive disorder Nervous systems. Ear disorders tinnitus, eye disorders Conjunctivitis.Vasculitis, hypotension, thromboembolic events vascular disorder	New	Germany	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj

নং	c¯ZKvi‡Ki bvg	JI‡ai bvg	†RvbviK bvg	‡_iwcDwUK Kvm	wb‡`kbv	Contraindication & Side-effect	Status (New Molecule/ Existing)	FSC/CPP	‡UKwbK¨vj mve-KvgwUi mfvi wm×vš	mfvi wm×vš
18.	**Manufacturer:** Haupt Pharma Wolfrastshausen GmbH, pfaffenrieder Str.5,82515, Germany **Local agent:** Zas Corporation 80/22, Mymensingh Road, Banglamotor , Dhaka-1000	Methotrexate PhaRes 1000mg/4ml (Solution for Injection)	Methotrexate 25mg/ml	Anticancer	Antineoplastic Chemotherapy; Breast Carcinoma, Leukaemia, Meningeal Leukaemia & Psoriasis Chemotherapy.	**Contraindication:** Patients with significantly impaired renal function, heparic function, Pre-existing blood dyscrasias, such as significant marrow hypoplasia, leucopenia, thrombocytopenia or anaemia, laboratory evidence of immunodeficiency syndrome methotrexate is contraindicated in pregnancy. Methotrexate in breast fed infants allergic hypersensitivity methotrexate. **Side Effect:** Skin severe, occasionally fatal, dermatologic reactions, hepatic toxicity resulting in significant elevations of liver enzymes, Urogenital systems renal failure, azotaemia, cystitis, haematuria, blood eosinophilia, headaches, drowsiness, blurred vision, aphasia cognitive disorder Nervous systems. Ear disorders tinnitus, eye disorders Conjunctivitis.Vasculitis ,hypotension, thromboembolic events vascular disorder	New	Germany	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

বং	c¯ZKvi‡Ki bvg	JI‡ai bvg	†RvbviK bvg	‡_iwcDwUK Kvm	wb‡`kbv	Contraindication & Side-effect	Status (New Molecule/ Existing)	FSC/CPP	‡UKwbK¨vj mve-KgwUi mfvi wm×vš	mfvi wm×vš
19.	**Manufacturer:** Kedrion S.P.A Via Provincial localita` Bolognana 55027 Gallicano (LU)- Italy. **Local agent:** Zas Corporation 80/22, Mymensingh Road, Banglamotor, Dhaka-1000	IMMUNOHBs (Solution for Injection for Intramuscular use)	Human Hepatitis B Immunoglobulin 180IU/ml	Immunoglobulin	Prophylaxis against Hepatitis B infection in adults and children over 2 years of age who have not been vaccinated against Hepatitis B. Prophylaxis in neonates born to HBsAG positive mother.	**Contraindication:** Hypersensitivity to the active substances or any of the excipients. **Side effect:** Skin reaction, Erythema, Fever, Malaise, Chill, pain at injection site.	New	Italy, Switzerland	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj
20.	**Manufacturer:** Xellia Pharmaceuticals ApS, Dalslandsgade 11 2300 Copenhagen, s, Denmark **Local agent:** Zas Corporation 80/22, Mymensingh Road , Banglamotor , Dhaka-1000	Colistimethate Sodium, 1 Million International Units (IU) Powder for Solution for Injection or Infusion	Colistimethate Sodium, 1 Million International Units (IU)	Antibiotic	Intravenous administration for the treatment of some serious infections caused by Gram-negative bacteria, including those of the lower respiratory tract and urinary tract, when more commonly used systemic antibacterial agents may be contra-indicated or may be ineffective because of bacterial resistance	**Contraindication:** Hypersensitivity to Colistimethate sodium or to polymyxin B In cystic fibrosis patient's neurological events have been reported in up to 27% of patients. These are generally mild and resolve during or shortly after treatment.	New	Denmark & MHRA-UK	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj

Annex-C: Propose Product List for Import (Veterinary)

bs	c¯ZKvi‡Ki bvg I wVKvbv	JI‡ai bvg	†Rvbv‡iK bvg	‡_‡K wcDwUK Kvm	wb‡`kbv	Contraindication & Side-effect	Status (New/Existing)	FSC/CPP	‡UKwbK¨vj mve Kvgv‡Ui mfvi wm×vš	mfvi wm×vš
1.	Manufacturer: SP veterinaria, S.A.,Spain **Local Agent:** Haychem (Bangladesh)Limited, Plot-18/19 BSCIC Industrial area , Maskanda,Mymensing	COLIMICINA Polvo Hidrosoluble 205,000,000IU/100gm Water Soluble Powder	Colistin sulphate 205,000,000IU/100gm	Antibacterial	Prevention and treatment of infections in the gastrointestinal tract, caused by Gram-negative microorganisms: *E. coli*, *Salmonella*, *Shigella*, *Proteus*, *Pseudomonas*, *Klebsiella*, etc. • *Poultry: colibacillosis, salmonellosis, unspecific diarrhoea.* • *Pigs, cattle*: colibacillosis, salmonellosis, bacterial enteritis.	**Contra-indication:** Do not administer in animals with renal insufficiency. **Side-effects:** Not reported.	New	Spain	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
2.	Laboratorios Calier, S.A., Spain **Local Agent:** NEXUS Distributor House no. 49/K jail Road, Mymensingh, Bangladesh	AQUACOLI 2000,000 IU/ml Solution for use in drinking water (Veterinary)	Colistin (as Colistin sulfate) 2MIU/ml	Antibacterial	Treatment and metaphylasix of enteric infections caused by non-invasive Escherichia coli suspectible to colistin. The presence of the disease in the hard should be established before metaphylactic treatment	Contra-indication & Side-effect : Do not use in horse, particularly in foals since colistin, due to to a shift in the gastrointestinal microflora balance could lead to the development of antimicrobial associated colitis (Colitis X), typically associated tieh Clostridium difficile, which may be fatal. Do not use in cases of known hypersensitivity to the active substance or to any of the excipients. Do not use in case of resistance to the polymyxin.	New	Spain	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
3.	**Manufacturer**: SP veterinaria, S.A.,Spain **Local Agent:** Haychem (Bangladesh)Limited, Plot-18/19 BSCIC Industrial area, Maskanda, Mymensing	COCCIVEX Oral solution `	Amprolium Chlorhidate 226.2mg eq. to Amprolium 200 mg/per ml	Anticoccidial	Treatment of coccidiosis in broilers, turkeys, hens, ovine and caprine.	**Contra-indication:** Not reported **Side-effects :** If thiamine-deficient diets, paresia and polyneuritis can occur. It is corrected by administering thiamine	New	Spain	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

bs	c¯ZKvi‡Ki bvg I wVKvbv	Jl‡ai bvg	†Rvbv‡iK bvg	‡_iwcDwUK Kvm	wb‡`kbv	Contraindication & Side-effect	Status (New/Existing)	FSC/CPP	‡UKwbK¨vj mve Kvgv‡Ui mfvi wm×vš	mfvi wm×vš
4.	Laboratorios Calier, S.A., Spain **Local Agent:** NEXUS Distributor House no. 49/K jail Road, Mymensingh, Bangladesh	INMUNAIR 17.5 Solution (Veterinary)	Inactivated Cells of propionibacterium acnes 0.17mg + Lipopolysaccharide from E Coli 0.05 mg/ml	Immunomodulator	In broilers, pullets and hens (layers and breeders : INMUNAIR 17.5 improves the immune status of vaccinated poultry against Marek's disease when applied at three week interval after vaccination. It increases the protection index of vaccinated poultry against infectious bursitis. It reduces mortality, clinical symp0toms and/or lesions caused by the virus of mareik's disease. It reduces lesions and damage of zootechnical parameters produce by the infection of Mycoplasma gallisepticum. It reduces mortality and damage of zootechnial parameters produced by subgroup J avian leukosis virus. It reduces lesions produced by the virus of infectious bursitis. INMUNAIR 17.5 is recommended in critical periods of poultry's productive life when they are under stress conditions and are more susceptible to infectious disease.	**Contraindications:** Have not been described **Side-effect:** Have not been described	New	Spain	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
5.	IZO S.r.l. a socio unico, S.S. 234 Km 28,2-27013 Chignolo Po (PV), ITALY **Local Agent:** Renata Ltd., Mirpur, Dhaka	**IZOVAC H120 – CLONE** freeze-dried Live Vaccine	Live attenuated Newcastle Disease Virus, strain CLONE: 10^6 EID_{50} + Live attenuated Avian Infectious Bronchitis Virus, strain H120: 10^3 EID_{50}	Vaccine	Prevention against Newcastle Disease and Infectious Bronchitis in chickens.	**Contraindications:** There are not contra-indications. **Side Effects:** There are not side-effects.	New	Italy	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
6.	IZO S.r.l. a socio unico, S.S. 234 Km 28,2-27013 Chignolo Po (PV), ITALY **Local Agent:** Renata Ltd., Mirpur, Dhaka	**IZOVAC ND-IBD** Emulsified inactivated Vaccine	Inactivated ND virus: 50 PD_{50} + Inactivated IBD virus strain Winterfield 2512:RP≥1* *RP= Relative potency (Test ELISA), compared to a reference vaccine	Vaccine	Prevention of Newcastle Disease and Gumboro Disease	**Contraindications:** There are not contra-indications. **Side Effects:** There are not side-effects.	New	Italy	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj

bs	c¯ZKvi‡Ki bvg I wVKvbv	JI‡ai bvg	†Rwbw¡K bvg	‡_‡iwcDwUK Kvm	wb‡`kbv	Contraindication & Side-effect	Status (New/Existing)	FSC/CPP	‡UKwbK¨vj mve Kvgv‡Ui mfvi wm×vš	mfvi wm×vš
7.	IZO S.r.l. a socio unico, S.S. 234 Km 28,2-27013 Chignolo Po (PV), ITALY **Local Agent:** Renata Ltd., Mirpur, Dhaka	**IZOVAC ND-EDS-IB** Emulsion for Injection for chickens	Inactivated Newcastle Disease Virus, strain La Sota:≥50 PD50 + Inactivated EDS '76 virus strain 127: as from phEu + Inactivated Infectious Bronchitis virus, strain M41, D274, D1466: as from	Vaccine	Prevention of Newcastle Disease, EDS '76 and Infectious Bronchitis in Breeders and Layers	**Contraindications:** There are not contra-indications. **Side Effects:** There are not side-effects.	New	Italy	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
8.	IZO S.r.l. a socio unico, S.S. 234 Km 28,2-27013 Chignolo Po (PV), ITALY **Local Agent:** Renata Ltd., Mirpur, Dhaka	**IZOVAC MG** Emulsified inactivated Vaccine against Mycoplasma gallisepticum	Inactivated cultures of *Mycoplasma gallisepticum* strain S-6:1.5x10^{10} CFU	Vaccine	Prevention of Mycoplasma gallisepticum infection	There are not contra-indications. There are not side-effects.	New	Italy	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
9.	LAPROVET Hungary Veterinary Pharmaceuticals Ltd. Hungary **Local Agent:** Navana Pharmaceuticals Ltd.	**ITA ND+IB+EDS+COR ABC (Inactivated) Vaccine**	Inactivated Newcastle disease virus, strain NDV-SZ LaSota….min PD50 + infectious bronchitis virus, strain M-41…..induced min.4.4log2HI + egg drop syndrome 76 virus, Strain B8/78…. induced min.7 log2 HI + Avibacterium paragallinarum, serotype A min. 7log10CFU + Avibacterium paragallinarum, serotype B min. 7log10CFU + Avibacterium paragallinarum, serotype C min. 7log10CFU	Vaccine	It is recommended for the vaccination of breeder and laying type chicken flocks, previously immunized against ND, IB, EDS and infectious Coryza, in order to prevent mortality, reduce clinical signs and lesions of the diseases.	**Contra-indications:** No contraindications are known **Side effects:** Vaccination does not cause systemic reactions at site of injection a swelling of pea size may develop; majority of cases this disappears In 2-3 weeks.	New	Hungary	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

bs	c¯ZKvi‡Ki bvg I wVKvbv	Jl‡ai bvg	†Rvbv‡iK bvg	‡_‡iwcDwUK Kvm	wb‡`kbv	Contraindication & Side-effect	Status (New/Existing)	FSC/CPP	‡UKwbK¯'vj mve Kvgv‡Ui mfvi wm×vš	mfvi wm×vš
10.	Lohmann Animal Health GmbH Heinz-Lohmann-Str. 4, D-27472 Cuxhaven Germany **Local Agent:** Elanco Bangladesh Limited 6 Kemal Ataturk Avenue, Banani, Dhaka-1213	AviPro ND C131 Lyophilisate for suspension Vaccine(Vet)	Live Newcastle Disease virus, strain Clone 131 ≥$10^{6.0}$ EID_{50}/Dose	Vaccine	For active immunization of chickens against Newcastle disease to reduce clinical signs, mortality, organ colonization and virus excretion. Onset of immunity: 1 week (3 weeks in MDA positive chickens) Duration of immunity: - broilers: at least 4 weeks after vaccination - layers/ breeders: at least 20 weeks after the application of the complete vaccine scheme performed accordingly with the recommended scheme described under point 4.9 (based on serological levels)	**Contra-indication:** Do not use in clinically ill or weakened animals. **Side Effects:** Transient mild respiratory reactions (coughing or sneezing) may be observed.	New	Germany	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
11.	Lohmann Animal Health GmbH Heinz-Lohmann-Str. 4, D-27472 Cuxhaven Germany **Local Agent:** Elanco Bangladesh Limited, 6 Kemal Ataturk Avenue, Banani, Dhaka-1213	AviPro IBD Xtreme (vet) Lyophilisate for suspension With Diluent Vaccine	Live Infectious Bursal Disease (IBD) Virus, Strain V217 ≥$10^{1.5}$ EID50/Dose	Vaccine	For active immunization of chickens to reduce clinical disease, weight loss, mortality and acute lesions of the bursa of Fabricius associated with infection caused by very virulent avian Infectious Bursal Disease (IBD) viruses. Onset of immunity: at least 1 week Duration of immunity in broilers: at least 4 weeks based on serology Duration of immunity in layers: at least 12 weeks	**Contra-indication:** Do not use in clinically ill or weakened animals **Side Effects:** On day 7 post-vaccination, severe lymphocyte depletion is seen in the bursae of the majority of birds. Lymphocyte repopulation commences 7 days after vaccination but notable lesions still remain in the bursae of the birds at day 28 post vaccination.	New	Germany	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

bs	c¯ZKvi‡Ki bvg I wVKvbv	JI‡ai bvg	†Rvbv‡iK bvg	‡_iwcDwUK Kvm	wb‡`kbv	Contraindication & Side-effect	Status (New/Existing)	FSC/CPP	‡UKwbK¨vj mve Kvgv‡Ui mfvi wm×vš	mfvi wm×vš
12.	Lohmann Animal Health GmbH Heinz-Lohmann-Str. 4, D-27472 Cuxhaven Germany **Local Agent:** Elanco Bangladesh Limited, 6 Kemal Ataturk Avenue, Banani, Dhaka-1213	AviPro Salmonella DUO Lyophilisate for suspension	Live attenuated Salmonella Enteritidis strain Sm24/Rif12/Ssq ≥ 1 x 10^8 CFU + Live attenuated Salmonella Typhimurium strain Nal2/Rif9/Rtt ≥ 1 x 10^8 CFU/ Dose	Vaccine	**Chickens (future breeders and future layer hens):** For active immunisation of chickens to reduce faecal excretion and colonisation of internal organs with *Salmonella* Enteritidis and *Salmonella* Typhimurium field strains and to reduce colonisation of eggs with *Salmonella* Enteritidis field strains. Onset of immunity: 15 days Duration of immunity after last vaccination: 52 weeks against *Salmonella* Enteritidis field strains and 46 weeks against *Salmonella* Typhimurium field strains. **Turkey breeders and turkeys for meat production:** For active immunisation of turkeys to reduce colonization of internal organs with *Salmonella* Enteritidis and *Salmonella* Typhimurium field strains. Onset of immunity: 21 days Duration of immunity after last vaccination: Future breeders: 30 weeks against *Salmonella* Enteritidis field strains and 28 weeks against *Salmonella* Typhimurium field strains. Turkeys for meat production: 10 weeks against *Salmonella* Enteritidis field strains and against *Salmonella* Typhimurium field strains. **Ducks for meat production:** For active immunisation of ducks to reduce colonisation of internal organs with *Salmonella* Enteritidis and Typhimurium field strains. Onset of immunity: 22 days Duration of immunity: 50 days	**Contra-indication:** Do not use in clinically ill or weakened animals. **Side Effects:** None known.	New	Germany	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

bs	c¯ZKvi‡Ki bvg I wVKvbv	Jl‡ai bvg	†Rvbv‡iK bvg	‡_‡iwcDwUK Kvm	wb‡`kbv	Contraindication & Side-effect	Status (New/Existing)	FSC/CPP	‡UKwbK¨vj mve Kvgv‡Ui mfvi wm×vš	mfvi wm×vš
13.	**Manufacturer :** SCI CO. Ltd. 593, Seonggok- RI,, Gyeolseong- Myeon, Hongseong- GUn, Chungcheongnam-DO, korea **Local agent:** M/S. rafique Medicine, College 55/B, Hazi zahir Bhaban, Sobahanbag, Saver, Dhaka.	**GIT Amino Acid**	Amino Acid, Fatty acid & minerals **Amino Acid (Per Kg)** Alanine 22,400 mg Arginine 17,100 mg Aspartam Acid 30,000 mg Cystein 4,400 mg Glutamic Acid 63,100 mg Glycine 24,400 mg Histidine 8,000 mg Isoleucine 15,100 mg Leucine 31,000 mg Lysine 19,000 mg Methionine 7,600 mg Phenylalanine 14,500 mg Proline 17,700 mg Serube 13,800 mg Threonine 14,200 mg Tryptophan 2,200 mg Tyrosine 11,000 mg Valine 20,000 mg **Fatty Acid (Per Kg)** Linoleic Acid (%) 2,216 Oleic Acid (%) 1,316 Palmitoletic Acid (%) 1,916 Steric Acid (%) 1,100 Satyrated Fatty Acid (%) 4,655 Unsanturated Fatty Acid (%) 5,495 Total Fatty Acid (%) 9,150 **Minerals (Per Kg)** Calcium 8,000 mg Phosphorus 9,000 mg Potassium 8,500 mg Sodium 8,800 mg Magnesium 800 mg Iron 10400 mg Manganese 20 mg Zinc 1300 mg Copper 3100 mg	**Amino Acid**	GIT is a product with a maximum absorption by hydrolizing the protein down to molecular level. Its absorption is 20 times as good as normal protein preparations. Containing chelated amino acid and fatty acid minerals, it shows excellent effects on growth, recovery and increased immunity	**Contraindication:** Have not been described **Side-effect :** Have not been described	New	Korea	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

bs	c¯ZKvi‡Ki bvg I wVKvbv	Jl‡ai bvg	†Rwbwi‡K bvg	‡_iwcDwUK Kvm	wb‡`kbv	Contraindication & Side-effect	Status (New/Existing)	FSC/CPP	‡UKwbK¨vj mve KwgwUi mfvi wm×vš	mfvi wm×vš
14.	Lohmann Animal Health International 375, China Road, winslow, USA **Local Agent:** C.P. bangladesh CO. Ltd.House-28 Alaol avenue, Sector-6, Uttara, Dhaka-1230	Avipro 105 ND chick Vaccine	Newcastle Disease virus vaccine, killed virus	Vaccine	AVIPRO® 105 ND-CHICK has been developed specifically for use in young chicks in geographical areas where highly pathogenic Newcastle disease is endemic. It is recommended that it be injected subcutaneously into one day old chicks and that it be used in conjunction with the Hitchner B1 strain of live virus Newcastle disease vaccine. Depending on the prevalence of Newcastle disease, this vaccination program may be used as late as 10 days of age.	**Contraindication :** Do not vaccinate 42 days before slaughter. **Side effects :** Hypersensitivity reactions may occur.	New	USA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
15.	Interchemice Werken de Adelaar B.V., Nederland **Local Agent:** C.P. bangladesh CO. Ltd.House-28 Alaol avenue, Sector-6, Uttara, Dhaka-1230	DOXIN -200 WS Powder	Doxycycline Hyclate Ph. Eu. 100mg + Tylosin Tartrate Ph. Eu. 100mg/gm	Antibiotic	Gastrointestinal and respiratory infections caused by tylosin and doxycycline sensitive microorganisms, like bordetella, Campylobacter, Chlamydia, E. coli, Haemophilus, mycoplasma, Pasteurella, Rickettsia, Salmonella, Staphylococcus, Streptococcus and treponema spp. in calves, goats, poultry, sheep and swine.	**Contraindication:** Hypersensitivity to tetracyclines and/or tylosin. administration to animals with a seriously impaired hepatic function. Concurrent adiministration of penicillins, cephalosporins, quinolones and cycloserine. Administration to animals with an microbial digestion. **Side effects:** Discoloration of teeth in young animals. Hypersensitivity reactions. Diarrhoea may occur.	New	The Netherlands	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
16.	Interchemice Werken de Adelaar B.V., Nederland **Local Agent:** C.P. bangladesh CO. Ltd.House-28 Alaol avenue, Sector-6, Uttara, Dhaka-1230	DIMOSAN WS powder	Colistin Sulphate Ph. Eu. 1,200,000IU + Amoxycillin Trihydrate Ph. Eu. 200mg/gm	Antibiotic	Gastrointestinal, respiratory and urinary tract infections caused by amoxicillin and colistin sesensitive micro-organisms, like Campylobacter, Clostridium, c\orynebacterium, E. coli, Erysipelothrix, Haemophilus, Pasteurella, Salmonella, penicillinase negative Staphylococcus and Streptococcus spp. in calves, goats poultry, Sheep and swine.	**Contraindication:** Hypersensitivity to amoxicillin and/or colistin. Administration to aminals with a seriously impaired renal function, Concurrent and lincosamides. Administration to animals with an microbial digestion. **Side effects :** Hypersensitivity reactions, renal dysfunction, neurotoxicity and neuromuscular blockade.		The Netherlands	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

bs	c¯ZKvi‡Ki bvg I wVKvbv	Jl‡ai bvg	†Rwbwi‡K bvg	‡_iwcDwUK Kvm	wb‡`kbv	Contraindication & Side-effect	Status (New/Existing)	FSC/CPP	‡UKwbK¨vj mve Kvgvwi mfvi wm×vš	mfvi wm×vš
17.	Interchemice Werken de Adelaar B.V., Nederland **Local Agent:** C.P. bangladesh CO. Ltd.House-28 Alaol avenue, Sector-6, Uttara, Dhaka-1230	COLOXAN WS Powder	Doxycycline Hyclate Ph. Eu. 100mg + Colistin Sulphate Ph. Eu. 1,200,000 IU/gm	Antibiotic	Gastrointestinal, respiratory and urinary tract infections caused by amoxicillin and colistin sesensitive micro-organisms, like Campylobacter, Clostridium, c\orynebacterium, E. coli, Erysipelothrix, Haemophilus, Pasteurella, Salmonella, penicillinase negative Staphylococcus and Streptococcus spp. in calves, goats poultry, Sheep and swine.	**Contraindication:** Hypersensitivity to amoxicillin and/or colistin. Administration to aminals with a seriously impaired renal function, Concurrent and lincosamides. Administration to animals with an microbial digestion. **Side effects:** It can cause discoloration of the teeth Hypersensitivity reactions, Changes in the digestion can occur as well as intestinal dysbiosis, accumulation of gases or diarrhea easier.	Colistin Sulphate 500000 IU + Doxycycline 100mg/gm	The Netherlands	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
18.	Interchemice Werken de Adelaar B.V., Nederland **Local Agent:** C.P. bangladesh CO. Ltd.House-28 Alaol avenue, Sector-6, Uttara, Dhaka-1230	INTROFLOR-100 Oral liquid	Florfenicol 100mg/ml	Antibiotic	Introflor-100 Oral liquid is indicated for preventive and therapeutic treatment of gastrointestinal and respiratory tract infections, caused by florfenicol sensitive micro- organisms such as actinobaccillusspp. pasteurella spp. Salmonella Spp. and Steptoccus spp. in swine and poultry. The presence of the disease in the herd should be established before preventive treatment. Medication should be initiated promptly when respiratory disease is diagnosed.	**Contraindication:** Not to be used in board intended for breeding purposes, or in animals producing egg or milk for human consumption. do not administer in cases of previous hypersensitivity to florfenicol the use of introflor 100 oral during pregnancy and lactation is not recommended. the product shuld not be used or stored in galvanized metal watering system or containers. **Side effects:** A decrease in food and water consumption and transient sofetening of the faeces or diarrhoea may occur during the treatment period. The treated animals recover quickly and competely upon terminiation of treatment. In swine, commonly observed adverse effects of the rectum. These effects are transient.	New	The Netherlands	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

bs	c¯ZKvi‡Ki bvg I wVKvbv	Jl‡ai bvg	†Rvbv‡iK bvg	‡_iwcDwUK Kvm	wb‡`kbv	Contraindication & Side-effect	Status (New/Existing)	FSC/CPP	‡UKwbK¨vj mve Kvgv‡Ui mfvi wm×vš	mfvi wm×vš
19.	**Manufacturer:** WooGene B&G Co., Ltd. 230, Jeongmunsongsan-ro, Yanggam-myeon, Hwaseong-si, Gyeonggi-do, Republic of korea. **Local Agent:** Cebac Animsal care, 23/1 Dilu Road, New Eskaton, Dhaka, Bangladesh.	**Wellclin-C** Liquid Disinfectant	Quaternary ammonium chloride as QAC (n-Alkyl dimenthyl Benzyl Ammonium Chloride, n- Alkyl Dimenthyl Ethylbenzyl Ammonium Chloride) 100gm + Citric Acid Hydrate 200gm/kg	*Disinfectant*	Disinfection against bacteria (Salmonella Typhimurium, Brucella Ovis) and Virus (FMDV, Avian Influenza)	**Contraindications:** Gastric Lavage, neutralization, activated charcol. **Side effects:** Not Known	new	Korea	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*
20.	**Manufacturer:** WooGene B&G Co., Ltd. 230, Jeongmunsongsan-ro, Yanggam-myeon, Hwaseong-si, Gyeonggi-do, Republic of korea. **Local Agent:** Cebac Animsal care, 23/1 Dilu Road, New Eskaton, Dhaka, Bangladesh.	Florject 400/ml Injection	Florfenicol 400mg/1ml	Antibiotic	It is indicated for preventive and therapeutic treatment of gastrointestinal and respiratory tract infections, caused by florfenicol sensitive micro- organisms such as actinobaccillusspp. pasteurella spp. Salmonella Spp. and Steptoccus spp. in swine and poultry. The presence of the disease in the herd should be established before preventive treatment. Medication should be initiated promptly when respiratory disease is diagnosed.	**Contraindication:** Not to be used in board intended for breeding purposes, or in animals producing egg or milk for human consumption. do not administer in cases of previous hypersensitivity to florfenicol the use of introflor 100 oral during pregnancy and lactation is not recommended. the product shuld not be used or stored in galvanized metal watering system or containers. **Side effects:** A decrease in food and water consumption and transient sofetening of the faeces or diarrhoea may occur during the treatment period. The treated animals recover quickly and competely upon terminiation of treatment. In swine, commonly observed adverse effects of the rectum. These effects are transient.	new	Korea	*Abyjgv`b Kiv †h‡Z cv‡i |*	*Abyjgv`b Kiv nj |*

Annex-D: Proposed Product for Local (Medical Devices)

Sl No	c¯ZKvi‡Ki bvg	ewYR¨K bvg	‡gvW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv/e¨envi	†UKwbK¨vj mve-Kwgw U mfvi wm×vš	mfvi wm×vš
1.	JMI hospital Requisite Mfg. Ltd., 7/A Shantibag, Dhaka-1217	JMI Sterile Disposable Scalpel	Scalpel handle (Acrynonitrile Butadine Styrine Medical grade 4.89gm) + Blade (Stainless steel Medical grade) + Protective cap (Liner low density polyethylene Medical grade 2.303gm)	B	Sterile Disposable Scalpel use for surgery, anatomical dissection in surgical operation. Also used to make small incision in skin and muscle.	c`wU wb¤ ewYZ k‡Z Abyjgv`b Kiv †h‡Z cv‡i: 1| ‡iwR‡ók‡bi c‡e c`wUi †mdwU WvUv `wLj Ki‡Z n‡e| 2| †iwR‡÷k‡bi c‡e ey‡qU KZK mwUwd‡KU MnY Ki‡Z n‡e|	c`wU wb¤ ewYZ k‡Z Abyjgv`b Kiv nj: 1| ‡iwR‡ók‡bi c‡e c`wUi †mdwU WvUv `wLj Ki‡Z n‡e| 2| †iwR‡÷k‡bi c‡e ey‡qU KZK mwUwd‡KU MnY Ki‡Z n‡e|
2.	JMI hospital Requisite Mfg. Ltd., 7/A Shantibag, Dhaka-1217	**JMI Sononned** Atraumitic Needle for peripheral nerve block	Cannula (Stainless steel Medical grade) + Tubing (Poly vinyl Chloride Medical grate 2.73gm) + Wings hub (Poly vinyl Chloride Medical grate 0.3gm) + Needle cap (Poly vinyl Chloride Medical grate 0.5gm) + Luer hub (Acrynonitrile Butadine Styrine Medical grade 0.238gm) + Thread Stopper (High Density polyethylene Medical grade 0.35gm + Master Batch Medical grade 0.014gm) + Adhesive (Epoxy resin Medical grade 0.001ml + Epoxy Hardener Medical Grade 0.001ml) + Cylohexanone Medical Grade 0.01ml + Siliconization (Silicoat Medical grade 0.01ml + n-hexana Medical grade 0.01ml)	B	Sononned used for facilitating a principal nerve block procedure. The needle has several fenestration along it's length to allow and efflux of local anesthetic into particular facial compartment to treat a corresponding peripheral nerve.	c`wU wb¤ ewYZ k‡Z Abyjgv`b Kiv †h‡Z cv‡i: 1| ‡iwR‡ók‡bi c‡e c`wUi †mdwU WvUv `wLj Ki‡Z n‡e| 2| †iwR‡÷k‡bi c‡e ey‡qU KZK mwUwd‡KU MnY Ki‡Z n‡e|	c`wU wb¤ ewYZ k‡Z Abyjgv`b Kiv nj: 1| ‡iwR‡ók‡bi c‡e c`wUi †mdwU WvUv `wLj Ki‡Z n‡e| 2| †iwR‡÷k‡bi c‡e ey‡qU KZK mwUwd‡KU MnY Ki‡Z n‡e|
3.	JMI hospital Requisite Mfg. Ltd., 7/A Shantibag, Dhaka-1217	JMI Alcohol prer pads	Non Woven Fabric + Isopropyl Alcohol (70% v/v) (Medical Grade 0.335gm)	A	It is used for antiseptic skin preparation prior to injection or venipuncture.	Abyjgv`b Kiv †h‡Z cv‡i|	Abyjgv`b Kiv nj|

Annex-E: Proposed Product for Import (Medical Devices)

Sl No	c¯ZKvi‡Ki bvg	ewYR¨K bvg	‡gvW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve KwgwUi mfvi wm×vš	mfvi wm×vš
4.	**Manufacturer:** Smith & Nephew Medical Limited, UK **Local Agent:** UniMed Ltd., House No. 46, Sheikh Kamal Saroni, Road No. 16, Rangs Nasim (7th Floor) Square, Dhanmondi	Allevyn Gentle Border	Dressing	C	ALLEVYN Gentle Border dressings are indicated for use on: ➢ Shallow, granulating wounds ➢ Chronic and acute exudative wounds ➢ Full and partial thickness wounds such as pressure ulcers, leg ulcers and diabetic foot ulcers ➢ Infected wounds ➢ Malignant wounds ➢ Surgical wounds ➢ First and second degree burns ➢ Donor sites ➢ Oncological wounds ➢ Epidermolysis bullosa wounds	Contraindications: Not known Side Effects: Not known	MHRA-UK	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
5.	**Manufacturer:** Bioelectronics CORPORATION, USA **Local Agent:** Multiple Health Pharma Ltd, 33 kawran Bazar, Shah Ali Tower (11th Floor), Dhaka-1212	**ActiPatch**	Electromagnetic Therapy (Muskoskeletal Pain relief)	**B**	ActiPatch provides advanced long-lasting chronic pain relief using Electromagnetic Pulse Therapy and works great for back pain, knee pain, muscle & joint pain, arthritis, sciatica, fibromyalgia, strains, sprains and more. ActiPatch® is drug and ingredient-free so is safe for continuous use and can be used while taking any other medication. Unlike a TENS machine, you will not feel heat or vibration. In fact, ActiPatch® is completely sensation free, so the only thing you are going to feel is better.	**Contraindications:** None **Side Effects:** None **Hazards and Cautions:** ActiPatch should not be used by persons with a cardiac pacemaker, cardioverter defibillator, neurostimulator or any active medical or metallic implant that is within the area of application. ActiPatch should not be used by women who are pregnant or think they may be pregnant. See product insert for more information. Can be used in conjunction with medications.	USA EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
6.	**Manufacturer:** Alcon Laboratories Inc, USA **Local Agent:** Globex Marketing Company Ltd. RH Home Center (6th Floor) 74/B/1, Green Road, Dhaka-1215	Alcon Cutting Instruments	Ophthalmic Knives and Blades	**B**	It is used to vitreoretinal surgery of human eye.	Contraindication: Not Known Side Effects: Not Known	USA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

Sl No	c¯ZKvi‡Ki bvg	ewYwR¨K bvg	‡gwW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve Kwgkb mfvi wm×vš	mfvi wm×vš
7.	**Manufacturer:** BioElectronics Corporation, USA **Local Agent:** Multiple Health Pharma Ltd, 33 kawran Bazar, Shah Ali Tower (11th Floor), Dhaka-1212	**ALLAY**	Electromagnetic Therapy (Menstrual Pain relief)	**B**	Allay® is drug-free micro medical device that uses Electromagnetic Pulse Therapy to reduce menstrual pain and discomfort. Allay® is a wafer-thin, discreet, lightweight, and comfortable and is designed to provide 24/7 continual use throughout your entire period.	**Contraindications:** None **Side Effects:** None	USA EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj	
8.	**Manufacturer:** Becton Dickinson and Company, UK **Branch Office:** Becton Dickinson India Pvt. Ltd.80, Kakrail Dhaka	BD A-line Arterial Blood collection Syringe	Arterial Blood Gas Syringe	**B**	Arterial Blood Gas syringe is sterile, single use, medical devices specifically intended to be used for the collection, primary containment and preservation of blood specimens derived from the human body for the purposes of in Vitro diagnostic Examination. It is also used to collect blood either by aspiration or by pre-setting to a desired volume and allowing the syringe to fill under arterial blood pressure. The syringes contain calcium balanced lithium heparin to prevent clotting of the sample.	Contraindication: Not Known Side Effects: Not Known	MHRA-UK EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj	
9.	**Manufacturer:** GE Medical Systems Information Technologies, Inc. 8200 West Tower Avenue, Milwaukee, WI 53223 USA **Agent**: GE Healthcare Bangladesh Ltd. Shanta Western Tower, OS # 804, Level # 08, 186 Tejgaon I/ A Dhaka	**CardioSoft** Cardiac Testing System	Cardiac Testing System	B (P-141)	CardioSoft is designed to acquire, process, record, archive, analyzeand output ECG data (12 and 15 leads) during a period of physiologic stress or during a resting ECG test and acquire data from ancillary devices, such as spirometry and ambulatory blood pressure devices. Furthermore, it provides median morphology recordings and records ECG in real-time with and without arrhythmia detection. The arrhythmia detection algorithm of CardioSoft is provided to the user for the convenience of automatic detection of arrhythmias but does not provide alarms. CardioSoft provides the control of external devices. These are typically treadmills and bicycle ergometers. Additionally, CardioSoft communicates with centralized electronic/digital storage systems via data networks.	**Contra-Indications**: None. **Side Effect**: None.	USA EC ECtificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj	

Sl No	c¯ZKvi‡Ki bvg	ewYR¨K bvg	‡gvW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve KwgwUi mfvi wm×vš	mfvi wm×vš
10.	**Manufacturer:** GE Medical Systems Information Technologies, Inc. 8200 West Tower Avenue, Milwaukee, WI 53223 USA **Local Agent**: GE Healthcare Bangladesh Ltd. Shanta Western Tower, OS # 804, Level # 08, 186 Tejgaon I/ A Dhaka	**CASE** Cardiac Testing System	Cardiac Testing System	**B** (P-29)	CASE Cardiac Testing System is intended to be used by trained operators under direct supervision of alicensed health care practitioner on adult and pediatric patients. The CASE Cardiac Testing System isdesigned to acquire process, record, archive, analyze and output (12 and 15 lead) ECG data during a period ofphysiologic stress or during a resting ECG test, acquire data from ancillary devices (such as Spirometry andAmbulatory Blood Pressure), provide median morphology recordings and record ECG in real-time with and withoutarrhythmia detection. The arrhythmia detection portion of CASE Cardiac Testing System is provided to theuser for the convenience of automatic detection of arrhythmias but does not provide alarms. CASE Cardiac Testing System provides the control of external device (typically a treadmill or Ergometer)and communicates with centralized electronic/digital storage system via network. CASE Cardiac TestingSystem provides a user selectable option for printouts of prognostic scores on select reports. Vector loops are alsoavailable. CASE Cardiac Testing System can be configured in a network environment for multiple CASE stations allowing the user to create a central database of patient demographics and collected patientphysiological data. CASE Cardiac Testing System is intended to be used primarily in the hospital but can be used in clinics, physician offices, outreach centers or wherever exercise, stress testing, ECG, Spirometry or ambulatory blood pressuretesting is performed. CASE Cardiac Testing System offers no diagnostic opinion to the user. Instead it provides interpretivestatements of morphology, rhythm, and conduction for which the physician renders his/her own medical opinion.CASE Cardiac Testing System is not intended to be used as a transport device or for home use. CASE Cardiac Testing System is not intended for the use as a vital signs physiological monitor or forintracardiac use.	**Contraindication:** None, as identified in the Clinical Evaluation Summary and User Manual. **Side effect:** None	USA EC ECtificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

Sl No	c¯ZKvi‡Ki bvg	ewYR¨K bvg	‡gvW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve Kwgkb Ui mfvi wm×vš	mfvi wm×vš
11.	**Manufacturer:** Echosens, Paris, France **Local Distributor:** KNS Canada (BD), 42, West Tejturi Bazar, KaziNazrul Islam Avenue, 4th Floor, Dhaka-1215	FibroScan	Ultrasound Diagnostic Device & associated probes	**B**	Examination with FibroScan is also called transient elastography which is a technique used to asses Liver Stiffness (measured in kPa correlated to Fibrosis) and Controlled Attenuation Parameter CAP (measured in dB correlated to steatosis) without invasive investigation. The result is immediate. It shows the condition of the liver and allows physicians to diagnose and monitor disease evaluation in conjunction with treatment and collateral factors.	**Contraindication:** The FibroScan examination should not be used on: any organ other than the liver, wounds, patients with ascites. **Side-effects:** None	France EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
12.	**Manufacturer:** Fujinomiya Factory of Terumo Corporation, Japan **Local Agent :** UniMed Ltd., House-46, Sheikh Kamal Soroni, Road-16, Rans Nasim Sqaure, Dhanmondi, Dhaka	IMUGARD III-PL	Blood Transfusion Filter	B (P-114)	Leukocyte removal filter for Platelet Preparation	**Contraindication:** None **Side Effects:** None	Japan	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
13.	**Manufacturer:** Fujinomiya Factory of Terumo Corporation, Japan **Local Agent :** UniMed Ltd., Dhaka	IMUGARD III-RC	Blood Transfusion Filter	B (P-114)	Leukocyte removal filter for Red blood Cell Preparation	**Contraindication:** None **Side Effects:** None	Japan	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
14.	**Manufacturer:** Kehr Surgical Pvt. Ltd. Kanpur, India **Local Agent:** JMI Hospital Requisite Mfg. Ltd. 7/A, Shantibag, Dhaka-1217	JMS Brand CS Sterile Surgical Blades	Blades	B (P-58)	Used for orthopaedic surgery	**Contraindications:** Not Known **Side Effects:** Not Known	India	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

Sl No	c¯ZKvi‡Ki bvg	ewYRK bvg	‡gvW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve KwgwUi mfvi wm×vš	mfvi wm×vš
15.	**Manufacturer:** Kehr Surgical Pvt. Ltd. Kanpur, India **Local Agent:** JMI Hospital Requisite Mfg. Ltd. 7/A, Shantibag, Dhaka-1217	JMS Brand SS Sterile Disposable Scalples	Scalples	B (P-58)	Used for orthopaedic surgery	**Contraindications:** Not Known **Side Effects:** Not Known	India	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj	
16.	**Manufacturer:** Romsons Scientific & Surgical Industries Pvt. Ltd., India **Local Agent:** Barisal Surgical Rizia Manson, 34/1, Mitford Road, Dhaka	LARYNGEAL MASK AIRWAYS-EXCELL	Laryngeal Mask	**B**	Laryngeal Mask Airway –Excell is sterile, single use device intended to provide elective ventilation in the operating room & also for use in emergency as an important accessory device for management of difficult airway Laryngeal Mask Airways Excell is a shaped like a larged endotracheal tube on the proximal end that connects to an elliptical mask on the distal end. It is designed to sit in the patients hypopharynx & cover the supraglottic structures, thereby allowing relative isolation of the trachea Laryngeal Mask Airways Excell is an acceptable alternative to mask anesthesia in the operating room. it is often used for short procedures, when endotracheal intubation is not necessary.	No side effects	India EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj	
17.	**Manufacturer:** Wipro GE healthcare Private Ltd., India **Local Agent**: GE Healthcare Bangladesh Ltd. Shanta Western Tower, OS # 804, Level # 08, 186 Tejgaon I/ A Dhaka	Lullaby LED Phototherapy	LED Phototherapy	B	It is used for the treatment of indirect hyperbilirubinemia in term and pre-term infants, in a hospital environment - NICUs, PICUs and Well-baby Nurseries - administered by trained, professional medical staff, on the order of a licensed medical practitioner. The Lullaby™LED Phototherapy System is intended for use under the direct supervision of a licensed healthcare practitioner. The Lullaby™LED Phototherapy System device is not intended to be operated in mobile vehicles including ambulances or other vehicles associated with health care facilities.	**Contraindication:** None. **Side effect:** None	India EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj	

Sl No	c¯ZKvi‡Ki bvg	ewYR¨K bvg	‡gvW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve Kwgkb mfvi wm×vš	mfvi wm×vš
18.	**Manufacturer:** Wipro GE healthcare Private Ltd., India **Local Agent**: GE Healthcare Bangladesh Ltd. Shanta Western Tower, OS # 804, Level # 08, 186 Tejgaon I/ A Dhaka	Lullaby Warmer Prime	Infant Warmer	B	Infant warmers provide heat in a controlled manner to infants who are unable to maintain thermoregulation based on their own physiology. Infant warmers may be used to facilitatethe newborn's transition to the external environment or to provide a controlled open microenvironment	**Contraindication:** None. **Side effect:** None	India EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
19.	**Manufacturer:** Boston Scientific Corporation, 300 Boston Scientific Way, Marlborough, MA 01752, USA **Local Agent:** M/s. Medi Graphic Trading Ltd., 14 Purana Paltan, Dhaka, Bangladesh	Malecot nephrostomy Catheter set	Nephrostomy Catheters and Catheter set	B	It is intended to establish percutaneous nephrostomy drainage and is indicated for the following: Diagnostic Indications: Antegrade pyelography, Pressure/perfusion study (Whitaker test), Therapeutic Indications: Nephrostomy catheter drainage, Perfusion chemolysis of renal stones, Percutaneous nephrolithotomy, Percutaneous resection and coagulation of urothelial tumors	**Contraindications:** The Malecot Nephrostomy Catheter Set is not indicated for use in patients with blood clotting anomalies due to coagulopathies or pharmacological anticoagulations. **Adverse Event:** Hemorrhage, Subcapsular renal hematoma, Bleeding from iatrogenic arteriovenous-caliceal fistulas, Edema, Extravasation, Loss of renal function, Sepsis	USA EC Certificate	**Abyjgv`b Kiv †h‡Z cv‡i**	Abyjgv`b Kiv nj
20.	**Manufacturer:** GE Medical Systems Information Technologies, Inc. 8200 West Tower Avenue, Milwaukee, WI 53223 USA **Local agent:** GE Healthcare Bangladesh Ltd. Shanta Western Tower, OS # 804, Level # 08, 186 Tejgaon I/ A Dhaka	**MUSE** Cardiology Information System Application Software	Cardiology Information System Application Software	**B** (P-141)	The MUSE Cardiology Information System is intended to store, access and manage cardiovascular information on adult and pediatric patients. The information consists of measurements, text, and digitized waveforms. The MUSE Cardiology Information System provides the ability to review and edit electrocardiographic procedures on screen, through the use of reviewing, measuring, and editing tools including ECG serial comparison and interpretive 12-lead analysis. The MUSE Cardiology Information System is intended to be used under the direct supervision of a licensed healthcare practitioner, by trained operators in a hospital or facility providing patient care. The MUSE Cardiology Information System is not intended for real-time patient monitoring. The MUSE Cardiology Information System is not intended for pediatric serial comparison.	**Contraindication:** The MUSE is not intended for primary monitoring, pediatric serial comparison, real time patient monitoring or for the transfer of time-sensitive data. **Side effect:** None	USA EC Certificate	**Abyjgv`b Kiv †h‡Z cv‡i**	Abyjgv`b Kiv nj

Sl No	c¯ZKvi‡Ki bvg	ewYRwK bvg	‡gvW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve Kwgkb mfvi wm×vš	mfvi wm×vš
21.	**Manufacturer:** Susruta Instruments Pvt. Ltd., India **Local Agent:** Access Medical Services R.H Home Center 74/B/1, Green Road Dhaka-1215	Ophthalmic Microsurgical Knives, Blades	Ophthalmic Microsurgical Knives, Blades	B (P-119)	It is used for Eye Surgery	**Contraindication:** Not Known. **Side Effects:** None	India EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj	
22.	**Manufacturer:** Haemonetics Malaysia Sdn. Bhd., Malaysia **Local Agent:** MBioLogix 205, Shahid Syed Nazrul Islam Swarani Bijoy Nagar (3rd Floor) Dhaka-1000	PLT & PLS SetW/Filter	Platelet & Plasma Therapeutics	B (P-114)	Closed set for automated Platelet collection Plasma collection is optional. Platelet Bags allows 7 days storage. Leukodeplation filter for continuos filtration during procedure and it is one ar procedure.	**Contraindication:** None **Side Effects:** None	Swiszerland EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj	
23.	**Manufacturer:** Romsons Scientific & Surgical Industries Pvt. Ltd., India **Local Agent:** Barisal Surgical Rizia Manson, 34/1, Mitford Road, Dhaka	Romo ADK	Abdominal Drainage Kit	B	Intended for management of the post-operative abdominal drainage	No side effects	India EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj	
24.	**Manufacturer:** Romsons Scientific & Surgical Industries Pvt. Ltd., India **Local Agent:** Barisal Surgical Rizia Manson, 34/1, Mitford Road, Dhaka	Romsons ECG Electrodes	ECG Electrodes	B	Single Use Disposable ECG electrodes for recording & Monitoring ECG.	No side effects	India	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj	

Sl No	c¯ZKv‡i Kiv bvg	ewYR¨K bvg	‡gvW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve Kwgkb mfvi wm×vš	mfvi wm×vš
25.	**Manufacturer:** Romsons Scientific & Surgical Industries Pvt. Ltd., India **Local Agent:** Barisal Surgical Rizia Manson, 34/1, Mitford Road, Dhaka	Romsons Nasopharyngeal Airways	Nasopharyngeal Airways	B	Nasopharyngeal Airways are suitable for use as an alternative when the oropharyngeal airways are occluded.	No side effects	India EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj	
26.	**Manufacturer:** GE Healthcare (Tianjin) Company Limited No.266 Jingsan Road Tianjin Airport Economic Area,Tianjin, P.R. China 300308 **Local Agent**: GE Healthcare Bangladesh Ltd. Shanta Western Tower, OS # 804, Level # 08, 186 Tejgaon I/ A Dhaka-1208	**SIGNA Creator** (8 Channel) **/ SIGNA Explorer** (16 Channel)	Nuclear Magnetic Resonance Imaging System	**B** (P-102)	The systems are whole body magnetic resonance scanner designed to support high resolution, high signal-to-noise ratio, and short scan times. It is indicated for use as a diagnostic imaging device to produce axial, sagittal, coronal, and oblique images, spectroscopic images, parametric maps, and/or spectra, dynamic images of the structures and/or functions of the entire body, including, but not limited to, head, neck, TMJ, spine, breast, heart, abdomen, pelvis, joints, prostate, blood vessels, and musculoskeletal regions of the body. Depending on the region of interest being imaged, contrast agents may be used. The images produced by the system reflect the spatial distribution or molecular environment of nuclei exhibiting magnetic resonance. These images and/or spectra when interpreted by a trained physician yield information that may assist in diagnosis.	**Contraindication:** The use of MR systems is contraindicated (i.e., not advised) for patient and MR workers with any ofthe following, since the magnetic and electromagnetic fields produced by the MR system mayinterfere with the operation of these devices: electrically, magnetically, or mechanically activated implants (e.g., cardiac pacemakers and ferrous/electrically activated cardiac catheters); Intracranial aneurysm clips. In general, patients with conductive (e.g. metallic) implants are contraindicated for MR scans. Some implantable devices have been labeled as MR Conditional under certain operating conditions. MR Safe implants will have the MR Safe symbol in their implant documentation. When evaluating whether to proceed with MR scanning on patients with such implants, consultthe implantable device's labeling. **MR Safe**: For patients with implants that are labeled as MR Safe, consult the implantable device'slabeling. **MR Conditional**: For patients with implants that are labeled as MR Conditional, consult theimplantable device's labeling. **MR Unsafe**: Patients with implantable devices that are MR Unsafe are contraindicated. If the levelof MR compatibility is not known, then an implantable device should be considered MRUnsafe. **Side effect:** None	USA EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj	

Sl No	c¯ZKvi‡Ki bvg	ewYRK bvg	‡gvW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve KwgwUi mfvi wm×vš	mfvi wm×vš
27.	**Manufacturer:** GE Medical Systems Information Technologies, Inc. 8200 West Tower Avenue Milwaukee, WI 53223 USA **Local Agent**: GE Healthcare Bangladesh Ltd. Shanta Western Tower, OS # 804, Level # 08, 186 Tejgaon I/ A Dhaka-1208	**T2100 Treadmill**	Treadmill	**B** (P-89)	It is intended for use with any of the several GE Medical Systems Information Technologies exercise testing systems, for administering a controlled exercise load during a diagnostic stress test. It is intended to be used primarily in the hospital but can be used in clinics, physician offices, outreach centers or wherever exercise stress testing with a treadmill is performed. The T2100 Treadmill is not suitable for use in domestic establishments and those directly connected to the public low-voltage power supply network that supplies buildings used for domestic purposes. It is intended for use by qualified medical personnel who have received proper product training. It will be used typically for 120 diagnostic stress tests per week.	**Contraindication:** None. **Side effect:** None	USA EC Certificate	Abyôgv`b Kiv †h‡Z cv‡i	Abyôgv`b Kiv nj|
28.	**Manufacturer:** Haemonetics Malaysia Sdn. Bhd., Malaysia **Local Agent:** MBioLogix 205, Shahid Syed Nazrul Islam Swarani Bijoy Nagar (3rd Floor) Dhaka-1000	Therapeutic Plasma Exchange Sets (TPE)	Plasma Therapeutics	B (P-114)	The TPE disposable set is used to collect plasma from a patient and replace it with fluid, such as Albumin, FFp or balanced electrolyte solution Replacement solution administration can be machine controlled or optionally operator control through a second venous access.	**Contraindication:** None **Side Effects:** None	Swiszerland EC Certificate	Abyôgv`b Kiv †h‡Z cv‡i	Abyôgv`b Kiv nj|
29.	Romsons International (Unit-II), India **Local Agent:** Barisal Surgical Rizia Mansion 34/1 Mitford Road Dhaka-1100	Male Cath	Condom Catheter	B	Male Cath is a sterile single use Male Comdom catheter (External or Penile Sheath intended for management of urine incontinence.	No side effects	India Ec Certificate	Abyôgv`b Kiv †h‡Z cv‡i	Abyôgv`b Kiv nj|

Sl No	c¯ZKvi‡Ki bvg	ewYwR¨K bvg	‡gvW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve Kwgkb Ui mfvi wm×vš	mfvi wm×vš
30.	Datex-Ohmeda, Inc. 3030 Ohmeda Drive PO Box 7550 Madison, WI 53707-7550, USA **Local agent:** GE Healthcare Bangladesh Ltd., Shanta Western Tower, 186, Tejgaon. Dhaka	Carestation 620/650/650c (**Three Gas system:** Oxygen, Air, Nitrous Oxide)	GAS-Machine, Anesthesia	**C** (P-29)	The Carestation 620/650/650canesthesia systems are intended toprovide general inhalation anesthesiaand ventilatory support to a wide range of patients (neonate, pediatric, andadult). The anesthesia systems aresuitable for use in a patientenvironment, such as hospitals, surgicalcenters, or clinics. The systems areintended to be operated by a clinicianqualified in the administration ofgeneral anesthesia.	**Contraindication:** None. **Side effect:** None	USA EC Certificate	Abygv`b Kiv †h‡Z cv‡i |	Abygv`b Kiv nj |
31.	**Manufacturer :** Boston Scientific Limited, Ballybrit Business Park, Galaway, Ireland **Legal Manufacturer:** Boston Scientific Corporation, USA **Local Agent :** Vastech Limited, Nurjehan Tower (6th Floor), 80/22 Mymensingh Road Dhaka	**2D Helical**	Fibered Platinum Coil	C	2D Helical - 35 Fibered Platinum Coils are intended for arterial and venous embolizations in the peripheral vasculature.	**Contraindications:** None known. **Adverse Events:** The complications that may result from a peripheral embolization procedure include, but are not limited to: • Complications related to catheterization (e.g., hematoma at the site of entry, vessel injury, etc.) Death, Emboli, Foreign body reactions necessitating medical intervention, Hemorrhage, Infection necessitating medical intervention, Ischemia, Pain, Recanalization, Temporary neurological deficit, Tissue, necrosis, Undesirable clot formation of the vasculature, Vasospasm	Ireland EC Certificate	Abygv`b Kiv †h‡Z cv‡i |	Abygv`b Kiv nj |
32.	**Manufacturer :** Boston Scientific Limited, Ballybrit Business Park, Galaway, Ireland **Legal Manufacturer:** Boston Scientific Corporation, USA **Local Agent :** Vastech Limited, Nurjehan Tower (6th Floor), 80/22 Mymensingh Road Dhaka	**Complex Helical**	Fibered Platinum Coil	C	Boston Scientific's 0.018 Fibered Platinum Coils are intended for arterial and venous embolizations in the peripheral vasculature. The Coil Pusher-16 is intended to be used in conjunction with a microcatheter to deliver and deploy 0.018 pushable occlusion coils.	**CONTRAINDICAT IONS:** None known. **ADVERSE EVENTS:** The complications that may result from a peripheral embolization procedure include, but are not limited to: • Complications related to catheterization (e.g., hematoma at the site of entry, vessel injury, etc.) Death, Emboli, Foreign body, reactions necessitating medical intervention, Hemorrhage, Infection necessitating medical intervention, Ischemia, Pain, Recanalization, Temporary neurological deficit, Tissue, necrosis, Undesirable clot formation of the vasculature, Vasospasm	Ireland EC Certificate	Abygv`b Kiv †h‡Z cv‡i |	Abygv`b Kiv nj |

Sl No	c¯ZKvi‡Ki bvg	ewYwR¨K bvg	‡gvW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve KwgwUi mfvi wm×vš	mfvi wm×vš
33.	**Manufacturer :** Boston Scientific Limited, Ballybrit Business Park, Galaway, Ireland **Legal Manufacturer:** Boston Scientific Corporation, USA **Local Agent :** Vastech Limited, Nurjehan Tower (6th Floor), 80/22 Mymensingh Road Dhaka	**Direxion**	Torqueable Microcatheter	C	The Direxion and Direxion HI-FLO Torqueable Microcatheters are intended for peripheral vascular use. The pre-loaded Fathom and Transend Guidewires can be used to selectively introduce and position the microcatheter in the peripheral vasculature. The microcatheter can be used for controlled and selective infusion of diagnostic, embolic, or therapeutic materials into the vessel.	**Contraindications:** None known. **Adverse events:** The Adverse Events include, but are not limited to: Allergic reaction, Death, Embolism, Hemorrhage/Hematoma, Infection, Pseudoaneurysm, Stroke, Vascular, thrombosis, Vessel occlusion, Vessel spasm, Vessel trauma (dissection, perforation, rupture)	Ireland EC Certificate	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj	
34.	**Manufacturer :** Boston Scientific Limited, Ballybrit Business Park, Galaway, Ireland **Legal Manufacturer:** Boston Scientific Corporation, USA **Local Agent :** Vastech Limited, Nurjehan Tower (6th Floor), 80/22 Mymensingh Road Dhaka	Epic Over-the Wire Self-Expanding Nitinol Stent with Delivery System	Peripheral Artery Stent	C	It is indicated for the treatment of peripheral vascular lesions and obstructions.	**Contraindication:** Generally, contraindications for percutaneous transluminal angioplasty (PTA) are also contraindications for stent placement. Contraindications for the use of the Epic Nitinol Vascular Stent System include, but may not be limited to: • Patients with highly calcified lesions resistant to PTA. • Patients with a target lesion with a large amount of adjacent acute or subacute thrombus. • Patients with uncorrected bleeding disorders or patients who cannot receive anticoagulation or antiplatelet aggregation therapy. • Patients with perforated vessels evidenced by extravasation of contrast media. • A lesion that is within or adjacent to an aneurysm. • Patients with excessive vessel tortuosity. **Adverse events:** Potential adverse events that may occur following intravascular stent implantation include, but are not limited to: Abscess, Allergic reaction (to drug, contrast, device or other), Amputation, Aneurysm, Angina/coronary ischemia, Arrhythmia, Arteriovenous fistula, Death, Drug reactions, Embolization (air, plaque, thrombus, device, or other), Entanglement of delivery system indeployed stent, Fever, GI bleeding, Hemorrhage/hematoma, Hypotension/hypertension, Myocardial Infarction (MI), Need for urgent intervention or surgery, Pseudoaneurysm, Renal insufficiency or failure, Restenosis of stented artery, Sepsis/infection, Stent fracture, Stent migration, Stent misplacement/jumping, Stent thrombosis with possible neurological injury, Stroke, Thrombosis/thrombus,Tissue ischemia/necrosis, Vasospasm, Vessel injury, examples include perforation, dissection, intimal tear, rupture, Vessel occlusion	Ireland EC Certificate	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj	

Sl No	c¯ZKvi‡Ki bvg	ewYRK bvg	‡gvW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve Kwgkb mfvi wm×vš	mfvi wm×vš
35.	**Manufacturer :** Boston Scientific Limited, Ballybrit Business Park, Galaway, Ireland **Legal Manufacturer:** Boston Scientific Corporation, USA **Local Agent :** Vastech Limited, Nurjehan Tower (6th Floor), 80/22 Mymensingh Road Dhaka	FATHOM Steerable Guidewire	Guidewire	C	The FATHOM -16 Steerable Guidewire is intended for general intravascular use in the peripheral vasculature. It can be used to selectively introduce and position catheters and other interventional devices within the peripheral vasculature. This device should be used only by physicians trained in percutaneous, intravascular techniques and procedures.	**Contraindication:** Not Known. **Adverse events:** Complications attributed to endovascular procedures are the following: Vessel trauma, Vessel damage, Embolism (catheter/device, air bubble, plaque, thrombus, air embolism, thromboembolism), Pseudoaneurysm, Seizure/stroke, Vessel dissection, Hematoma at the puncture site, Nerve injury, Infection, Perforation of the vessel, Vessel spasm, Hemorrhage, Vascular thrombosis, Vessel occlusion, Death, Bleeding, Failed treatment, Inability to position guidewire, Damage to the catheter	Ireland EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj|
36.	**Manufacturer :** Boston Scientific Limited, Ballybrit Business Park, Galaway, Ireland **Legal Manufacturer:** Boston Scientific Corporation, USA **Local Agent :** Vastech Limited, Nurjehan Tower (6th Floor), 80/22 Mymensingh Road Dhaka	Innova Self-Expanding Stent System	Peripheral Artery Stent	C	The Innova Self-Expanding Stent System is indicated for the treatment of peripheral vascular lesions.	**Contraindication:** The Innova Self-Expanding Stent System is contraindicated for use in any situation in which percutaneous transluminal angioplasty (PTA) is contraindicated and also, but not limited to: • Patients with highly calcified lesions resistant to PTA. • Patients with a persistent, intraluminal thrombus at the target lesion. • Patients with uncorrected bleeding disorders or patients who cannot receive anticoagulation or antiplatelet aggregation therapy. • Patients with perforated vessels evidenced by extravasation of contrast media. • A lesion that is within or adjacent to an aneurysm. • Patients with excessive vessel tortuosity. • Patients that present with a lesion that cannot be completely dilated with an angioplasty balloon or do not allow for proper placement of a stent. **Adverse events:** Potential adverse events that may occur following intravascular stent implantation include, but are not limited to: Abscess, Allergic reaction (to drug, contrast, device or other), Amputation, Aneurysm, Angina/coronary ischemia, Arrhythmia, Arteriovenous fistula, Death, Drug reactions, Embolization (air, plaque, thrombus, device, or other), Entanglement of delivery system indeployed stent, Fever, GI bleeding, Hemorrhage/hematoma, Hypotension/hypertension, Myocardial Infarction (MI), Need for urgent intervention or surgery, Pseudoaneurysm, Renal insufficiency or failure, Restenosis of stented artery, Sepsis/infection, Stent fracture, Stent migration, Stent misplacement/jumping, Stent thrombosis with possible neurological injury, Stroke, Thrombosis/thrombus,Tissue ischemia/necrosis, Vasospasm, Vessel injury, examples include perforation, dissection, intimal tear, rupture, Vessel occlusion	Ireland EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj|

Sl No	c¯ZKvi‡Ki bvg	ewYRwRK bvg	‡gvW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve Kwgkb mfvi wm×vš	mfvi wm×vš
37.	**Manufacturer :** Boston Scientific Limited, Ballybrit Business Park, Galaway, Ireland **Legal Manufacturer:** Boston Scientific Corporation, USA **Local Agent :** Vastech Limited, Nurjehan Tower (6th Floor), 80/22 Mymensingh Road Dhaka	**Renegade**	Fiber Braided Microcatheter	C	The Renegade Fiber Braided Microcatheter is intended for peripheral vascular use. The microcatheter can be coaxially tracked over a steerable guidewire in order to access distal, tortuous vasculature. Once the subselective region has been accessed, the microcatheter can be used for the controlled and selective infusion of diagnostic, embolic, or therapeutic materials into vessels. Diagnostic, embolic, or therapeutic agents to be used in accordance with specifications outlined by the manufacturer.	**CONTRAINDICAT IONS:** None Known. **ADVERSE EVENTS:** The Adverse Events include, but are not limited to: • Vessel trauma • Embolism • Hemorrhage/Hematoma • Vasospasm • Infection • Air embolism • Allergic reaction	Ireland EC Certificate	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
38.	**Manufacturer :** Boston Scientific Limited, Ballybrit Business Park, Galaway, Ireland **Legal Manufacturer:** Boston Scientific Corporation, USA **Local Agent :** Vastech Limited, Nurjehan Tower (6th Floor), 80/22 Mymensingh Road Dhaka	Renegade HI-FLO	Microcatheter	C	The Renegade HI-FLO Microcatheter is intended for peripheral vascular use. The microcatheter can be coaxially tracked over a steerable guidewire in order to access distal, tortuous vasculature. Once the subselective region has been accessed, the microcatheter can be used for the controlled and selective infusion of diagnostic, embolic, or therapeutic materials into vessels. Diagnostic, embolic, or therapeutic agents to be used in accordance with specifications outlined by the manufacturer.	**Contraindication:** Not Known. **Side Effects:** The Adverse Events include, but are not limited to: Vessel trauma, Embolism, Hemorrhage/Hematoma, Vasospasm, Infection, Air embolism, Allergic reaction	Ireland EC Certificate	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj
39.	**Manufacturer :** Boston Scientific Limited, Ballybrit Business Park, Galaway, Ireland **Legal Manufacturer:** Boston Scientific Corporation, USA **Local Agent :** Vastech Limited, Nurjehan Tower (6th Floor), 80/22 Mymensingh Road Dhaka	Renegade HI-FLO FATHOM System	Microcatheter	C	The Renegade HI-FLO FATHOM System is intended for peripheral vascular use. The FATHOM Guidewire can be used to selectively introduce and position the Renegade HI-FLO Microcatheter in the peripheral vasculature. The microcatheter can be used for controlled and selective infusion of diagnostic, embolic, or therapeutic materials into vessels.	**Contraindication:** Not Known. **Adverse events:** The Adverse Events include, but are not limited to: Allergic reaction, Death, Embolism, Hemorrhage/Hematoma, Infection, Pseudoaneurysm, Stroke, Vascular thrombosis, Vessel occlusion, Vessel spasm, Vessel trauma (dissection, perforation, rupture)	Ireland EC Certificate	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj

Sl No	c¯ZKvi‡Ki bvg	ewYR¨K bvg	‡gvW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve Kwgkb Ui mfvi wm×vš	mfvi wm×vš
40.	**Manufacturer :** Boston Scientific Corporation, USA **Local Agent :** Vastech Limited, Nurjehan Tower (6th Floor), 80/22 Mymensingh Road Dhaka	**Rubicon**	Support Catheter	C	The Rubicon Support Catheter is intended to facilitate placement and support of guidewires and other interventional devices within the peripheral vasculature and to allow for exchange of guidewires, and provide a conduit for the delivery of saline or contrast solutions.	**Contraindications:** None known **ADVERSE EVENTS:** Vascular catheterization and/or vascular interventions may result in complications including but not limited to: Access site pain, Allergic reaction (drug, contrast, device & other) and neurological reactions, Death, Hemorrhage or hematoma, Impaired blood flow due to thrombosis, embolism, or vasospasm that could lead to tissue infarction, limb amputation, and other thrombo-embolic organ damage such as renal infarction, Infection/Sepsis, Vessel injury (dissection, perforation, trauma & rupture), Vasospasm	USA EC Certificate	Abyôgv`b Kiv †h‡Z cv‡i	Abyôgv`b Kiv nj
41.	**Manufacturer:** GE Medical Systems (China) Co., Ltd. No. 19, Changjiang Road, Wuxi National Hi-Tech Dev. Zone, 214028 Jiangsu, China **Agent**: GE Healthcare Bangladesh Ltd. Shanta Western Tower, OS # 804, Level # 08, 186 Tejgaon I/ A Dhaka	**9100c** (**Three Gas system:** Oxygen, Air, Nitrous Oxide)	Anesthesia System	**C** (P-29)	It is intended to provide general inhalation anesthesia and ventilatory support to a wide range of patients. The devices are intended for volume or pressure control ventilation.	**Contraindication:** There aren't any specific contraindications for these anesthesia machines, however, these systems are designed for hospital use and should only be used under the orders of a clinician. **Side effect:** None	China EC Certificate	Abyôgv`b Kiv †h‡Z cv‡i	Abyôgv`b Kiv nj
42.	**Manufacturer:** Datex-Ohmeda, Inc. 3030 Ohmeda Drive PO Box 7550 Madison, WI 53707-7550, USA **Local Agent**: GE Healthcare Bangladesh Ltd. Shanta Western Tower, OS # 804, Level # 08, 186 Tejgaon I/ A Dhaka- 1208	**Aespire View** (Anesthesia Machine)	GAS-Machine, Anesthesia	C (P-49)	The Aespire View anesthesia system is intended to provide general inhalation anesthesia and ventilatory support to a wide range of patients. The device is intended for volume or pressure control ventilation.	**Contraindication:** There are no contraindications for Aespire. **Side effect:** None	USA EC Certificate	Abyôgv`b Kiv †h‡Z cv‡i	Abyôgv`b Kiv nj

Sl No	c¯ZKvi‡Ki bvg	ewYRK bvg	‡gvW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve KwgwUi mfvi wm×vš	mfvi wm×vš
43.	**Manufacturer:** Smith & Nephew Medical Limited, UK **Local Agent:** UniMed Ltd., House No. 46, Sheikh Kamal Saroni, Road No. 16, Rangs Nasim (7th Floor) Square, Dhanmondi	Allevyn Adhesive	Adhesive Hydrocellular Foam Dressing	C	ALLEYVN Adhesive dressings are indicated for use on: Shallow, granulating wounds, Chronic and acute exudative wounds, Full and partial thickness wounds such as pressure ulcers, leg ulcers and diabetic foot ulcers, Infected wounds, Malignant wounds, Surgical wounds, First and second degree burns, Donor sites Fungating wounds	**Contraindications:** Not known **Side Effects:** Not known	MHRA-UK EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj	
44.	Manufacturer: Smith & Nephew Medical Limited, UK Local Agent: UniMed Ltd., House No. 46, Sheikh Kamal Saroni, Road No. 16, Rangs Nasim (7th Floor) Square, Dhanmondi	Allevyn Gentle Border Sacrum	Silicone gel adhesive hydrocellular foam dressing	C	ALLEVYN Gentle Border Lite is indicated for wound management by secondary intention on shallow, granulating wounds, chronic and acute low exuding wounds full and partial thickness wounds including: Skin tears Pressure ulcers Diabetic foot ulcers Leg ulcers Fungating wounds Infected wounds Malignant wounds Surgical wounds Donor sites First and second degree burns	Contraindications: Not known Side Effects: Not known **Precautions:** Do not use ALLEVYN Gentle Border Sacrum dressing with oxidizing agents such as hypochlorite solutions (e.g. EUSOL) or hydrogen peroxide, as these can break down the absorbent polyurethane component of the dressing. If reddening or sensitization occurs discontinue use.	MHRA-UK	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj	
45.	Manufacturer: Smith & Nephew Medical Limited, UK Local Agent: UniMed Ltd., House No. 46, Sheikh Kamal Saroni, Road No. 16, Rangs Nasim (7th Floor) Square, Dhanmondi	Allevyn Heel	Hydrocellular Heel Dressing	C	ALLEVYN Heel is indicated for exudate absorption and the management of partial to full thickness wounds. Some typical wounds are: Ulcers (venous, arterial, diabetic), Pressure Sores, Surgical Incisions, Surgical Excisions, Burns (1st and 2nd degree),. Creation and maintenance of a moist wound environment. Moist wound environments have been established as optimal environments for the management of the wound. Provides physical separation between the wound and external environments to assist in preventing bacterial contamination of the wound.	**Contraindications:** Not known **Side Effects:** Not known **Precautions:** Do not use ALLEVYN dressings with oxidising agents such as hypochlorite solutions (e.g. Dakins) or hydrogen peroxide solutions, as these can break down the absorbent hydrocellular component of the dressing. Reddening of the skin around the wound has been reported rarely following the use of ALLEVYN dressings. In some cases this relates to irritation of fragile skin, in others, wound exudate remaining in contact with peri-wound skin for prolonged periods may be the cause. Infrequently, cases of sensitivity to the dressing have also been reported. If reddening or sensitisation occur, discontinue use and consult a healthcare professional. ALLEVYN Heel is not indicated for use on third degree burns.	MHRA-UK	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj	

Sl No	c¯ZKvi‡Ki bvg	ewYRK bvg	‡gvW‡Kj vWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve KvgvUi mfvi wm×vš	mfvi wm×vš
46.	Manufacturer: Smith & Nephew Medical Limited, UK Local Agent: UniMed Ltd., House No. 46, Sheikh Kamal Saroni, Road No. 16, Rangs Nasim (7th Floor) Square, Dhanmondi	Allevyn Non Adhesive	Non-adhesive hydrocellular polyurethane dressing	C	ALLEVYN Non-Adhesive dressings are indicated for use on: • Shallow, granulating wounds • Chronic and acute exudative wounds • Full and partial thickness wounds such as pressure ulcers, leg ulcers and diabetic foot ulcers • Infected wounds • Malignant wounds • Surgical wounds • First and second degree burns • Donor sites • Fungating ulcers • Oncological Wounds • Epidermolysis bullosa wounds	**Contraindications:** Not known **Side Effects:** Not known	MHRA-UK	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj	
47.	**Manufacturer:** Smith & Nephew Medical Limited, UK **Local Agent:** UniMed Ltd., House No. 46, Sheikh Kamal Saroni, Road No. 16, Rangs Nasim (7th Floor) Square, Dhanmondi	Allevyn Sacrum	Hydrocellular Adhesive Sacrum Dressing	C	Exuding granulating wounds (for example: pressure sores) It can be used in conjunction with INTRASITE◊ Gel - Hydrogel Wound Dressing for necrotic or sloughy wounds	**Contraindications:** Not known **Side Effects:** Not known	MHRA-UK	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj	
48.	**Manufacturer:** Boston Scientific Corporation, 300 Boston Scientific Way, Marlborough, MA 01752, USA **Local Agent:** M/s. Medi Graphic Trading Ltd., 14 Purana Paltan, Dhaka, Bangladesh	Amplatz Super Stiff Guide wire	Peripheral Guidewires	C	The Amplatz Super Stiff guidewire facilitates catheter placement and exchange during diagnostic or interventional procedures.	**Contraindications:** None known. **Adverse Events:** Allergic Reaction, Amputation, Arteriovenous (AV) Fistula, Death, Embolism, Hematoma, Hemorrhage, Hemoglobinuria, Infection or Sepsis/Infection, Myocardial Ischemia and/or Infarction, Pseudoaneurysm, Stroke (CVA)/Transient Ischemic Attacks (TIA), Thrombus, Vessel Occlusion, Vessel Perforation, Dissection, Trauma, or Damage, Vessel Spasm,Wire Entrapment/Entanglement, Foreign body/Wire Fracture	USA EC Certificate	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj	

Sl No	c¯ZKvi‡Ki bvg	ewYR¨K bvg	‡gvW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve Kwgkb Ui mfvi wm×vš	mfvi wm×vš
49.	**Manufacturer:** GE Hualun Medical Systems, Co. Ltd, China **Local Agent**: GE Healthcare Bangladesh Ltd. Shanta Western Tower, OS # 804, Level # 08, 186 Tejgaon I/ A Dhaka	1. Brivo OEC 715 2. Brivo OEC 785 3. Brivo OEC 865	Mobile C-arm X-ray Product	C	The OEC® Brivo Mobile C-Arm X-Ray Products are designed to provide digital spot-film imaging and fluoroscopic image guidance for all adult and pediatric populations for orientations between patient anatomy and surgical instruments. The product is used for general surgical applications and musculoskeletal procedures to visualize, for example, implant localization/s or needle positions for aspirations, injections or biopsy. The OEC® Brivo® is not indicated for interventional use.	**Contraindication:** Pregnant or lactating women, persons who can't tolerate surgery, persons who have mental disorders and can't cooperate in surgery. **Side effect:** None	China USA EC Certificate	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj|
50.	**Manufacturer:** GE Healthcare Finland Oy, Kuortaneenkatu 2 FI-00510 Helsinki Finland **Local agent:** GE Healthcare Bangladesh Ltd., Shanta Western Tower., 186, Tejgaon. Dhaka	**CARESCAPE Monitor** B450/ B650/B850	Patient Monitoring system	C (P-115)	The CARESCAPE Monitor is a multi-parameter patient monitor intended for use in multiple areas and intra-hospital transport within a professional healthcare facility. The CARESCAPE Monitor B450, B650, B850 is intended for use on adult, pediatric, and neonatal patients and on one patient at a time. The CARESCAPE Monitor B450, B650, B850 is indicated for monitoring of: • hemodynamic (including ECG, ST segment, arrhythmia detection, ECG diagnostic analysis and measurement, invasive pressure, non-invasive blood pressure, pulseoximetry, cardiac output (thermodilution and pulse contour), temperature, mixed venous oxygen saturation, central venous oxygen saturation, and Surgical Pleth Index), • respiratory (impedance respiration, airway gases (CO2, O2, N2O, and anesthetic agents), spirometry, gas exchange), and • Neurophysiological status (including electroencephalography, Entropy, Bispectral Index (BIS), andneuromuscular transmission). The CARESCAPE Monitor B450, B650, B850 also provides alarms, trends, snapshots and events, and calculations, and can be connected to displays, printers and recording devices. The CARESCAPE Monitor B450, B650, B850 can be a stand-alone monitor or interfaced to other devices. It can also be connected to other monitors for remote viewing and to data management software devices via a network. The CARESCAPE Monitor B450, B650, B850 is intended for use under the direct supervision of a licensed healthcare practitioner, or by personnel trained in proper use of the equipment in a professional healthcare facility. The CARESCAPE Monitor B450, B650, B850 is not intended for use during MRI.	**Contraindication:** The system is not intended for use in a MRI environment. **Side Effects:** None	Finland EC Certificate	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj|

Sl No	c¯ZKvi‡Ki bvg	ewYwR¨K bvg	‡gvW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve KwgwUi mfvi wm×vš	mfvi wm×vš
51.	**Manufacturer:** Datex-Ohmeda, Inc. 3030 Ohmeda Drive PO Box 7550 Madison, WI 53707-7550, USA **Local Agent**: GE Healthcare Bangladesh Ltd. Shanta Western Tower, OS # 804, Level # 08, 186 Tejgaon I/ A Dhaka	CARESCAPE R860 Ventilator	Ventilator	C	It is designed to provide mechanical ventilation or support to neonatal, pediatric, and adult patients weighing 0.25 kg and above. The CARESCAPE R860 ventilator is a microprocessor based, electronically controlled, pneumatically driven ventilator that includes integrated monitoring of FiO2, airway pressure, flow, and volume. Additional respiratory gas monitoring capabilities are supported through the use of optional GE patient monitoring modules. Not all features are available for all patient types or product configurations. The CARESCAPE R860 ventilator is not a pulmonary function calculation device. The system is designed for facility use, including within-facility transport, and should only be used under the orders of a clinician.	**Contraindication:** There are no contraindications for CARESCAPE R860 system. **Side effect:** None	USA EC Certificate	Abyôgv`b Kiv †h‡Z cv‡i	Abyôgv`b Kiv nj
52.	**Manufacturer:** GE Medical Systems (China) Co., Ltd. No. 19, Changjiang Road, Wuxi National Hi-Tech Dev. Zone, 214028 Jiangsu, China **Agent**: GE Healthcare Bangladesh Ltd. Shanta Western Tower, OS # 804, Level # 08, 186 Tejgaon I/ A Dhaka	Carestation 30 (**Two Gas system:** Oxygen & Air/Nitrous Oxide)	Anesthesia Machine	**C** (P-29)	It is intended to provide general inhalation anesthesia and ventilatory support to a wide range of patients. The devices are intended for volume or pressure control ventilation.	**Contraindication:** There aren't any specific contraindications for these anesthesia machines, however, these systems are designed for hospital use and should only be used under the orders of a clinician. **Side effect:** None	China EC Certificate	Abyôgv`b Kiv †h‡Z cv‡i	Abyôgv`b Kiv nj
53.	**Manufacturer:** Alcon Laboratories Inc, USA **Local Agent:** Globex Marketing Company Ltd. RH Home Center (6th Floor) 74/B/1, Green Road, Dhaka-1215	Centurion Vision System	Centurion Vision System	**C**	It is an ophthalmic microsurgical system that is indicated for both anterior segment (i.e. phacoemulsification, removal of cataracts, and intraocular lens injection) and posterior segment (i.e., vitreoretinal) ophthalmic surgery in addition to the indications included with the optional Next Generation laser. The AutoSert® IOL Injector Handpiece is intended to deliver qualified ACRYSOF® intraocular lenses into the eye following cataract removal. The AutoSert® IOL Injector Handpiece achieves the functionality of injection of intraocular lenses. The AutoSert® is indicated for use with ACRYSOF lenses SN60WF, SN6AD1, SN6AT3 through SN6AT9, as well as approved AcrySof® lenses that are specifically indicated for use with this inserter, as indicated in the approved labeling of those lenses.	**Contraindication:** None **Side Effects:** None	USA EC Certificate	Abyôgv`b Kiv †h‡Z cv‡i	Abyôgv`b Kiv nj

Sl No	c¯ZKvi‡Ki bvg	ewYR¨K bvg	‡gvW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve Kwgkb Ui mfvi wm×vš	mfvi wm×vš
54.	**Manufacturer:** Alcon Laboratories Inc, USA **Local Agent:** Globex Marketing Company Ltd. RH Home Center (6th Floor) 74/B/1, Green Road, Dhaka-1215	Constellation Vision System, Accessories & Consumables	Vision System Accessories & Consumables	**C**	The Constellation® Vision System is an ophthalmic microsurgical system that is indicated for both anterior segment (i.e. phacoemulsification and removal of cataracts) and posterior segment (i.e., vitreoretinal) ophthalmic surgery in addition to the indications included with the optional Next Generation laser.	Contraindication: Not Known Side Effects: Not Known	USA EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj	
55.	**Manufacturer:** GE MEDICAL SYSTEMS SCS, 283 RUE DE LA MINIERE 78530 BUC- FRANCE **Local Agent**: GE Healthcare Bangladesh Ltd. Shanta Western Tower, OS # 804, Level # 08, 186 Tejgaon I/ A Dhaka	Discovery IGS 730/740	Angiographic X-ray System	**C** (P-93)	The angiographic X-ray systems are indicated for use in generating fluoroscopic and rotational images of human anatomy for cardiovascular, vascular and non-vascular, diagnostic andinterventional procedures. Additionally, with the OR table, the angiographic X-ray systems ore indicated for use in generatingfluoroscopic and rotational images of human anatomy for image-guided surgical procedures. The OR table is suitable for interventional and surgical procedures.	**Contraindication:** These devices are not intended for mammography applications. **Side effect:** None	France EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj	
56.	**Manufacturer:** Ohmeda Medical A Division of Datex - Ohmeda Inc. 8880 Gorman Road Laurel Maryland 20723 USA **Local Agent**: GE Healthcare Bangladesh Ltd. Shanta Western Tower, OS # 804, Level # 08, 186 Tejgaon I/ A Dhaka	Giraffe Incubator Carestation CS1	Baby Incubator	C	It is an Infant Incubator. Incubators provide heat in a controlled manner to neonates who are unable to thermo-regulate based on their own physiology. They achieve this by providing an enclosed temperature controlled environment to the infant. This device may incorporate a Servo Controlled Oxygen Delivery System. This is indicated to provide a stable oxygen concentration within the infant compartment at the value set by the operator (21-65%).	**Contraindication:** None. **Side effect:** None	USA EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj	

Sl No	c¯ZKvi‡Ki bvg	ewYRwK bvg	‡gvW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve Kwgkb mfvi wm×vš	mfvi wm×vš
57.	**Manufacturer:** OHMEDA Medical 8880 Gorman Road Laurel Maryland, MD USA 20723 **Local Agent**: GE Healthcare Bangladesh Ltd. Shanta Western Tower, OS # 804, Level # 08, 186 Tejgaon I/ A Dhaka	Giraffe Warmer	Infant Radiant Warmer	C (P-99)	Infant radiant warmer provide infrared heat in a controlled manner to neonates who are unable to thermo-regulate based in their own physiology. Infant radiant warmersmay be used to facilitate the neonate's transition to the external environment or to provide acontrolled open environment. An optional integrated SpO2 monitoring feature may be used for continuous noninvasive monitoring of functional oxygen saturation of arterial hemoglobin. (SpO2) and pulse rate (measured by an Sp02 sensor). An optional integrated resuscitation system may be used to provide the basic equipmentrequiredforpulmonaryresuscitation of infants. Pulmonary resuscitation includes practices necessary to establish a clear airway and provide oxygen or air/oxygen mixture and/or manual ventilation to the infant.	**Contraindication:** There are no known contraindications to the use of either radiant warmers or resuscitation components in the intended population discussed in the Warmers technical files and based upon equivalent devices in the marketplace. **Side effect:** None	USA EC Certificate	Abyôgv`b Kiv †h‡Z cv‡i	Abyôgv`b Kiv nj
58.	**Manufacturer:** Alcon Laboratories Inc, USA **Local Agent:** Globex Marketing Company Ltd. RH Home Center (6th Floor) 74/B/1, Green Road, Dhaka-1215	INFINITI Vision System Accessories and Spare Parts	Vision System, Accessories and Spare Parts	**C**	The Infinitre Vision System is indicated for emulsification, separation, and removal of cataracts, the removal of residual cortical material and lens epithelial cells, vitreous aspiration and cutting associated with anterior vitrectomy, bipolar coagulation, and intraocular lens injection. The AutoSert TM IOL Injector Handpiece is intended to deliver qualified AcrySofo intraocular lenses into the eye following cataract removal. The following system modalities additionally support the described indications: - Ultrasound with UltraChopper Tip achieves the functionality of cataract separation. - AquaL-ase achieves the functionality for removal of residual cortical material and lens epithelial cells - Th Aut~ertM CLInjetor andiece achieves the functionality of injection of intraocular lenses. The AutoSertT is indicated for use with ACRYSOF lenses SN60WF and SN6AD1, as well as approved Acrysof lenses that are specifically indicated for use with this inserter, as indicated in the approved labeling of those lenses.	**Contraindication:** Not known. **Side Effects:** Not known.	USA EC Certificate	Abyôgv`b Kiv †h‡Z cv‡i	Abyôgv`b Kiv nj

Sl No	c¯ZKvi‡Ki bvg	ewYRK bvg	‡gvW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve KwgwUi mfvi wm×vš	mfvi wm×vš
59.	**Manufacturer:** GE MEDICAL SYSTEMS SCS 283 RUE DE LA MINIERE, 78530 BUC-FRANCE **Local Agent**: GE Healthcare Bangladesh Ltd. Shanta Western Tower, OS # 804, Level # 08, 186 Tejgaon I/ A Dhaka	Innova IGS 520/530/540/620/630	Angiographic X-ray System	**C** (P-93)	The angiographic X-ray systems are indicated for use for patients from newborn to geriatric in generating fluoroscopic and rotational images of human anatomy for cardiovascular, vascular and non-vascular, diagnostic and interventional procedures. Additionally, with the OR table, the angiographic X-ray systems are indicated for use in generating fluoroscopic and rotational images of human anatomy for image-guided surgical procedures. The OR table is suitable for interventional and surgical procedures.	**Contraindication:** These devices are not intended for mammography applications. **Side effect:** None	France USA EC Certificate	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj
60.	**Manufacturer:** Alcon Laboratories Inc, USA **Local Agent:** Globex Marketing Company Ltd. RH Home Center (6th Floor) 74/B/1, Green Road, Dhaka-1215	Laureate World Phaco System, and Accessories	Phaco System and Accessories	**C**	The LAUREATE® World Phaco System is a traditional phacoemulsification system intended for use in anterior segment procedures that require simultaneous lens extraction, irrigation, and aspiration, as well as associated procedures such as anterior vitrectomy and coagulation.	Contraindication: Not Known Side Effects: Not Known	USA	c`wU ¯wMZ Kiv nj Ges Gi Safety data `wLj Kivi Rb¨ ejv †h‡Z cv‡i	c`wU ¯wMZ Kiv nj Ges Gi Safety data `wLj Kivi Rb¨ ejv nj
61.	**Manufacturer:** Alcon Laboratories Inc, USA **Local Agent:** Globex Marketing Company Ltd. RH Home Center (6th Floor) 74/B/1, Green Road, Dhaka-1215	LenSx Laser System	Laser System	**C**	The LenSx Laser System is indicated for use: • In the creation of corneal cuts/incisions (single-plane, multi-plane and arcuate), anterior capsulotomy and laser phacofragmentation during cataract surgery. Each of these procedures may be performed either individually or consecutively during the same surgery. • In the creation of a corneal flap in patients undergoing LASIK surgery or other treatment requiring initial lamellar resection of the cornea	Contraindication: Not Known Side Effects: Not Known	USA EC Certificate	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj

Sl No	c¯ZKvi‡Ki bvg	ewYRK bvg	‡gvW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve Kwgkb Ui mfvi wm×vš	mfvi wm×vš
62.	**Manufacturer:** GE Healthcare GE Medical Systems (China) Co., Ltd. China **Local Agent**: GE Healthcare Bangladesh Ltd. Shanta Western Tower, OS # 804, Level # 08, 186 Tejgaon I/ A Dhaka	LOGIQ e Ultrasound System	Ultrasound Diagnostic system	C	The LOGIQ e is intended for ultrasound imaging, measurement and analysis of the human body for multiple clinical applications including: ophthalmic: fetal/ob; abdominal (gyn& urology); pediatric; small organ (breast, testes, thyroid); neonatal and adult cephalic; cardiac (adult & pediatric); peripheral vascular; musculoskeletal conventional & superficial;transrectal; transvaginal; transesophageal; intraoperative (abdominal, thoracic and peripheral); thoracic/pleural for motionand fluid detection and imaging guidance of interventional procedures (e.g. Nerve block; vascular access).	**Contraindication:** The System is not intended for ophthalmic use or a use causing the acoustic beam to pass through the eye. Note the LOGIQ E (as of R6) with 8C-RS, 12L-RS, L4-12t-RS, and 3SC-RS; and Venue 50 with the 10C-SC,12L-SC and 3S-SC probes have been tested and meet the requirements for Ophthalmic use and are not included in this contraindication. **Side effect:** None	China USA EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
63.	**Manufacturer:** GE Medical Systems (China) Co., Ltd. China **Local Agent**: GE Healthcare Bangladesh Ltd. Shanta Western Tower, OS # 804, Level # 08, 186 Tejgaon I/ A Dhaka	LOGIQ F6, LOGIQ F8,	Ultrasound Diagnostic System	C	The LOGIQ F8 / LOGIQ F6 are general purpose ultrasound imaging and analysis systems providing digital acquisition, processing and display capability, clinical applications including: Abdominal, Obstetrical, Gynecological, Small parts, Vascular/Peripheral Vascular, Transcranial, Pediatric, Musculoskeletal, Urological, Cardiac, Transvaginal, Transrectal, and Biopsy.	**Contraindication:** The System is not intended for ophthalmic use or a use causing the acoustic beam to pass through the eye. **Side effect:** None	China USA EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
64.	**Manufacturer:** GE Ultrasound Korea, Ltd.Republic of Korea **Local Agent**: GE Healthcare Bangladesh Ltd. Shanta Western Tower, OS # 804, Level # 08, 186 Tejgaon I/ A Dhaka	**LOGIQ P7; LOGIQ P9**	Ultrasound Imaging System, General Purpose	C	The device is intended for use by a qualified physician for ultrasound evaluation of Fetal; Abdominal; Pediatric; Small Organ (breast, testes, thyroid); Neonatal Cephalic; Adult Cephalic; Cardiac (adult and pediatric); Peripheral Vascular; Musculo-skeletal Conventional and Superficial; Urology (including prostate);Transrectal; Transvaginal; and Intraoperative (abdominal, thoracic, vascular).	**Contraindication:** The System is not intended for ophthalmic use or a use causing the acoustic beam to pass through the eye. **Side effect:** None	Korea USA EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

Sl No	c¯ZKvi‡Ki bvg	ewYRK bvg	‡gvW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve Kwgkb mfvi wm×vš	mfvi wm×vš
65.	**Manufacturer:** GE Medical Systems (China) Co., Ltd, China **Local Agent**: GE Healthcare Bangladesh Ltd. Shanta Western Tower, OS # 804, Level # 08, 186 Tejgaon I/ A Dhaka	LOGIQ V1(Black & White)/ LOGIQ V2 (Color)	Ultrasound Diagnostic system	C	These are general purposed ultrasound imaging and analysis systems providing digital acquisition, processing and display capability and clinical applications including: Abdominal, Obstetrical, Gynecological, Small parts, Vascular/Peripheral Vascular, Adult Cephalic, Pediatric, Musculoskeletal, Transcranial, Neonatal Cephalic, Transvaginal, Urological, Cardiac, Transrectal, and Tissue Biopsy/Fluid Drainage.	**Contraindication:** The System is not intended for ophthalmic use or a use causing the acoustic beam to pass through the eye. **Side effect:** None	China USA EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
66.	**Manufacturer:** GE Medical Systems (China) Co., Ltd, China **Local Agent**: GE Healthcare Bangladesh Ltd. Shanta Western Tower, OS # 804, Level # 08, 186 Tejgaon I/ A Dhaka	LOGIQ V3; LOGIQ V5	Ultrasound Diagnostic system	C	These are general purposed ultrasound imaging and analysis systems providing digital acquisition, processing and display capability and clinical applications including: Abdominal, Obstetrical, Gynecological, Small parts, Vascular/Peripheral Vascular, Adult Cephalic, Pediatric, Musculoskeletal, Transcranial, Neonatal Cephalic, Transvaginal, Urological, Cardiac, Transrectal, and Tissue Biopsy/Fluid Drainage.	**Contraindication:** The System is not intended for ophthalmic use or a use causing the acoustic beam to pass through the eye. **Side effect:** None	China USA EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
67.	**Manufacturer:** Wipro GE healthcare Private Ltd., India **Local Agent**: GE Healthcare Bangladesh Ltd. Shanta Western Tower, OS # 804, Level # 08, 186 Tejgaon I/ A Dhaka	Lullaby Warmer	Infant Radiant Warmer	C	Infant radiant Warmers provide infrared heat in a controlled manner to infants who are unable to maintain thermoregulation based on their own physiology. Infant radiant Warmers may be used to facilitate the newborn's transition to the external environment or to provide a controlled open microenvironment.	**Contraindication:** None **Side effect:** None	India USA EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

Sl No	c¯ZKvi‡Ki bvg	ewYR¨K bvg	‡gvW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve Kwgkb mfvi wm×vš	mfvi wm×vš
68.	**Manufacturer:** GE HUALUN MEDICAL SYSTEMS Co., LTD.,China **Local Agent**: GE Healthcare Bangladesh Ltd. Shanta Western Tower, OS # 804, Level # 08, 186 Tejgaon I/ A Dhaka-1208, Bangladesh. Tel No. : 880 961 0887800. email ID: abidur.chowdhury@ge.com	Optima IGS 320/ Optima IGS 330	**Angiographic X-Ray Systems**	**C** (P-93)	The angiographic X-ray systems are indicated for use for patients from newborn to geriatric in generating fluoroscopic and rotational images of human anatomy for cardiovascular, vascular and non-vascular, diagnostic and interventional procedures.	**Contraindication:** These devices are not intended for mammography applications. **Side effect:** None	China USA EC Certificate	Abyôgv`b Kiv †h‡Z cv‡i	Abyôgv`b Kiv nj
69.	**Manufacturer:** Alcon Laboratories Inc, USA **Local Agent:** Globex Marketing Company Ltd. RH Home Center (6th Floor) 74/B/1, Green Road, Dhaka-1215	ORA System 2000, Accessories, Objectives Lens and Dovetail Kits	ORA System with VeriEye Plus	**C**	The ORA System uses wavefront aberrometry data in the measurement and analysis of the refractive power of the eye (i.e. sphere, cylinder, and axis measurements) to support cataract surgical procedures.	CONTRAINDICATIONS: The ORA System is contraindicated for patients: • who have progressive retinal pathology such as diabetic retinopathy, macular degeneration, or any other pathology that the physician deems would interfere with patient fixation; • who have corneal pathology such as Fuchs', EBMD, keratoconus, advanced pterygium impairing the cornea, or any other pathology that the physician deems would interfere with the measurement process; • whose preoperative regimen includes residual viscous substances left on the corneal surface such as lidocaine gel or viscoelastics; • with visually significant media opacity (such as prominent floaters or asteroid hyalosis) what will either limit or prohibit the measurement process; or • who have received retro or peribulbar block	USA EC Certificate	Abyôgv`b Kiv †h‡Z cv‡i	Abyôgv`b Kiv nj

						or any other treatment that impairs their ability to visualize the fixation light. In addition, utilization of iris hooks during an ORA System image capture is contraindicated, because the use of iris hooks will yield inaccurate measurements. WARNINGS AND PRECAUTIONS: • Significant central corneal irregularities resulting in higher order aberrations might yield inaccurate refractive measurements. • Post refractive keratectomy eyes might yield inaccurate refractive measurement. • The safety and effectiveness of using the data from the ORA System have not been established for determining treatments involving higher order aberrations of the eye such as coma and spherical aberrations. • The ORA System is intended for use by qualified health personnel only. • Improper use of this device may result in exposure to dangerous voltage or hazardous laser-like radiation exposure. • Do not operate the ORA System in the presence of flammable anesthetics or volatile solvents such as alcohol or benzene, or in locations that present an explosion hazard.			

Sl No	c¯ZKvi‡Ki bvg	ewYRK bvg	‡gvW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve Kwgkb mfvi wm×vš	mfvi wm×vš
70.	**Manufacturer:** GE Medical Systems (China) Co. Ltd., China **Local agent:** GE Healthcare Bangladesh Ltd., Shanta Western Tower, 186, Tejgaon. Dhaka	Patient Monitor B20/B40	Patient Monitoring system	C (P-115)	It is a portable multi-parameter unit to be used for monitoring and recording of, and to generate alarms for, multiple physiological parameters of adult, pediatric, and neonatal patients in a hospital environment and during intra-hospital transport. The device is intended for use under the direct supervision of a licensed health care practitioner. The device is not intended for use during MRI. The device can be a stand-alone monitor or interfaced to other devices via a network. The device monitors and displays : ECG (including ST segment, arrhythmia detection), invasive blood pressure, heart/pulse rate, oscillometric non-invasive blood pressure (systolic, diastolic and mean arterial pressure), functional oxygen saturation (SpO2) and pulse rate via continuous monitoring (including monitoring during conditions of clinical patient motion or low perfusion), temperature with a reUSAble or disposable electronic thermometer for continual monitoring Esophageal/Nasopharyngeal/Tympanic/Rectal/Bladder/ Axillary/Skin/Airway/Room/Myocardial/Core/Surface temperature, impedance respiration, respiration rate, airway gases (CO2, O2, N2O, anesthetic agents, anesthetic agent identification and respiratory rate), and Entropy.	**Contraindication:** The system is not intended for use in a MRI environment. **Side Effects:** None	China USA	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
71.	**Manufacturer:** ST. Stone Medical Devices Pvt. Ltd, India **Local Agent:** New Vision Medisystem 17/2, 3rd Floor, New Eskaton Road, Mogbazar, Dhaka	PLATICATH	Dialysis Catheter	C	It is used for exchanging blood to and from a hemodialysis machine and a patient.	**Contraindications:** Not Known **Side Effects:** Not Known	India EC Certifiacte	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

Sl No	c¯ZKvi‡Ki bvg	ewYR¨K bvg	‡gvW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve Kwgkb Ui mfvi wm×vš	mfvi wm×vš
72.	**Manufacturer:** Ethicon, LLC, USA **Distributor:** Johnson & Johnson PVt. Ltd., India **Local Agent:** Modern Surgical (pvt) Ltd., 34/1, (2nd Floor) Mitford Road, Dhaka	PROLENE Propylene Soft Mesh	Soft Mesh	**C**	This mesh may be used for the repair of hernia or other fascial defects that require the addition of a reinforcing or bridging material to obtain the desired surgical result.	**Contraindications:** When this mesh is used in infants or children with future growth potential, the surgeon should be aware that this product will not stretch significantly as the patient grows, PROLENE Soft mesh in contaminated wounds should be used with the understanding that subsequent infection may require removal of the material **Adverse Reactions:** Potential adverse reactions are those typically associated with surgically implantable materials, including infection potentiation, inflammation, adhesion formation, fistula formation, and extrusion.	USA EC Certificate	Abyjgvb Kiv †h‡Z cv‡i	Abyjgvb Kiv nj
73.	**Manufacturer:** Alcon Laboratories Inc, USA **Local Agent:** Globex Marketing Company Ltd., RH Home Center (6th Floor) 74/B/1, Green Road, Dhaka-1215	PurePoint Laser System	Laser System	**C**	The PUREPOINT® Laser is indicated for use in photocoagulation of both anterior and posterior segments of the eye including: • Retinal photocoagulation, panretinal photocoagulation and intravitreal endophotocoagulation of vascular and structural abnormalities of the retina and choroid including: Proliferative and nonproliferative retinopathy (including diabetic); choroidal neovascularization secondary to age-related macular degeneration; retinal tears and detachments; macular edema, retinopathy of prematurity; choroidal neovascularization; leaking microaneurysms. • Iridotomy/Iridectomy for treatment of chronic/primary open angle glaucoma, acute angle closure glaucoma and refractory glaucoma. • Trabeculoplasty for treatment of chronic/primary open angle glaucoma and refractory glaucoma. • And other laser treatments including: internal sclerostomy; lattice degeneration; central and branch retinal vein occlusion; suturelysis; vascular and pigment skin lesions.	Contraindications: Patients with a condition that prevents visualization of target tissue (cloudy cornea, or extreme haze of the aqueous humor of the anterior chamber of vitreous humor) are poor candidates for LIO delivered laser treatments. **Complications:** Corneal burns, inflammation, loss of best-corrected visual acuity, loss of visual field and transient elevations in intraocular pressure can occur as a result of ophthalmic laser treatment. Unintentional retinal burns can occur if excessive treatment beam power or duration is used.	USA EC Certificate	Abyjgvb Kiv †h‡Z cv‡i	Abyjgvb Kiv nj

Sl No	c¯ZKvi‡Ki bvg	emvYvi¨K bvg	‡gvW‡Kj vWfvB‡mi bvg	Kvm	vb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKvbK¨vj mve Kvgvi mfvi vm×vš	mfvi vm×vš
74.	**Manufacturer:** GE Healthcare Japan Corporation, Japan, GE Medical System, LLC, USA	Revolution EVO Optima CT660	Computed Tomography System	C	These systems are intended to produce cross-sectional images of the body by computer reconstructionof x-ray transmission data taken at different angles and planes, including Axial, Cine, Helical (Volumetric), Cardiac, and Gated acquisitions. These images may be obtained either with or withoutcontrast. This device may include signal analysis and display equipment, patient and equipmentsupports, components and accessories. This device may include data and image processing to produce images in a variety of trans-axialand reformatted planes. Further the images can be post processed to produce additional imagingplanes or analysis results. The system is indicated for head, whole body, cardiac and vascular X-ray Computed Tomographyapplications in patients of all ages. The device output is a valuable medical tool for the diagnosis of disease,trauma, or abnormalityand for planning, guiding, and monitoring therapy.	**Contraindication:** There are no absolute contraindications for use of it. As with all procedures, the relative benefits and risks of the procedure should be evaluated prior to the performance of CT, with or without the administration of intravenous iodinated contrast or oral contrast media. Appropriate precautions should be taken to minimize patient risks, including radiation exposure. **Side effect:** None	Japan USA EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
75.	**Manufacturer:** Echo-Son S.A, Poland **Local Agent:** Access Medical Services, R.H Home Center, 74/B/1, Green Road, Dhaka-1215	Ultrasound Biometric Scanner; Model: PIROP	Ultrasound Biometric Scanner	C (P-156)	Non-invasive, modern device for fast and accurate tissue thickness measurements. It allows to measure and monitor of periodontal mucosa thickness in prophylaxis and periodontal treatment. Identification of a periodontal biotype in patients has a fundamental meaning in optimal planning of preventive and therapeutic management mainly inperiodontology, orthodontics, implantologyand dental prosthetics.	**Contraindication:** Not Known. **Side Effects:** None	India EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
76.	**Manufacturer:** GE Vingmed Ultrasound AS Strandpromenaden 45 N-3191 Horten Norway **Local Agent**: GE Healthcare Bangladesh Ltd. Shanta Western Tower, OS # 804, Level # 08, 186 Tejgaon I/ A Dhaka	Vivid E9	Ultrasound system, Imaging, cardiovascular	C	It is a general-purpose ultrasound system, specialized for usein cardiac imaging. It is intended for use by, or under the direction of a qualified physicianfor ultrasound imaging and analysis of Fetal; Abdominal (including renal and GYN); Pediatric; Small Organ (breast, testes, thyroid); Neonatal Cephalic; Adult Cephalic; Cardiac(adult and pediatric); Peripheral Vascular (PV); Musculo-skeletal Conventional; Urology(including prostate), Transesophageal; Transrectal (TR); Transvaginal (TV); andIntraoperative (abdominal, thoracic, & vascular).	**Contraindication:** The products are not intended for ophthalmic use or any use causing the acoustic beam to pass through the eye. **Side effect:** None	Norway EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

Sl No	c¯ZKvi‡Ki bvg	ewYR¨K bvg	‡gvW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve Kwgkb Ui mfvi wm×vš	mfvi wm×vš
77.	**Manufacturer:** GE Medical Systems (China) Co., Ltd. China **Local Agent**: GE Healthcare Bangladesh Ltd. Shanta Western Tower, OS # 804, Level # 08, 186 Tejgaon I/ A Dhaka	Vivid T8	General Purpose Ultrasound imaging and analysis system	C	The Vivid TB/Vivid T8 Pro is Multipurpose Cardiovascular System designed for cardiac and Shared Service Imaging, the system supports following applications: Fetal/OB, Abdominal, Pediatric, Small Organ, Cardiac, Peripheral Vascular, Adult Cephalic, Neonatal Cephalic, Musculoskeletal Superficial/Conventional, Transcranial, Transrectal, Transvaginal and Transesophageal.	**Contraindication:** The products are not intended for ophthalmic use or any use causing the acoustic beam to pass through the eye. **Side effect:** None	China USA EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |
78.	**Manufacturer:** GE Helathcare Austria GmbH & Co OG, Tiefenbach 15, 4871 Zipf/Austria **Local agent:** GE Healthcare Bangladesh Ltd., Shanta Western Tower, 186, Tejgaon. Dhaka	VOLUSON E6/E8/E8 Expert/E10	Ultrasound System	C	The device is a general purpose ultrasound system. Specific clinical applications remain the same as previously cleared: 20 GE Healthicare 510(k) Premarket Notification Submission Fetal/OS; Abdominal (including GYN, pelvic and infertility monitoring/follicle development); Pediatric; Small Organ (breast, testes, thyroid etc.); Neonatal and Adult Cephalic; Cardiac (adult and pediatric); Musculo-skeletal Conventional and Superficial; Peripheral Vascular Tranavaginal; Transrectal; and Intraoperative (abdominal, PV and neurological).	Contraindications: It is not intended for: • Ophthalmic use or any use where the probe is directly applied to the eye. • intra-operative use **Side effect:** None	Austria	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |
79.	**Manufacturer:** GE Ultrasound Korea, Ltd., Republic of Korea **Local Agent**: GE Healthcare Bangladesh Ltd. Shanta Western Tower, OS # 804, Level # 08, 186 Tejgaon I/ A Dhaka	VOLUSON S6/S8/S10	General Purpose Ultrasound Imaging System	C	The device is a general – purpose ultrasound system. Specific clinical application and exam types include: Fetal (Obstetrics); Abdominal (including renal and GYN/pelvic); Pediatric; Small Organ (breast, testes, thyroid, salivary glands, lymph nodes, pediatric and neonatal patients); Neonatal Cephalic; Adult Cephalic; Cardiac (adult and pediatric); Peripheral Vascular (PV); Musculo-Skeletal Conventional and Superficial; Transcrectal (TR); Transvaginal (TV).	**Contraindication:** The System is not intended for ophthalmic use or a use causing the acoustic beam to pass through the eye. The System is not intended for intra-operative use that is defined as introducing aprobe into a surgical incision or burr hole. Abdominal and Linear probes are not intended for endocavity use. IOTA LR2 model measurement tool should only be used with women with adnexaltumors that have been selected to undergo surgery. IOTA states that use outside of the intended population can overestimate risk of a malignant tumor. **Side effect:** None	Korea USA EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |

Sl No	c¯ZKvi‡Ki bvg	ewYR¨K bvg	‡gvW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve Kwgkb mfvi wm×vš	mfvi wm×vš
80.	**Manufacturer:** Boston Scientific Limitwd Ballybrit Business park. Galway, Ireland **Local Agent:** M/s. Medi Graphic Trading Ltd., 14 Purana Paltan, Dhaka, Bangladesh	2cm Peripheral Cutting Balloon Microsurgical Dilatation Device	Peripheral Cutting Balloon	D	It is indicated for Percutaneous Transluminal Angioplasty (PTA) of obstructive lesions in peripheral vessels. The target lesion should possess the following characteristics: minimal tortuosity of proximal vessel segment, and be a non-angulated lesion segment (≤ 45°).	**Contraindications:** Use of the Cutting Balloon Device is contraindicated in situations where the Cutting Balloon Device would be passed through the struts of a previously placed stent as the deflated Cutting Balloon Device could become entangled in the stent. **Adverse Events:** The Peripheral Cutting Balloon Device is not for use in the coronary arteries and carotid arteries, The Peripheral Cutting Balloon Device is not intended for the expansion or delivery of stents.	Ireland EC Certificate	Abjgv`b Kiv †h‡Z cv‡i |	Abjgv`b Kiv nj |
81.	**Manufacturer:** Smith & Nephew Medical Limited, UK **Local Agent:** UniMed Ltd., House No. 46, Sheikh Kamal Saroni, Road No. 16, Rangs Nasim (7th Floor) Square, Dhanmondi	**Acticoat**	Silver-coated Antimicrobial Barrier Wound dressings	D (P-30)	It is an antimicrobial barrier layer over partial and full-thickness wounds such as: first and second degree burns and donor and recipient graft sites, pressure ulcers, venous ulcers, and diabetic ulcers.	**Contraindications:** Not known **Side Effects:** Not known	MHRA-UK EC Certificate	Abjgv`b Kiv †h‡Z cv‡i |	Abjgv`b Kiv nj |
82.	**Manufacturer:** Smith & Nephew Medical Limited, UK **Local Agent:** UniMed Ltd., House No. 46, Sheikh Kamal Saroni, Road No. 16, Rangs Nasim (7th Floor) Square, Dhanmondi	Allevyn Ag Heel	Absorbent Silver Barrier Dressing	D	ALLEVYN Ag Heel Absorbent Silver Barrier Dressing is indicated for exudate absorption and the management of partial to full thickness wounds. Some typical wounds are: ➢ Ulcers (venous, arterial, diabetic) ➢ Pressure Sores ➢ Donor Sites ➢ Surgical Incisions ➢ Surgical Excisions ➢ Burns (1st and 2nd degree) Creation and maintenance of a moist wound environment. Moist wound environments have been established as optimal environments for the management of the wound. Provides physical separation between the wound and external environments to assist in preventing bacterial contamination of the wound.	Contraindications: Not known Side Effects: Not known **Precautions** For external use only ALLEVYN Ag Heel is not compatible with oxidizing agents (e.g. EUSOL) or hydrogen peroxide, as these can break down the absorbent polyurethane component of the dressing ALLEVYN Ag Heel dressings are not intended to provide treatment for infected wounds. ALLEVYN Ag Heel may be used on infected wounds being managed in accordance with institutional clinical protocols for infection abatement as an adjunct to the standard treatment regimen to provide a barrier to bacterial penetration. ALLEVYN Ag Heel may not be compatible with topical anti-microbials Reddening of the skin around the wound may be observed. This may be related to irritation of fragile skin or wound exudates remaining in contact with normal skin for prolonged periods of time. If reddening or	MHRA-UK EC Certificate	Abjgv`b Kiv †h‡Z cv‡i |	Abjgv`b Kiv nj |

						sensitization occur, discontinue use. Avoid contact with electrodes or conductive gels during electronic measurements e.g. EEG and ECG. When ALLEVYN Ag Non Adhesive is used on a patient during MRI (Magnetic Resonance Imaging) examination some warming may be experienced ALLEVYN Ag Heel is not compatible with oil based products such as petrolatum. ALLEVYN Ag Heel is not suitable for use on cavity wounds but may be used over cavity wounds as a secondary dressing. As with all products containing silver sulfadiazine the following precautions should be noted, especially when covering a large surface area with the dressing, 1. Use caution in patients with significant hepatic or renal impairment, 2. Use caution in individuals known to have glucose-6-phosphate dehydrogenase deficiency, 3. Effects of systemically administered drugs may be altered. This can especially apply to oral hypoglycaemic agents and to phenytoin. In the case of these drugs, it is recommended that blood levels should be monitored as their effect can be potentiated.			
83.	**Manufacturer:** Eucare Pharmaceuticals Pvt Ltd., INDIA **Local Agent:** Kulic Limited 9/7, Eastern Plaza (8th Floor), 70, BirUttom C.R Datta Road, Hatirpul, Dhaka	**BIOFIL-AB**-Sterile Medicated collagen particles (Fish Origin) Collagen with Mupirocin USP 2% w/w Metronidazole I%	Biological Skin Dressing	D	• Biofil-AB particles are intended specifically to be used as collagen-based wound dressing for wound management. • Non-healing ulcers • Diabetic foot ulcers • Pressure ulcers • Venous ulcers • Infected and non-infected wounds • Traumatic wounds Minimal to heavily draining wounds • Surgical wounds • Tunneled and undermined wounds	**PRECAUTIONS & WARNINGS:** • Wound may appear larger during the first several days of treatment due to the reduction of edema • An increase in drainage may be seen in the first several days of treatment • Treatment of the underlying conditions (Venous or arterial flow, external pressure etc.) as well as nutritional support should proceed concurrently with the use of BioFil®-AB Particles • All opened and unused portions of BioFil®-AB Particles must be discarded • Avoid on patients hypersensitive to Collagen. Mupirocin and Metronidazole • Avoid topical agents like Povidone Iodine, H2O2, and Sodium Hypochlorite for cleansing the wound as their toxicity level may hamper the new tissue growth • BioFil®-AB Particles is a single use product, which should not be re-sterilized	India EC Certificate	Abjgv`b Kiv †h‡Z cv‡i	Abjgv`b Kiv nj

Sl No	c¯ZKvi‡Ki bvg	ewYRK bvg	‡gvW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve KwgwUi mfvi wm×vš	mfvi wm×vš
84.	**Manufacturer:** Boston Scientific Limitwd Ballybrit Business park. Galway, Ireland **Local Agent:** M/s. Medi Graphic Trading Ltd., 14 Purana Paltan, Dhaka, Bangladesh	Blazer II HTD Temperature ablation Catheter	Ablation catheter	D	It is indicated for interruption of accessory atrioventricular conduction pathways associated with tachycardia, for treatment of AV nodal reentrant tachycardia and for creation of complete AV block in patients with a rapid ventricular response to an atrial arrhythmia - typically chronic, drug refractory atrial fibrillation.	**Contraindications:** The use of this device is contraindicated in patients who have vena cava embolic protection filters devices and/or known femoral thrombus who require catheter insertion from the femoral approach. The use of the device is contraindicated in patients with active systemic infection. The transseptal approach is contraindicated in patients with left atrial thrombus or myxoma, or interatrial baffle or patch. The retrograde transaortic approach is contraindicated in patients with aortic valve replacement. **Adverse Event:** Allergic reaction, Arrhythmias (new or exacerbation of existing arrhythmias) and complications of sedative agents/anesthesia, Cardiac or respiratory arrest, Cardiac valve damage, Catheter entrapment/entanglement, Chest pain, Damage to vessel intima or cardiac ultrastructures, Death, Embolus, air embolus, Hematoma/ecchymosis, Hemorrhage, Hypotension, Infection, Myocardial infarction, Perforation, Pericardial/pleural effusion, Pericarditis/pleuritis, Phrenic or intercostal nerve damage, Pneumothorax, Pseudoaneurysm, Pulmonary edema, Sinus or AV node injury, Skin burns, Stroke, Stroke or cerebral vascular accident, Tamponade, Thrombosis, Vasovagal reaction, X-ray exposure	Ireland EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

Sl No	c¯ZKvi‡Ki bvg	ewYRK bvg	‡gvW‡Kj vWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve Kwgwi mfvi wm×vš	mfvi wm×vš
85.	**Manufacturer:** Boston Scientific Limitwd Ballybrit Business park. Galway, Ireland **Local Agent:** M/s. Medi Graphic Trading Ltd., 14 Purana Paltan, Dhaka, Bangladesh	Blazer II Temperature ablation Catheter	Ablation catheter	D (P-45)	It is indicated for interruption of accessory atrioventricular conduction pathways associated with tachycardia, for treatment of AV nodal reentrant tachycardia and for creation of complete AV block in patients with a rapid ventricular response to an atrial arrhythmia - typically chronic, drug refractory atrial fibrillation.	**Contraindications:** The use of this device is contraindicated in patients who have vena cava embolic protections filter devices and/or known femoral thrombus, who requires catheter insertion from the femoral approach. The use of the device is contraindicated in patients with active systemic infection. The transseptal approach is contraindicated in patients with left atrial thrombus or myxoma, or interatrial baffle or patch. The retrograde transaortic approach is contraindicated in patients with aortic valve replacement. **Adverse Event:** Allergic reaction, Arrhythmias (new or exacerbation of existing arrhythmias) and complications of sedative agents/anesthesia, Cardiac or respiratory arrest, Cardiac valve damage, Catheter entrapment/entanglement, Chest pain, Damage to vessel intima or cardiac ultrastructures, Death, Embolus, air embolus, Hematoma/ecchymosis, Hemorrhage, Hypotension, Infection, Myocardial infarction, Perforation, Pericardial/pleural effusion, Pericarditis/pleuritis, Phrenic or intercostal nerve damage, Pneumothorax, Pseudoaneurysm, Pulmonary edema, Sinus or AV node injury, Skin burns, Stroke, Stroke or cerebral vascular accident, Tamponade, Thrombosis, Vasovagal reaction, X-ray exposure	Ireland EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

Sl No	c¯ZKv i‡Ki bvg	ewYRK bvg	‡gvW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve Kwgkb mfvi wm×vš	mfvi wm×vš
86.	**Manufacturer:** Boston Scientific Limitwd Ballybrit Business park. Galway, Ireland **Local Agent:** M/s. Medi Graphic Trading Ltd., 14 Purana Paltan, Dhaka, Bangladesh	Blazer II XP Temperature ablation Catheter	Ablation catheter	D	It is indicated for interruption of accessory atrioventricular conduction pathways associated with tachycardia, for treatment of AV nodal reentrant tachycardia, for treatment of atrial flutter tachycardia, and for creation of complete AV block in patients with a rapid ventricular response to an atrial arrhythmia – typically chronic, drug refractory atrial fibrillation.	**CONTRAINDICATIONS:** The use of this device is contraindicated in patients who have vena cava embolic protection filter devices and/or known femoral thrombus who require catheter insertion from the femoral approach.The use of the device is contraindicated in patients with active systemic infection. The transseptal approach is contraindicated in patients with left atrial thrombus or myxoma, or interatrial baffle or patch. The retrograde transaortic approach is contraindicated in patients with aortic valve replacement. **Adverse Event:** Allergic reaction, Arrhythmias (new or exacerbation of existing arrhythmias) and complications of sedative agents/anesthesia, Cardiac or respiratory arrest, Cardiac valve damage, Catheter entrapment/entanglement, Chest pain, Damage to vessel intima or cardiac ultrastructures, Death, Embolus, air embolus, Hematoma/ecchymosis, Hemorrhage, Hypotension, Infection, Myocardial infarction, Perforation, Pericardial/pleural effusion, Pericarditis/pleuritis, Phrenic or intercostal nerve damage, Pneumothorax, Pseudoaneurysm, Pulmonary edema, Sinus or AV node injury, Skin burns, Stroke, Stroke or cerebral vascular accident, Tamponade, Thrombosis, Vasovagal reaction, X-ray exposure	Ireland EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

Sl No	c¯ZKvi‡Ki bvg	ewYRK bvg	‡gvW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve KwgwUi mfvi wm×vš	mfvi wm×vš
87.	**Manufacturer:** Boston Scientific Limitwd Ballybrit Business park. Galway, Ireland **Local Agent:** M/s. Medi Graphic Trading Ltd., 14 Purana Paltan, Dhaka, Bangladesh	Blazer™ Open-Irrigated Ablation Catheter	Ablation catheter	D	It is indicated for use in catheter-based cardiac electrophysiological mapping (stimulating and recording) and, when used in conjunction with a radiofrequency generator, for cardiac ablation.recording) and, when used in conjunction with a radiofrequency generator, for cardiac ablation.	**Contraindications:** The Blazer Open-Irrigated Ablation Catheter is contraindicated for use: in patients with active systemic infection, in patients with a mechanical prosthetic heart valve through which the catheter must pass, in patients with left ventricular thrombus; or with left atrial thrombus or myxoma via the transseptal approach, in patients who are unable to receive heparin or an acceptable alternative to achieve adequate anticoagulation, in patients who have vena cava embolic protection filter devices and/or known femoral thrombus who require catheter insertion from the femoral approach, in patients who are hydrodynamically unstable, in patients who have myxoma or an intracardiac thrombus, in patients who have had a ventriculotomy or atriotomy within the preceding eight weeks, via transseptal approach in patients with interatrial baffle or foramen ovale patches, via retrograde transaortic approach in patients who have had aortic valve replacement. **Adverse Event:** Allergic reaction, Arrhythmias (new or exacerbation of existing arrhythmias) and complications of sedative agents/anesthesia, Cardiac or respiratory arrest, Cardiac valve damage, Catheter entrapment/entanglement, Chest pain, Damage to vessel intima or cardiac ultrastructures, Death, Embolus, air embolus, Hematoma/ecchymosis, Hemorrhage, Hypotension, Infection, Myocardial infarction, Perforation, Pericardial/pleural effusion, Pericarditis/pleuritis, Phrenic or intercostal nerve damage, Pneumothorax, Pseudoaneurysm, Pulmonary edema, Sinus or AV node injury, Skin burns, Stroke, Stroke or cerebral vascular accident, Tamponade, Thrombosis, Vasovagal reaction, X-ray exposure	Ireland EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

Sl No	c¯ZKvi‡Ki bvg	ewYwR¨K bvg	‡gwW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve KwgwUi mfvi wm×vš	mfvi wm×vš
88.	**Manufacturer:** MicroVention, Inc. 1311 Valencia Ave Tustin, CA 92780 USA **Local Agent:** M/S, UniTrade Corporation 2/5, Humayun Road (3rdFl), Block-B Mohammadpur, Dhaka-1207	Chaperon Guiding Catheter system	Guiding Catheter for Cerebral Aneurysm	**D**	Chaperon Guiding Catheter is intended for General intravascular use, including the neuro and peripheral vasculature. It can be used to facilitate Introduction of diagnostic or therapeutic devices. It is not intended for use in coronary arteries.	**Contraindication:** There are no known contraindications. **Side Effect:** None	USA EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
89.	**Manufacturer:** Boston Scientific Corporation, 300 Boston Scientific Way, Marlborough, MA 01752, USA **Local Agent:** M/s. Medi Graphic Trading Ltd., 14 Purana Paltan, Dhaka, Bangladesh	Comet Pressure Guidewire	Pressure Guidewire	D	It is indicated to direct a catheter through a blood vessel and to measure physiological parameters in the coronary blood vessels.	Contraindications: The Comet Pressure Guidewire is contraindicated for use in the cerebral vasculature. **Adverse events**: Potential adverse events which may result from the use of the device include but are not limited to: Abrupt closure, Allergic reaction, Embolism, Exposure to biohazardous material, Infection, Prolonged procedure, Restenosis (reocclusion), Spasm, Stroke/cerebral vascular accident, (CVA)/transient ischemic attack (TIA), Vascular thrombus, Vessel trauma (dissection, perforation, rupture or injury) In addition, when used for interventional procedures: Angina or unstable angina, Arrhythmias, Cardiac tamponade/pericardial effusion, Contrast induced renal insufficiency or renal failure, Death, Myocardial infarction or ischemia. Some of the above potential adverse events may require additional surgical intervention.	USA EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

Sl No	c¯ZKvi‡Ki bvg	ewYwR¨K bvg	‡gvW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve Kwgkb mfvi wm×vš	mfvi wm×vš
90.	**Manufacturer:** Boston Scientific Limitwd Ballybrit Business park. Galway, Ireland **Local Agent:** M/s. Medi Graphic Trading Ltd., 14 Purana Paltan, Dhaka, Bangladesh	Crossboss Guide catheter	Vascular Guide Catheter	D	It is intended for use with a guidewire to access discrete regions of the coronary and peripheral vasculature. When used as part of the system consisting of the CrossBoss Catheter, Stingray™ Catheter, and Stingray Guidewire, the CrossBoss Catheter is indicated for use to facilitate the intraluminal placement of conventional guidewires beyond stenotic coronary lesions (including chronic total occlusions [CTOs]) prior to PTCA or stent intervention.	**Contraindications:** Do not use with guidewire extension systems with a coupling profile larger than 0.014 in (0.36 mm) diameter (i.e. wave pattern Coupling mechanism). **Adverse Events:** Potential adverse events include, but are not limited to, the following, Acute myocardial infarction, Vessel trauma requiring surgical repair or intervention, Hemorrhage or hematoma, Artery spasm, Embolism, Stroke, Neurological deficit, Drug reactions, allergic reaction to contrast media , Infection CrossBos	Ireland EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
91.	**Contract manufacture** ADESHWAR MEDITEX PVT LTD, INDIA **Local agent:** Janata Traders TCB Bhabon, 1 kawran Bazar,Dhaka	**Cuticell Plus** (Antibiotic Tulle Dressing) (Sterile)	Antibiotic Tulle Dressing Each GM Containing: Neomycin Sulphate USP 3400 Unit + Polymyxin B Sulphate USP 5000 units-Bacitracine Zinc USP 400 Unit/	D (P-30)	For superficial exudating wounds such as ➢ First and second-degree burns ➢ Cuts and abrasions ➢ Lacerations ➢ Radiation injuries ➢ Donor and recipient skin graft sites	**Contraindication:** Not Known. **Side Effects:** None	India	‡iwR‡ókb MvBWjvBb Abyjvqx c‡qvRbxq WKzgU `wLj Kivi Rb¨ ejv †h‡Z cv‡i	‡iwR‡ókb MvBWjvBb Abyjvqx c‡qvRbxq WKzgU `wLj Kivi Rb¨ ejv nj

Sl No	c¯ZKvi‡Ki bvg	ewYRK bvg	‡gvW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve Kwgkb mfvi wm×vš	mfvi wm×vš
92.	**Manufacturer:** Boston Scientific Corporation, 300 Boston Scientific Way, Marlborough, MA 01752, USA **Local Agent:** M/s. Medi Graphic Trading Ltd., 14 Purana Paltan, Dhaka, Bangladesh	Dynamic Tip Unidirectional Steerable Diagnostic Catheter	Unidirectional Steerable Diagnostic Catheter	D	It is intended for temporary Intracardiac sensing, recording, stimulation and temporary pacing during the evaluation of cardiac arrhythmias.	**Contraindications:** The transseptal approach is contraindicated in patients with left atrial thrombus or myxoma, or interatrial baffle patch. The retrograde transaortic approach is contraindicated in patients with aortic valve replacement. **Adverse Event:** Allergic reaction, Arrhythmias (new or exacerbation of existing arrhythmias) and complications of sedative agents/anesthesia, Cardiac or respiratory arrest, Cardiac valve damage, Catheter entrapment/entanglement, Chest pain, Damage to vessel intima or cardiac ultrastructures, Death, Embolus, air embolus, Hematoma/ecchymosis, Hemorrhage, Hypotension, Infection, Myocardial infarction, Perforation, Pericardial/pleural effusion, Pericarditis/pleuritis, Phrenic or intercostal nerve damage, Pneumothorax, Pseudoaneurysm, Pulmonary edema, Sinus or AV node injury, Skin burns, Stroke, Stroke or cerebral vascular accident, Tamponade, Thrombosis, Vasovagal reaction, X-ray exposure	USA EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
93.	**Manufacturer:** Boston Scientific Corporation, 300 Boston Scientific Way, Marlborough, MA 01752, USA **Local Agent:** M/s. Medi Graphic Trading Ltd., 14 Purana Paltan, Dhaka, Bangladesh	Dynamic XT Unidirectional Steerable Diagnostic Catheter	Unidirectional Steerable Diagnostic Catheter	D	It is intended for temporary Intracardiac sensing, recording, stimulation and temporary pacing during the evaluation of cardiac arrhythmias.	**Contraindications:** The transseptal approach is contraindicated in patients with left atrial thrombus or myxoma, or interatrial baffle patch. The retrograde transaortic approach is contraindicated in patients with aortic valve replacement. **Adverse Event:** Allergic reaction, Arrhythmias (new or exacerbation of existing arrhythmias) and complications of sedative agents/anesthesia, Cardiac or respiratory arrest, Cardiac valve damage, Catheter entrapment/entanglement, Chest pain, Damage to vessel intima or cardiac ultrastructures, Death, Embolus, air embolus, Hematoma/ecchymosis, Hemorrhage, Hypotension, Infection, Myocardial infarction, Perforation, Pericardial/pleural effusion, Pericarditis/pleuritis, Phrenic or intercostal nerve damage, Pneumothorax, Pseudoaneurysm, Pulmonary edema, Sinus or AV node injury, Skin burns, Stroke, Stroke or cerebral vascular accident, Tamponade, Thrombosis, Vasovagal reaction, X-ray exposure	USA EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

Sl No	c¯ZKvi‡Ki bvg	ewYRK bvg	‡gvW‡Kj vWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve Kvgvwi mfvi wm×vš	mfvi wm×vš
94.	**Manufacturer:** Boston Scientific Corporation, 300 Boston Scientific Way, Marlborough, MA 01752, USA **Local Agent:** M/s. Medi Graphic Trading Ltd., 14 Purana Paltan, Dhaka, Bangladesh	FilterWire EZ Embolic Protection System	Embolic Protection system	D	It is indicated for use as a guidewire and embolic protection system to contain and remove embolic material thrombus/debris) while performing angioplasty and stenting procedures in coronary saphenous vein bypass grafts and carotid arteries. The diameter of the vessel at the site of filter loop lacement should be between 2.25 mm and 5.5 mm for coronary saphenous vein bypass graft procedures and between 3.5 mm and 5.5 mm for carotid procedures.	**Contraindications:** Patients with severe allergy to heparin Patients with bleeding diathesis or other disorders that limit the use of anticoagulant therapy **Adverse Event:** Angina, Bleeding complications, Bradycardia or arrhythmias, including ventricular fibrillation or Tachycardia, Congestive heart failure, Damage to or dislocation of the implanted stent(s), Death, Detachment and/or implantation of a component of the system, Drug reaction, allergic reaction to contrast media, medications or device materials, Embolization of air, tissue, thrombus or other embolic debris, Emergent surgery,End organ ischemia/infarction, Headache Hypotension/hypertension, Infection (local or systemic), Myocardial infarction, No-reflow resulting from reduced blood flow through the FilterWire EZ System filter, Pain, Puncture site complications (i.e., vessel occlusion, hemorrhage,hematoma, pseudoaneurysm or arteriovenous fistula), Renal insufficiency, kidney failure, hematuria, Stroke/cerebrovascular accident (CVA), transient ischemic attack (TIA) or seizure, Vessel damage, dissection, occlusion, aneurysm, perforation, rupture, injury, thrombosis, or spasm	USA EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

Sl No	c¯ZKvi‡Ki bvg	ewYwR¨K bvg	‡gwW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve KwgwUi mfvi wm×vš	mfvi wm×vš
95.	**Manufacturer:** B. Braun Surgical, S.A, Spain **Local Agent:** Asia Pacific Medical Ltd., 775, Satmasjid Road (2nd Floor), Dhanmondi, Dhaka	Gelita Tampon	Local Gelatin Haemostat	**D**	**Indicated in:** ➢ **Dentistry and Maxilar surgery:** To fill dead spaces and control bleeding ➢ **Thoracic surgery:** To fill dead spaces after lymph node removal ➢ Ear Nose **Throat Surgery:** To support the tympanic membrane in tympanoplasties and to control nose bleeding (epistaxis, after nasal polyps resection, etc) ➢ To stop venous-capillary or parenchymatous haemorrhage in: Neurosurgery, Orthopedic surgery, Gynecology, Vascular Surgery, General Surgery and Urology	**Contraindication:** ➢ Gelita Tampon may not be applied to infected area , since it can increase the severity of infections. ➢ It should not be use d in conjunction with cemented endoprotheses since it reduces the adhesive strength of the bone cement. ➢ Known hypersensitivity to product. **Side Effects:** Tissue adhesion can occur occasionally.	Spain EC certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
96.	**Manufacturer:** Boston Scientific Corporation, 300 Boston Scientific Way, Marlborough, MA 01752, USA **Local Agent:** M/s. Medi Graphic Trading Ltd., 14 Purana Paltan, Dhaka, Bangladesh	Guidezilla Guide Extension catheter	Guide Extension catheter	D	It is intended to be used in conjunction with guide catheters to access discrete regions of the coronary and/or peripheral vasculature, and to facilitate placement of interventional devices.	**Contraindications:** Vessels less than 2.5 mm in diameter. Vessels in the neurovasculature and the venous system… **Adverse events**:	USA EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
97.	**Manufacturer:** MicroVention, Inc. 1311 Valencia Ave Tustin, CA 92780 USA **Local Agent:** M/S, UniTrade Corporation 2/5, Humayun Road (3rdFl), Block-B Mohammadpur, Dhaka-1207	Headway Microcatheter	Microcatheter for Cerebral Aneurysm	**D**	It is intended for general intravascular use, including the peripheral, coronary and neuro vasculature for the infusion of diagnostic agents, such as contrast media, and therapeutic agents, such as occlusion coils.	**Contraindication:** There are no known contraindications. **Side Effects:** None	USA EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

Sl No	c¯ZKvi‡Ki bvg	ewYRK bvg	‡gvW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve Kwgkb mfvi wm×vš	mfvi wm×vš
98.	**Manufacturer:** MicroVention, Inc. 1311 Valencia Ave Tustin, CA 92780 USA **Local Agent:** M/S, UniTrade Corporation 2/5, Humayun Road (3rdFl), Block-B Mohammadpur, Dhaka-1207	HydroCoil Embolic System (HES)	Embolic System for Cerebral Aneurysm	**D**	It is intended for the endovascular embolization of intracranial aneurysms and other neurovascular abnormalities such as arteriovenous malformations and arteriovenous fistulae. The HES is also intended for vascular occlusion of blood vessels within the neurovascular system to permanently obstruct blood flow to an aneurysm or other vascular malformation and for arterial and venous embolization in The peripheral vasculature.	**Contraindication:** There are no known contraindications. Side Effect: None	USA EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
99.	**Manufacturer:** Boston Scientific Corporation, 300 Boston Scientific Way, Marlborough, MA 01752, USA **Local Agent:** M/s. Medi Graphic Trading Ltd., 14 Purana Paltan, Dhaka, Bangladesh	IntellaMap Orion High Resolution Mapping Catheter	Mapping Catheter	D	It is indicated for electrophysiological mapping (recording or stimulating only) of the cardiac structures of the heart.	**Contraindications:** The IntellaMap Orion Catheter should not be used in: Patients who are not candidates for transvascular catheter procedures. Patients with a hypercoagulable state or who cannot tolerate heparin anticoagulation therapy. Patients with prosthetic or stenotic valves, in the chamber where the prosthetic or stenotic valve reside. Patients with active systemic infection. Pediatric patients. Pregnant and/or nursing patients. Patients with any other condition where catheter manipulation may not be safe. The IntellaMap Orion™ Catheter should not be used for radio frequency (RF) ablation. The IntellaMap Orion Catheter should not be used inside an MRI machine. **Adverse Event:** Allergic reaction, Arrhythmias (new or exacerbation of existing arrhythmias) and complications of sedative agents/anesthesia, Cardiac or respiratory arrest, Cardiac valve damage, Catheter entrapment/entanglement, Chest pain, Damage to vessel intima or cardiac ultrastructures, Death, Embolus, air embolus, Hematoma/ecchymosis, Hemorrhage, Hypotension, Infection, Myocardial infarction, Perforation, Pericardial/pleural effusion, Pericarditis/pleuritis, Phrenic or intercostal nerve damage, Pneumothorax, Pseudoaneurysm, Pulmonary edema, Sinus or AV node injury, Skin burns, Stroke, Stroke or cerebral vascular accident, Tamponade, Thrombosis, Vasovagal reaction, X-ray exposure	USA EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

Sl No	c¯ZKvi‡Ki bvg	ewYwR¨K bvg	‡gvW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve Kwgkb mfvi wm×vš	mfvi wm×vš
100.	**Manufacturer:** Eucare Pharmaceuticals Pvt Ltd., INDIA **Local Agent:** Kulic Limited 9/7, Eastern Plaza (8th Floor), 70, BirUttom C.R Datta Road, Hatirpul, Dhaka	**KOLLAGEN** Sterile collagen sheets **Origin:** Bovine	Biological Skin Dressing	D (P-73)	➢ Kollagen is intended specifically to be used as collagen-based wound dressing for wound management. ➢ 2nd degree superficial and deep dermal burns ➢ 3rd degree burns as a temporary cover after Escharectomy/ Tangential excision ➢ Skin Donor Sites ➢ Traumatic loss of skin cover ➢ Temporary wound cover in major open fractures before definitive flap cover ➢ Chronic skin ulcers ➢ Shallow pressure sores ➢ Leprosy ulcers ➢ Dermabrasion areas ➢ Protective cover over widely meshed autografts	**PRECAUTIONS & WARNINGS:** ➢ Grossly infected wound may reject the collagen cover ➢ On application of Collagen sheet on infected sites, care should be taken to inspect the wound condition frequently and change of dress is recommended after cleansing with normal saline. ➢ Avoid in patients hypersensitive to Collagen ➢ The product will not be sterile when the packing is damaged during transit ➢ Store at normal room temperatures.	India	‡iwR‡ókb MvBWjvBb Abjyqx c‡qvRbxq WKzgU `wLj Kivi Rb¨ ejv †h‡Z cv‡i	‡iwR‡ókb MvBWjvBb Abjyqx c‡qvRbxq WKzgU `wLj Kivi Rb¨ ejv nj
101.	**Manufacturer:** Boston Scientific Corporation, 300 Boston Scientific Way, Marlborough, MA 01752, USA **Local Agent:** M/s. Medi Graphic Trading Ltd., 14 Purana Paltan, Dhaka, Bangladesh	Leveen Standard Needle Electrode System	Leveen Electrode System	D	The LeVeen Needle Electrode Family is intended to be used in conjunction with the RF 3000 Generator for the thermal coagulation necrosis of soft tissues, including partial or complete ablation of nonresectable liver lesions. These procedures should only be performed by physicians and staff familiar with the equipment and techniques involved.	**Contraindications:** Before using, inspect the package for any breach to the sterile barrier and inspect product for any damage. If package is broken or product is damaged DO NOT USE. Immediately return package and product to Boston Scientific. **Adverse Events:** Abscess, ARDS (Acute Respiratory Distress Syndrome), Arrhythmia, Ascites, Biloma, Burn, Death, Delayed hemorrhage into ablated tissue, Diarrhea, Electric Shock, Fistula, including biliary fistula, Hematoma, Hemorrhage, Infection, Liver Failure, Liver Insufficiency, Pain, Perforation, Peritonitis, Persistent Fever > 39° C, Pleural Effusion, Renal Failure, Tumor Recurrence, Tumor Seeding	USA EC Certificate	Abjyqv`b Kivi †h‡Z cv‡i	Abjyqv`b Kivi nj

Sl No	c¯ZKvi‡Ki bvg	ewYRK bvg	‡gvW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve KwgwUi mfvi wm×vš	mfvi wm×vš
102.	**Manufacturer:** Boston Scientific Corporation, 300 Boston Scientific Way, Marlborough, MA 01752, USA **Local Agent:** M/s. Medi Graphic Trading Ltd., 14 Purana Paltan, Dhaka, Bangladesh	Leveen Super slim Needle Electrode System	Radiofrequency Ablation Needle Electrode	D	It is intended to be used in conjunction with the RF 3000 Generator for the thermal coagulation necrosis of soft tissues, including partial or complete ablation of nonresectable liver lesions. These procedures should only be performed by physicians and staff familiar with the equipment and techniques involved.	**Contraindications:** The LeVeen Needle Electrode Family is intended to be used in conjunction with the RF 3000 Generator for the thermal coagulation necrosis of soft tissues, including partial or complete ablation of nonresectable liver lesions. These procedures should only be performed by physicians and staff familiar with the equipment and techniques involved. **Adverse Events:** Abscess, ARDS (Acute Respiratory Distress Syndrome), Arrhythmia, Ascites, Biloma, Burn, Death, Delayed hemorrhage into ablated tissue, Diarrhea, Electric Shock, Fistula, including biliary fistula, Hematoma, Hemorrhage, Infection, Liver Failure, Liver Insufficiency, Pain, Perforation, Peritonitis, Persistent Fever > 39° C, Pleural Effusion, Renal Failure, Tumor Recurrence	USA EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
103.	**Manufacturer:** MicroVention, Inc. 1311 Valencia Ave Tustin, CA 92780 USA **Local Agent:** M/S, UniTrade Corporation 2/5, Humayun Road (3rdFl), Block-B Mohammadpur, Dhaka-1207	MicroPlex Coils System (MCS)	Coiling System for Cerebral Aneurysm	**D**	It is intended for the endovascular embolization of intracranial aneurysms and other neurovascular abnormalities such as arteriovenous malformations and arteriovenous fistulae. The MCS is also intended for vascular occlusion of blood vessels within the neurovascular system to permanently obstruct blood flow to an aneurysm or other vascular malformation and for arterial and venous embolizations in the peripheral vasculature.	**Contraindication:**There are no known contraindications. Side Effect: None	USA EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

Sl No	c¯ZKvi‡Ki bvg	ewYwR¨K bvg	‡gwW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve KwgwUi mfvi wm×vš	mfvi wm×vš
104.	**Manufacturer:** Boston Scientific Corporation, 300 Boston Scientific Way, Marlborough, MA 01752, USA **Local Agent:** M/s. Medi Graphic Trading Ltd., 14 Purana Paltan, Dhaka, Bangladesh	Orbiter PV Diagnostic Mapping Catheter	Mapping Catheter	D	It is intended for temporary intracardiac sensing, recording, stimulation and temporary pacing during the evaluation of cardiac arrhythmias.	**Contraindications:** The catheter should not be used in conditions where manipulation of the catheter would be unsafe (e.g. intracardiac mural thrombus). The transseptal approach is contraindicated in patients with left atrial thrombus or myxoma, or interatrial baffle patch. The retrograde transaortic approach is contraindicated in patients with aortic valve replacement. The retrograde transaortic approach is contraindicated due to the risk of entrapping the tip in the left ventricle. **Adverse Event:** Allergic reaction, Arrhythmias (new or exacerbation of existing arrhythmias) and complications of sedative agents/anesthesia, Cardiac or respiratory arrest, Cardiac valve damage, Catheter entrapment/entanglement, Chest pain, Damage to vessel intima or cardiac ultrastructures, Death, Embolus, air embolus, Hematoma/ecchymosis, Hemorrhage, Hypotension, Infection, Myocardial infarction, Perforation, Pericardial/pleural effusion, Pericarditis/pleuritis, Phrenic or intercostal nerve damage, Pneumothorax, Pseudoaneurysm, Pulmonary edema, Sinus or AV node injury, Skin burns, Stroke, Stroke or cerebral vascular accident, Tamponade, Thrombosis, Vasovagal reaction, X-ray exposure	USA EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

Sl No	c¯ZKvi‡Ki bvg	ewYRwK bvg	‡gvW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve Kwgkb mfvi wm×vš	mfvi wm×vš
105.	**Manufacturer:** Boston Scientific Corporation, 300 Boston Scientific Way, Marlborough, MA 01752, USA **Local Agent:** M/s. Medi Graphic Trading Ltd., 14 Purana Paltan, Dhaka, Bangladesh	Orbiter ST Bidirectional Steerable Diagnostic Catheter	Bidirectional Steerable Diagnostic Catheter	D	It is intended for temporary intracardiac sensing, recording, stimulation and temporary pacing during the evaluation of cardiac arrhythmias.	**Contraindications:** The catheter should not be used in conditions where manipulation of the catheter would be unsafe (e.g. intracardiac mural thrombus). The transseptal approach is contraindicated in patients with left atrial thrombus or myxoma, or interatrial baffle patch. The retrograde transaortic approach is contraindicated in patients with aortic valve replacement. **Adverse Event:** Allergic reaction, Arrhythmias (new or exacerbation of existing arrhythmias) and complications of sedative agents/anesthesia, Cardiac or respiratory arrest, Cardiac valve damage, Catheter entrapment/entanglement, Chest pain, Damage to vessel intima or cardiac ultrastructures, Death, Embolus, air embolus, Hematoma/ecchymosis, Hemorrhage, Hypotension, Infection, Myocardial infarction, Perforation, Pericardial/pleural effusion, Pericarditis/pleuritis, Phrenic or intercostal nerve damage, Pneumothorax, Pseudoaneurysm, Pulmonary edema, Sinus or AV node injury, Skin burns, Stroke, Stroke or cerebral vascular accident, Tamponade, Thrombosis, Vasovagal reaction, X-ray exposure	USA EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
106.	**Manufacturer:** **St. Jude Medical Inc.,** 14901 Deveau PL. Minnetonka, MN USA 55345 **Local Agent:** **M/s. The Spondon Ltd**. , BSEC Bhaban, Level-9, 102 Kazi Nazrul Islam Avenue, Kawran Bazar, Dhaka-1215	Pacel Bipolar Pacing Catheter	Bipolar Pacing Catheter	D	Pacel Bipolar Pacing Catheters are indicated for use intracardiac pacing and/or ECG recording.	**Contraindication:** Patients with recurrent sepsis or with a hypercoagulable state should not be considered candidates for transvenous catheters since the catheter could serve as a focal point for septic or bland thrombus formation. In addition, patients with tricuspid valve prosthesis should not be considered for ventricular pacing. **Side-effect:** • Do not alter this device • This device should only be used with equipment that complies with international safety standards. • The presence of pre-existing left bundle branch block predisposes the patient to the risk of transient complete heart block when positioning a catheter in the right ventricle. Isolated or battery powered equipment is recommended for use with intracardiac catheters.	USA EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

Sl No	c¯ZKvi‡Ki bvg	ewYRK bvg	‡gvW‡Kj vWfvB‡mi bvg	Kvm	vb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKvbK¨vj mve Kvgvi mfvi vm×vš	mfvi vm×vš
107.	**Manufacturer:** Boston Scientific Corporation, 300 Boston Scientific Way, Marlborough, MA 01752, USA **Local Agent:** M/s. Medi Graphic Trading Ltd., 14 Purana Paltan, Dhaka, Bangladesh	Radia Bidirectional Steerable Diagnostic Catheter	Bidirectional Steerable Diagnostic Catheter	D	It is intended for temporary intracardiac sensing, recording, stimulation and temporary pacing during the evaluation of cardiac arrhythmias.	**Contraindications:** The catheter should not be used in conditions where manipulation of the catheter would be unsafe (e.g. intracardiac mural thrombus). The transseptal approach is contraindicated in patients with left atrial thrombus or myxoma, or interatrial baffle patch. The retrograde transaortic approach is contraindicated in patients with aortic valve replacement. **Adverse Event:** Allergic reaction, Arrhythmias (new or exacerbation of existing arrhythmias) and complications of sedative agents/anesthesia, Cardiac or respiratory arrest, Cardiac valve damage, Catheter entrapment/entanglement, Chest pain, Damage to vessel intima or cardiac ultrastructures, Death, Embolus, air embolus, Hematoma/ecchymosis, Hemorrhage, Hypotension, Infection, Myocardial infarction, Perforation, Pericardial/pleural effusion, Pericarditis/pleuritis, Phrenic or intercostal nerve damage, Pneumothorax, Pseudoaneurysm, Pulmonary edema, Sinus or AV node injury, Skin burns, Stroke, Stroke or cerebral vascular accident, Tamponade, Thrombosis, Vasovagal reaction, X-ray exposure	USA EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

Sl No	c¯ZKvi‡Ki bvg	ewYwR¨K bvg	‡gwW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve KwgwUi mfvi wm×vš	mfvi wm×vš
108.	**Manufacturer:** Boston Scientific Corporation, 300 Boston Scientific Way, Marlborough, MA 01752, USA **Local Agent:** M/s. Medi Graphic Trading Ltd., 14 Purana Paltan, Dhaka, Bangladesh	Rotalink Burr	Exchangeable Burr Catheter	D	Percutaneous rotational coronary angioplasty with the Rotablator Rotational Atherectomy System, as a sole therapy or with adjunctive balloon angioplasty, is indicated in patients with coronary artery disease who are acceptable candidates for coronary artery bypass graft surgery and who meet one of the following selection criteria: ➢ Single vessel atherosclerotic coronary artery disease with a stenosis that can be passed with a guidewire, ➢ Multiple vessel coronary artery disease that in the physician's judgment does, not pose undue risk to the patient, ➢ Certain patients who have had prior percutaneous transluminal coronary, angioplasty (PTCA), and who have a restenosis of the native vessel; or, Native vessel atherosclerotic coronary artery disease that is less than 25 mm in length.	**Contraindications:** Occlusions through which a guidewire will not pass. Last remaining vessel with compromised left ventricular function. Saphenous vein grafts. Angiographic evidence of thrombus prior to treatment with the RotablatorSystem. Such patients may be treated with thrombolytics (e.g., Urokinase). When the thrombus has been resolved for two to four weeks, the lesion may be treated with the Rotablator System. Angiographic evidence of significant dissection at the treatment site. The patient may be treated conservatively for approximately four weeks to permit the dissection to heal before treating the lesion with the Rotablator System. **Adverse Event:** Angina or unstable angina, Arrhythmias, Bailout stenting, Cardiac perforation, Cardiac tamponade, Conduction block, Coronary artery spasm, Death, Drug reaction, allergic reaction to contrast media, Embolism (coronary, cerebral, peripheral), Hemorrhage or hematoma, Infection, local infection, systemic infection, Myocardial ischemia, Myocardial infarction (Q-wave and non Q-Wave), Pericardial effusion, Pulmonary edema/cardiogenic shock, Slow flow, no flow, abrupt vessel closure, Stroke, Vascular thrombus, Vessel trauma (dissection, perforation, rupture or injury)	USA EC Certificate	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj

Sl No	c¯ZKvi‡Ki bvg	ewYRK bvg	‡gvW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve KwgwUi mfvi wm×vš	mfvi wm×vš
109.	**Manufacturer:** Boston Scientific Corporation, 300 Boston Scientific Way, Marlborough, MA 01752, USA **Local Agent:** M/s. Medi Graphic Trading Ltd., 14 Purana Paltan, Dhaka, Bangladesh	RotaLink™ Advancer	Catheter advancing device	D	Percutaneous rotational coronary angioplasty with the Rotablator Rotational Atherectomy System, as a sole therapy or with adjunctive balloon angioplasty, is indicated in patients with coronary artery disease who are acceptable candidates for coronary artery bypass graft surgery and who meet one of the following selection criteria: ➢ Single vessel atherosclerotic coronary artery disease with a stenosis that can be passed with a guidewire, ➢ Multiple vessel coronary artery disease that in the physician's judgment does, not pose undue risk to the patient, ➢ Certain patients who have had prior percutaneous transluminal coronary, angioplasty (PTCA), and who have a restenosis of the native vessel; or, ➢ Native vessel atherosclerotic coronary artery disease that is less than 25 mm in length.	**Contraindications:** Occlusions through which a guidewire will not pass. Last remaining vessel with compromised left ventricular function. Saphenous vein grafts. Angiographic evidence of thrombus prior to treatment with the Rotablator, System. Such patients may be treated with thrombolytics (e.g., Urokinase). When the thrombus has been resolved for two to four weeks, the lesion may be treated with the Rotablator System. . Angiographic evidence of significant dissection at the treatment site. The patient may be treated conservatively for approximately four weeks to permit the dissection to heal before treating the lesion with the Rotablator System. **Adverse Event:** Angina or unstable angina, Arrhythmias, Bailout stenting, Cardiac perforation, Cardiac tamponade, Conduction block, Coronary artery spasm, Death, Drug reaction, allergic reaction to contrast media, Embolism (coronary, cerebral, peripheral), Hemorrhage or hematoma, Infection, local infection, systemic infection, Myocardial ischemia, Myocardial infarction (Q-wave and non Q-Wave), Pericardial effusion, Pulmonary edema/cardiogenic shock, Slow flow, no flow, abrupt vessel closure, Stroke, Vascular thrombus, Vessel trauma (dissection, perforation, rupture or injury)	USA EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

Sl No	c¯ZKvi‡Ki bvg	ewYwR¨K bvg	‡gwW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve KwgwUi mfvi wm×vš	mfvi wm×vš
110.	**Manufacturer:** Boston Scientific Corporation, 300 Boston Scientific Way, Marlborough, MA 01752, USA **Local Agent:** M/s. Medi Graphic Trading Ltd., 14 Purana Paltan, Dhaka, Bangladesh	RotaLink™ Plus	Pre-Connected Exchangeable Burr Catheter and burr advancing device	D	Percutaneous rotational coronary angioplasty with the Rotablator Rotational Atherectomy System, as a sole therapy or with adjunctive balloon angioplasty, is indicated in patients with coronary artery disease who are acceptable candidates for coronary artery bypass graft surgery and who meet one of the following selection criteria: ➢ Single vessel atherosclerotic coronary artery disease with a stenosis that can be passed with a guidewire, ➢ Multiple vessel coronary artery disease that in the physician's judgment does, not pose undue risk to the patient, ➢ Certain patients who have had prior percutaneous transluminal coronary, angioplasty (PTCA), and who have a restenosis of the native vessel; or, ➢ Native vessel atherosclerotic coronary artery disease that is less than 25 mm in length.	**Contraindications:** Occlusions through which a guidewire will not pass, Last remaining vessel with compromised left ventricular function, Saphenous vein grafts. Angiographic evidence of thrombus prior to treatment with the Rotablator System. Such patients may be treated with thrombolytics (e.g., Urokinase). When the thrombus has been resolved for two to four weeks, the lesion may be treated with the Rotablator System. Angiographic evidence of significant dissection at the treatment site. The patient may be treated conservatively for approximately four weeks to permit the dissection to heal before treating the lesion with the Rotablator System. **Adverse Event:** Angina or unstable angina, Arrhythmias, Bailout stenting, Cardiac perforation, Cardiac tamponade, Conduction block, Coronary artery spasm, Death, Drug reaction, allergic reaction to contrast media, Embolism (coronary, cerebral, peripheral), Hemorrhage or hematoma, Infection, local infection, systemic infection, Myocardial ischemia, Myocardial infarction (Q-wave and non Q-Wave), Pericardial effusion, Pulmonary edema/cardiogenic shock, Slow flow, no flow, abrupt vessel closure, Stroke, Vascular thrombus, Vessel trauma (dissection, perforation, rupture or injury)	USA EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

Sl No	c¯ZKv i‡Ki bvg	ewYwR¨K bvg	‡gwW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve KwgwUi mfvi wm×vš	mfvi wm×vš
111.	**Manufacturer:** Boston Scientific Corporation, 300 Boston Scientific Way, Marlborough, MA 01752, USA **Local Agent:** M/s. Medi Graphic Trading Ltd., 14 Purana Paltan, Dhaka, Bangladesh	Rotawire and wireClip Torquer	Rotablator Rotational Atherectomy System Guidewires	D	These guidewires are intended for use with the Rotablator Rotational Atherectomy System.	**Contraindications:** Carefully read this document and refer to the Rotablator Console "Operations Manual" and Rotalink Exchangeable Burr Catheter "Instructions for Use", observing all Contraindications, Restrictions, Warnings, and Precautions for specific information on the use of these components. **Adverse Event:** Angina or unstable angina, Arrhythmias, Bailout stenting, Cardiac perforation, Cardiac tamponade, Conduction block, Coronary artery spasm, Death, Drug reaction, allergic reaction to contrast media, Embolism (coronary, cerebral, peripheral), Hemorrhage or hematoma, Infection, local infection, systemic infection, Myocardial ischemia, Myocardial infarction (Q-wave and non Q-Wave), Pericardial effusion, Pulmonary edema/cardiogenic shock, Slow flow, no flow, abrupt vessel closure, Stroke, Vascular thrombus, Vessel trauma (dissection, perforation, rupture or injury)	USA EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
112.	**Manufacturer:** MicroVention, Inc. 1311 Valencia Ave Tustin, CA 92780 USA **Local Agent:** M/S, UniTrade Corporation, 2/5, Humayun Road (3rdFl), Block-B Mohammadpur, Dhaka-1207	**Scepter C & XC** Occlusion Balloon Catheter	Occlusion Balloon Catheter	**D**	The Scepter C and Scepter XC Occlusion Balloon Catheters are intended: For use in the peripheral and neuro vasculature where temporary occlusion is desired. The balloon catheters provide temporary vascular occlusion which is useful in selectively stopping or controlling blood flow. The balloon catheters also offer balloon assisted embolization of intracranial aneurysms. For use in the peripheral vasculature for the infusion of diagnostic agents, such as contrast media, and therapeutic agents such as embolization materials. For neurovascular use for the infusion of diagnostic agents such as contrast media, and therapeutic agents, such as embolization materials, that have been approved or cleared for use in the neurovasculature and are compatible with the inner diameter of the Scepter C/XC Balloon Catheter.	**Contraindication:** ➢ Not intended for embolectomy or angioplasty procedures ➢ Not intended for use in coronary vessels ➢ Not intended for pediatric or neonatal use **Potential complication**: Potential complications include, but are not limited to: vessel or aneurysm perforation, vasospasm, hematoma at the site of entry, embolism, ischemia, ntracerebral/ intracranial hemorrhage, pseudo aneurysm, seizure, stroke, infection, vessel dissection, thrombus formation, and death.	USA EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

Sl No	c¯ZKvi‡Ki bvg	ewYwR¨K bvg	‡gwW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve KwgwUi mfvi wm×vš	mfvi wm×vš
113.	**Manufacturer:** **St. Jude Medical** Cardiac Rhythm Management Division 15900 Valley View Ct Sylmar, CA USA 91342 **Local Agent:** **M/s. The Spondon Ltd**. BSEC Bhaban, Level-9, 102 Kazi Nazrul Islam Avenue, Kawran Bazar, Dhaka-1215	SJM Implantable Cardiac Pacing Lead	Implantable Cardiac Pacing Lead	D (P-46)	Leads are designed for use in compbination with a compatible pulse generator to provide permanent pacing and sensing in either the atrium or ventricle.	**Contraindication:** In patient who are expected to be hypersensitive to a single dose of 1.0 milligram of dexamethasone sodium phosphate. In the presence of tricuspid atresia and in patients with mechanical tricuspid valves **Side-effect:** Implanted cardiac leads are subjected to a hostile environment within the body due to constant, complex flexural and tensional forces, interactions with leads and/or the pulse generator, or other forces associated with cardiac contractions and patient physical activity, posture, and anatomical influence. Cardiac leads functional lifetimes can be affected by these and other factors.	USA EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i|	Abyjgv`b Kiv nj|
114.	**Manufacturer:** Boston Scientific Corporation, 300 Boston Scientific Way, Marlborough, MA 01752, USA **Local Agent:** M/s. Medi Graphic Trading Ltd., 14 Purana Paltan, Dhaka, Bangladesh	Soloist Single Needle Electrode	Radiofrequency Ablation Needle Electrode	D	It is intended to be used in conjunction with the RF 3000 Generator for the thermal coagulation necrosis of soft tissues, including partial or complete ablation of nonresectable liver lesions. These procedures should only be performed by physicians and staff familiar with the equipment and techniques involved.	**Contraindications:** Before using, inspect the package for any breach to the sterile barrier and inspect product for any damage. If package is broken or product is damaged DO NOT USE. Immediately return package and product to Boston Scientific. **Adverse Events:** Abscess, ARDS (Acute Respiratory Distress Syndrome), Arrhythmia, Ascites, Biloma, Burn, Death, Delayed hemorrhage into ablated tissue, Diarrhea, Electric Shock, Fistula, including biliary fistula, Hematoma, Hemorrhage, Infection, Liver Failure, Liver Insufficiency, Pain, Perforation, Peritonitis, Persistent Fever > 39° C, Pleural Effusion, Renal Failure, Tumor Recurrence, Tumor Seeding	USA EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i|	Abyjgv`b Kiv nj|

Sl No	c¯ZKvi‡Ki bvg	ewYwR¨K bvg	‡gvW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve Kwgw Ui mfvi wm×vš	mfvi wm×vš
115.	**Manufacturer:** Boston Scientific Limitwd Ballybrit Business park. Galway, Ireland **Local Agent:** M/s. Medi Graphic Trading Ltd., 14 Purana Paltan, Dhaka, Bangladesh	Stingray Extension Wire	Extension wire	**D**	The attachment of the Stingray Extension Wire to the extendable guidewire creates a guidewire that can be used to exchange a catheter without removing the guidewire from the vessel. When the exchange is complete, the guidewire extension can be detached and the original guidewire can again be used in a conventional manner.	**Contraindications:** None known. **Adverse Events:** None known.	Ireland EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |
116.	**Manufacturer:** Boston Scientific Limitwd Ballybrit Business park. Galway, Ireland **Local Agent:** M/s. Medi Graphic Trading Ltd., 14 Purana Paltan, Dhaka, Bangladesh	Stingray Guidewire with Hydrophilic coating	Guidewire	D	The Stingray Guidewires are intended to facilitate placement of balloon dilatation catheters or other intravascular devices during percutaneous transluminal coronary angioplasty (PTCA) and percutaneous transluminal angioplasty (PTA). Stingray Guidewires are not to be used in cerebral blood vessels. When used as part of the system consisting of the CrossBoss™ Catheter, Stingray Catheter, and Stingray Guidewire, the Stingray Guidewire is indicated for use to facilitate the intraluminal placement of conventional guidewires beyond stenotic coronary lesions (including chronic total occlusions [CTOs]) prior to PTCA or stent Intervention.	**Contraindications:** None known. **Adverse Events:** None known.	Ireland EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |
117.	**Manufacturer:** Boston Scientific Limitwd Ballybrit Business park. Galway, Ireland **Local Agent:** M/s. Medi Graphic Trading Ltd., 14 Purana Paltan, Dhaka, Bangladesh	Stingray LP Catheter	Vascular Guide Catheter	D	The Stingray LP Catheter is indicated for directing, steering, controlling, and supporting a guidewire in order to access discrete regions of the coronary and peripheral vasculature. When used as part of the System consisting of the CrossBoss™ Catheter, Stingray LP Catheter and Stingray Guidewire, the Stingray LP Catheter is indicated for use to facilitate the intraluminal placement of conventional guidewires beyond stenotic coronary lesions (including chronic total occlusions [CTOs]) prior to PTCA or stent intervention.	**Contraindications:** Not intended for use in the cerebral vasculature. Unprotected left main coronary artery. Coronary artery spasm in the absence of a significant stenosis. **Adverse Events:** Acute myocardial infarction, Hemorrhage or hematoma, Vessel trauma that may require further intervention or surgical repair, Artery spasm. Embolism, Stroke/neurological deficit, Drug reactions, allergic reaction to contrast media, Infection, Cardiac tamponade, Recurrence of angina,	Ireland EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i |	Abyjgv`b Kiv nj |

Sl No	c¯ZKvi‡Ki bvg	ewYRK bvg	‡gvW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve Kwgkb mfvi wm×vš	mfvi wm×vš
118.	**Manufacturer:** Ethicon, LLC, USA **Distributor:** Johnson & Johnson PVt. Ltd., India **Local Agent:** Modern Surgical (pvt) Ltd., 34/1, (2nd Floor) Mitford Road, Dhaka-1100	SURGICEL Absorbable Hemostat	Hemostat	**D**	SURGICEL Haemostat is used adjunctively in surgical procedures to assist in the control of capillary, venous and small arterial haemorrhage when ligation or other conventional methods of control are impractical or ineffective. Surgicel Haemostat can be used in many aeras of surgery e.g. cardiovascular surgery, haemorrhaidectomy, implantation of vascular prostheses, biopsies, lung operations, surgery to the face and jaw, gasrik resection, operations to the throat or nose, liver and gall bladder operations, thoracic and abdominal sympathectomies, neurosurgery, especially cerebral operations, skin transplantations, treatment of superficial injuries. Surgicel Haemostat is indicated also for adjunctive use in dental application to assist in the control of bleeding in exodontia and oral surgery. It may also be used to help achieve haemostasis after single or multiple tooth extractions, gingival haemorrhage, impactions, biopsies and other procedures in the oral cavity.	**Contraindications:** • Although packing or wadding sometimes is medically necessary, Surgicel Haemostat should not be used in this manner, unless it is to be removed after haemostasis is achieved. • Surgicel Haemostat should not be used for implantation in bone defects, such as fractures, since there is possibility of interference with callus formulation and a theoretical chance of cyst formation. • When Surgicel Haemostat is used to help haemostatsis in, around, or in proximity to foramina in bony confine, the spinal cord, or the optic nerve and chiasm, it must always be removed after haemostasis is achieved since it will swell and could exert unwanted pressure. • Surgicel Haemostat should not be used to control haemorrhage from large arteries. • Surgicel Haemostat should not be used on non-haemorrhagic serous oozing surfaces, since body fluids other than whole blood, such as serum, do not react with Surgicel Haemostat to produce satisfactory haemostatic effect. • Surgicel Haemostat should not be used as an adhesion prevention product. **Warnings:** Surgicel Haemostat is supplied sterile and as the material is not compatible with autoclaving or ethylene oxide sterilization Surgicel Haemostat should not be re-sterilized. Surgicel Haemostat is not intended as a substitute for careful surgery and proper use of suture and ligatures. Closing Surgicel Haemostat in a contaminated wound without drainage may lead to complications and should be avoided.	USA EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

Sl No	c¯ZKv i‡Ki bvg	ewYwR¨K bvg	‡gwW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve KwgwUi mfvi wm×vš	mfvi wm×vš
119.	**Manufacturer:** Ferrosan Medical Devices A/S, Denmark **Distributor:** Johnson & Johnson PVt. Ltd., India **Local Agent:** Modern Surgical (pvt) Ltd., 34/1, (2nd Floor) Mitford Road, Dhaka-1100	**SURGIFLO** Haemostatics Matrix Kit thrombin	Haemostatics Matrix Kit thrombin	**D**	SURGIFLO™ is indicated for surgical procedures (except ophthalmic) for haemostasis, when control of capillary, venous and arteriolar bleeding by pressure, ligature, and other conventional procedures is ineffective or impractical.	**Contraindications:** • Do not inject or compress SURGIFLO™ into blood vessels. Do not use SURGIFLO™ in intravascular compartments because of the risk of thromboembolism, disseminated intravascular coagulation, and increased risk of anaphylactic reaction. • Do not use SURGIFLO™ in patients with known allergies to porcine gelatin. • Do not use SURGIFLO™ in closure of skin incisions because it may interfere with the healing of skin edges. This interference is due to mechanical interposition of gelatin and is not secondary to intrinsic interference with wound healing. **Side Effects:** Not Known	Denmark EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
120.	**Manufacturer:** MicroVention, Inc. 1311 Valencia Ave Tustin, CA 92780 USA **Local Agent:** M/S, UniTrade Corporation 2/5, Humayun Road (3rdFl), Block-B Mohammadpur, Dhaka-1207	Traxcess Guidewires & Docking Wire Cerebral Aneurysm	Guidewires and Docking Wire for	**D**	Traxcess Guidewire is intended for general intravascular use, including the neuro and peripheral vasculature. The guidewire can be steered to facilitate the selective placement of diagnostic or therapeutic catheters. This device is not intended for use in coronary arteries.	**Contradiction:** There are no known contraindications. **Side Effect**: None	USA EC Certificate	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj

Sl No	c¯ZKvi‡Ki bvg	ewYwR¨K bvg	‡gvW‡Kj wWfvB‡mi bvg	Kvm	wb‡`kbv	Contraindication & Side-effect	FSC/CPP	‡UKwbK¨vj mve KwgwUi mfvi wm×vš	mfvi wm×vš
121.	**Manufacturer:** Toshiki International Singapore Pte Ltd., Singapore **Local Agent:** Zuellig Pharma Bangladesh ltd., 191/A Haque Tower, Tejgaon Gulshan Link Road Dhaka	Dermatix ultra Gel	Silicone gel	C	It is a topical silicone gel that functions as a transparent physical barrier that dries quickly helps maintain the skins moisturize balance, while aiding the management of scarred skin suface resulting from surgery, burns and other injuries dertix ultra gel is used for the management and prevention caloids and hypertropic scares (such as those resulting from general surgical procedures, trauma, wounds and burns). It should be used after the wound has healed and when the skin surface is intact.	**Contradiction:** There are no known contraindications. **Side Effect**: None	Singapore	‡iwR‡ókb MvBWjvBb Abjyqx c‡qvRbxq WKzgv›U `wLj Kivi Rb¨ ejv †h‡Z cv‡i|	‡iwR‡ókb MvBWjvBb Abjyqx c‡qvRbxq WKzgv›U `wLj Kivi Rb¨ ejv nj|
122.	**Manufacturer:** Eucare Pharmaceuticals Pvt Ltd., INDIA **Local Agent:** Kulic Limited 9/7, Eastern Plaza (8th Floor), 70, BirUttom C.R Datta Road, Hatirpul, Dhaka	Lysil Gel	Silicone gel	C	Derived from high purity polysiloxane material and extremely useful in treating new and old scars and hypertrophic scarring.	**Contradiction:** There are no known contraindications. **Side Effect**: None	India	‡iwR‡ókb MvBWjvBb Abjyqx c‡qvRbxq WKzgv›U `wLj Kivi Rb¨ ejv †h‡Z cv‡i|	‡iwR‡ókb MvBWjvBb Abjyqx c‡qvRbxq WKzgv›U `wLj Kivi Rb¨ ejv nj|

Annex-F: Proposed Product for Import (IVD Reagent)

Sl No	c¯ZKvi‡Ki bvg	ewYR¨K bvg	AvBwfwW wi‡q‡R‡Ui bvg	Kvm	Intended use	FSC/CPP	†UKwbK¨vj mve-KwgwU mfvi wm×vš	mfvi wm×vš
123.	**Manufacturer:** Haemonetics Corporation, 400, Wood Road, Braintree, Massachusetts, MA 02184, USA **Local Agent :** MBiologix, 205 Shahid Syed Nazrul Islam Swarani, Dhaka-1000.	ACD-A Anticogulant Solution 500ml REF:426C Acid Citrate Dextrose- Adenine	Sterile, Non-pyrogenic Solution for the Anticoagulation of Whole Blood for Use in Automated Apheresis Procedures.	C	Sterile, Non-pyrogenic Solution for the Anticoagulation of Whole Blood for Use in Automated Aphaeresis Procedures.	Switzerland	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
124.	**Manufacturer:** Lifescan, Europe, Switzerland **Local Agent :** MF Consumers Limited, Anchor Tower, 108 Bir uttam CR Datta Road, Dhaka	OneTouch Ultra® Control Solution	Control	B (P-38)	OneTouch Ultra® Control Solution contains a known amount of glucose and is used to check that the meter and the test strips are working properly. Do a control solution test: ➢ To practice the test process without having to use blood ➢ Once a week to check your meter ➢ Whenever you open a new vial of test strips ➢ If you suspect the meter or test strips are not working properly ➢ If you are repeatedly getting unexpected blood sugar results ➢ If you drop or damage your meter	Switzerland	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
125.	**Manufacturer:** Meril Diagnostic Pvt. Ltd., India **Local Agent :** Meril Bangladesh Pvt. Ltd.,West Panthopath, Dhaka	Merisera Combo (Anti-A, Anti-B, Anti-D(IgM) Blood Grouping Reagent	Combo (Anti-A, Anti-B, Anti-D(IgM) Blood Grouping Reagent	D	It is intended for in-vitro determination of ABo group and Rh typing of Human red blood cell antigen.	India	‡iwR‡ókb MvBWjvBb Abyjvqx c‡qvRbxq WKzjg›U `wLj Kivi Rb¨ ejv †h‡Z cv‡i	‡iwR‡ókb MvBWjvBb Abyjvqx c‡qvRbxq WKzjg›U `wLj Kivi Rb¨ ejv nj

Sl No	c¯ZKvi‡Ki bvg	ewYR¨K bvg	AvBwfW wi‡q‡R‡Ui bvg	Kvm	Intended use	FSC/CPP	†UKwbK¨vj mve-KwgwU mfvi wm×vš	mfvi wm×vš
126.	**Manufacturer:** Beacon Diagnostic Pvt. Ltd., India **Local Agent :** Medi Need International, B-5/7, Ganda, Savar, Dhaka	Tuberculin purified protein derivative	Tuberculin purified protein derivative	C	Tuberculin purified protein derivative (PPD) is a diagnostic agent. It works by causing a mild, delayed allergic reaction in patients infected with TB or who have had a past infection, which allows for detection of TB.	India	‡iwR‡ókb MvBWjvBb Abjyvqx c‡qvRbxq WKzgvU `wLj Kivi Rb¨ ejv †h‡Z cv‡i	‡iwR‡ókb MvBWjvBb Abjyvqx c‡qvRbxq WKzgvU `wLj Kivi Rb¨ ejv nj
127.	**Manufacturer:** Beacon Diagnostic Pvt. Ltd., India **Local Agent :** Medi Need International, B-5/7, Ganda, Savar, Dhaka	Widal Antigen Slide Test	Staine Salmonella Antigen test	C	It is used for detection of Specific Antibodies produced in response to the stimulation by specific antigen of Salmonella (group)	India	‡iwR‡ókb MvBWjvBb Abjyvqx c‡qvRbxq WKzgvU `wLj Kivi Rb¨ ejv †h‡Z cv‡i	‡iwR‡ókb MvBWjvBb Abjyvqx c‡qvRbxq WKzgvU `wLj Kivi Rb¨ ejv nj
128.	**Manufacturer:** Becton Dickinson and company, USA **Local Agent :** Becton Dickinson India Pvt. Ltd.80, Kakrail Dhaka	BD FACScount Reagent Kit	CD4, CD8, and CD3 counting reagent	D	The BD FACSCount™ reagent kit is intended for in vitro diagnostic use in enumerating the absolute counts of CD4, CD8, and CD3 T-lymphocytes in unlysed whole blood, using the BD FACSCount™ Instrument.	USFDA	Abjyvb Kiv †h‡Z cv‡i	Abjyvb Kiv nj
129.	**Manufacturer:** Becton Dickinson and company, USA **Local Agent :** Becton Dickinson India Pvt. Ltd.80, Kakrail Dhaka	BD Facscount Control Kit	FACSCount™ instrument Control	B	The BD FACSCount™ control kit is intended for in vitro diagnostic use in setting up the BD FACSCount™ instrument and for checking linearity.	USFDA	Abjyvb Kiv †h‡Z cv‡i	Abjyvb Kiv nj
130.	**Manufacturer:** Becton Dickinson and company, USA **Local Agent :** Becton Dickinson India Pvt. Ltd.80, Kakrail Dhaka	BD FACS flow sheath Fluid	HIV Detection Reagent	B	Optimized sheath fluid for use on flow cytometry instrument	USFDA	Abjyvb Kiv †h‡Z cv‡i	Abjyvb Kiv nj

Annex-: G Proposed Products for Locally Manufacture (Herbal)

bs	c¯ZKvi‡Ki bvg	JI‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	‡UKwbK¨vj mve KwgwUi wm×vš	mfvi wm×vš
1.	Acme Laboratories Ltd. (Herbal & Nutraceuticals Division)	Organic Bitter Melon Fruit extract + Organic Bitter Melon Stem powder (*Momordica charantia Linn.*) Capsule Organic Bitter Melon Fruit extract (*Momordica charantia Linn.*) 250 mg + Organic Bitter Melon Stem powder (*Momordica charantia Linn.*) 410 mg Capsule	It is used as hypoglycemic (blood sugar lowering) agent	**Contraindication:** It is contraindicated during pregnancy, persons with hypoglycemia & in women who are breast feeding **Side effects**: There is a relatively low toxicity of all parts of the bitter melon	New Molecule	PDR for Herbal Medicines, 3rd Edition, P-88, 89	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj
2.	Acme Laboratories Ltd. (Herbal & Nutraceuticals Division)	Svarjiksara (Sodium Bicarbonate) + Nimbukamlam (Citric acid) (coated) Effervescent Powder Svarjiksara (Sodium Bicarbonate) 2.200 g + Nimbukamlam (Citric acid) (coated) 2.070 g/5 g Effervescent Powder	The symptomatic relief of indigestion, flatulence & nausea	**Contraindication:** It is contraindicated into patients on salt restricted diets **Side effects**: No known side effects	New Molecule	Eno Of GlaxoSmith Klines (GSK) product	No Approved Reference	bvgÄyi 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	‡UKwbK¨vj mve KwgwUi wm×vš	mfvi wm×vš
3.	Acme Laboratories Ltd. (Herbal & Nutraceuticals Division)	**Camphor Oil (Ext.) + Blue Gum Tree Oil + Mint (Ext.) + Turpentine Oil + Nutmeg Oil Cold Relief Balm** Camphor Oil (Ext.) 2.5 g + Blue Gum Tree Oil 1.25 g + Mint (Ext.) 1.25 g + Turpentine Oil 1.25 g + Nutmeg Oil 1.25 g/100 g Cold Relief Balm	It is a soothing balm that relieves nasal and chest congestion. Its counter-irritant effects relieve headaches and body aches associated with the common cold	**Contraindication:** There is no known contraindication **Side effects**: No known side effects	New Molecule	Cold Balm of Himalaya Drug Co. Ltd., India The German Commission E Monograph	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
4.	Acme Laboratories Ltd. (Herbal & Nutraceuticals Division)	**Nim (Ext.) Ph. Grade + Turmeric (Rz.) Face Wash** Nim (Ext.) Ph. Grade 50 mg + Turmeric (Rz.) Ph. Grade 50 mg/g Face Wash	This medicated face wash prevents pimple, acne & helps clean & clear skin naturally	**Contraindication:** There is no known contraindication **Side effects**: No known side effects	New Molecule	Himalaya Drug Co. Ltd., India PDR for Herbal Medicine 2nd edition. The German Commission E Monograph Mosbys drugs consultant	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
5.	Acme Laboratories Ltd. (Herbal & Nutraceuticals Division)	Tumeric (*Curcuma longa*) + Fenugreek (*Trigonellafoenum-graecum*) + Sal Tree (*Shorea robusta*) + Honey (*Mel despumatum*) Cream Tumeric (*Curcuma longa*) Ph. Grade 1.25 g + Fenugreek (*Trigonellafoenum- graecum*) Ph. Grade 1.25 g + Sal Tree (*Shorea robusta*) Ph. Grade 1.25 g + Honey (*Mel despumatum*) 10 g/100 g Cream	The cream is useful in dry & cracked heels/soles & rough feet	**Contraindication:** There is no known contraindication **Side effects**: There is a relatively low toxicity of all parts of the bitter melon	New Molecule	Foot Care Cream of Himalaya Drug Co. Ltd., India	No Approved Reference	bvgÄyi 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	‡UKwbK¨vj mve KwgwUi wm×vš	mfvi wm×vš
6.	Bexer Herbal & Nutraceuticals Matidali, 2nd Bypass Road, Manikchock, Bogra.	**Fructus Agni Casti** Fructus Agni Casti 162 mg tablet Vitex agnus-castus	Fructus Agni Casti consists of the dried, ripe fruits of Vitex agnus-castus L. (Lamiaceae) (1, 2). Used as an anaphrodisiac, calefacient, contraceptive, emmenagogue, seda-tive and as a tonic. Effect on prolactin secretion, Abnormal menstrual cycles and infertility, Acne treatment, Cyclic breast pain (mastalgia), Premenstrual syndrome, Effects on lactation.	**Contra-indication:** Fructus Agni Casti should not be used during pregnancy. **Side effect:** The major reactions reported included acne, changes to the menstrual cycle, dizziness, gastrointestinal distress, increased menstrual flow, nausea, skin reactions, urticaria and weight gain .		1. WHO monographs on selected medicinal plants. Volume 4	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
7.	Bexer Herbal & Nutraceuticals Matidali, 2nd Bypass Road, Manikchock, Bogra.	**Ginkgo biloba** Ginkgo biloba 70 mg capsule.	Cerebral insufficiency: memory deficit, depression, attention and memory loss that occur with Alzheimer's disease and multi-infarct dementia. Vertigo and tinnitus (ringing in the ear) of vascular and involutional origin Peripheral vascular disease: improvement of pain-free walking distance in Peripheral Arterial Occlusive Disease in Stage II according to Fontaine (intermittent claudication) in a regimen of physical therapeutic measures, in particular walking exercise. **OTHER POTENTIAL USES :** Protective action in hypoxia Acute cochlear deafness Sexual dysfunction associated with SSRI use.	**Contra-indication:** Some MEDICINES MAY INTERACT with ginkgo biloba. Tell your health care provider if you are taking any other medicines, especially any of the following: Nonsteroidal anti-inflammatory drugs (NSAIDs; eg, aspirin, ibuprofen) or warfarin because the risk of serious bleeding may be increased **Side effect:** No side effects following proper administration of designated therapeutic dosages.		1. PDR for Herbal Medicine 2nd edition. Page 342 - 346 2. Peng, F., H. Guo, M. Hao, J. Guo, Y. Yang & P. Tan (2012) Am. J. Plant Sci. 102-9. 3. Stan, I. (2009) Bull. UASVM Horticult. 66: 620- 4. Mori, M., K. Suzuki & R. Kohzaki (2000) J. Japan. Soc. Food Soc. Technol. 47: 448-51. 5. Barlow, P.W. & E.U. Kurczynska (2007) J. Plant Res. 120: 269-80.	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b 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	‡UKwbK¨vj mve KwgwUi wm×vš	mfvi wm×vš
8.	Bexer Herbal & Nutraceuticals Matidali, 2nd Bypass Road, Manikchock, Bogra.	Salix alba bark extract 500mg tablet	Used in back pain, join pain	**Contra-indication:** Cortex Salicis is contraindicated in cases of hypersensiticity or allergy to the plant material or to salicylates (e.g. asthma, bronchial spasm, rhinitis or urticaria) **Side effect:** Allergicreaction such as pruritus, urticaria and gastrointestinal symptoms may occur. One case of an allergic reaction in a 32- year-old atopic patient who showed a severe anaphylactic reaction after the ingestion of a pollen compound containing 9the crude drug has been reported.		Who monographs on Selected medicinal plants Volume 4	Abyjgv`b Kiv †h‡Z cv‡i|	Abyjgv`b Kiv nj|
9.	Bexer Herbal & Nutraceuticals Matidali, 2nd Bypass Road, Manikchock, Bogra.	**Glucosamine and Chondroitin Sulfate** Glucosamine 250 mg and Chondroitin Sulfate 200 mg Tablet	Osteoarthritis (OA), Rheumatoid arthritis (RA), Degenerative change in bone and joint. Prophylaxis of muscle and tendon injury.	**Contra-indication:** It is safe in recommended dose but caution should be taken in Diabetes, Glucose-6-phosphate dehydrogenase deficiency (G6PD deficiency), Iron-related disorders such as hemochromatosis, thalassemia or anemia and sickle cell disease. **Side effect:** A traditional and widely consumed food, the rosehip showed considerable safety. In very rare case acid regurgitation, diarrhea and constipation may occur.		WorksafeBC Evidence-Based Practice Group Dr. Craig W.Martin, Senior Medical Advisor. O'Donnell S, Lagacé C, McRae L, Bancej C. Life with arthritis in Canada: a personal and public health challenge Chronic Diseases and Injuries in Canada. 2011;Vol 31(3):135-6.	No Approved Reference	bvgÄyi 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	‡UKwbK¨vj mve KwgwUi wm×vš	mfvi wm×vš
10.	Bexer Herbal & Nutraceuticals Matidali, 2nd Bypass Road, Manikchock, Bogra.	**Maca** Maca 500 mg capsule *Lepidium meyenii*	sexual dysfunction in women antidepressants Male infertility sperm count blood pressure depression anxiety anemia Leukemia Sexual desire Depression Symptoms of menopause Osteoporosis	**Contra-indication:** Patients with thyroid conditions should avoid maca because glucosinolates taken in excess and combined with a low-iodine diet can cause goiter. **Side effect:** Maca is **LIKELY SAFE** for most people when taken in amounts found in foods.		Maca. Review of Natural Products. factsandcomparisons4.0 [online]. February 2008. Available from Wolters Kluwer Health, Inc. Accessed January 17, 2008. 2. TGA, Medsafe. Interim Joint Expert Advisory Committee on Complementary Medicine. Meeting 3 (September 14th, 2006) 2006. http://www.medsafe.govt.nz/Profs/class/classintro.asp (Accessed September 24th, 2009). 3. MHRA. Safety of herbal medicines. 2005. http://www.mhra.gov.uk/home/groups/is-pol/documents/websiteresources/con009277.pdf (Accessed September 24th,	No Approved Reference	bvgÄyi 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	‡UKwbK¨vj mve KwgwUi wm×vš	mfvi wm×vš
11.	Bexer Herbal & Nutraceuticals Matidali, 2nd Bypass Road, Manikchock, Bogra.	**Muira Puama** Muira Puama 500mg capsule *Ptychopetalum olacoides*	• sexual intercourse in women with a low sex drive. • Sexual disorders. Stomach upset. • Menstrual disorders. • Sore joints. • Loss of appetite. • Other conditions.	**Contra-indication:** Contraindications have not yet been identified. **Side effect:** It is not known if muira puama is safe or what the possible side effects might be.		1. PDR for Herbal Medicines . 2nd ed. Montvale, NJ: Thomson Medical Economics; 2000: 531-532. 2. Schultes R, et al. The Healing Forest: Medicinal and Toxic Plants of the Northwest Amazon. Portland, OR: Dioscorides Press,1990, p. 343. 3. Bucek E, et al. Volatile constituents of Ptychopetalum olacoides root oil. Planta Med 1987;53:231.	Abyjgv`b Kiv †h‡Z cv‡i|	Abyjgv`b Kiv nj|
12.	Bexer Herbal & Nutraceuticals Matidali, 2nd Bypass Road, Manikchock, Bogra.	**Folium Cynarae** Folium Cynarae 500 mg tablet *Cynara scolymus*	Folium Cynarae consists of the dried basal leaves of Cynara cardunculus. Irritable bowel syndrome. Oral treatment of anaemia, diabetes, fever, gout, rheumatism and urinary stones	**Contra-indication:** Hypersensitivity or allergies to artichokes and other plants from the Compositae/ Asteraceae, and obstruction of the bile ducts. **Side effect:** Gastrointestinal complaints included mild diarrhoea, accompanied by ab- dominal cramps, upper abdominal pain, nausea and heartburn. Allergic reactions may occur in sensitized patients		WHO monographs on selected medicinal plants. Volume 4	Abyjgv`b Kiv †h‡Z cv‡i|	Abyjgv`b 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	‡UKwbK¨vj mve KwgwUi wm×vš	mfvi wm×vš
13.	Bexer Herbal & Nutraceuticals Matidali, 2nd Bypass Road, Manikchock, Bogra.	**Tang-kuei** Tang-kuei 500 mg capsule *Angelica sinensis*	Premature ejaculation cinnamon Heart disease menstruation High blood pressure pulmonary hypertension Premenstrual syndrome Joint aches and pains, Ulcers, Anemia, Constipation, Skin discoloration and psoriasis, Allergies, Other conditions. More evidence is needed to rate the effectiveness of Tang-kuei for these uses.	**Contra-indication:** We have no information for Tang-kuei contraindications. **Side effect:** Not to be used during pregnancy due to the stimulating effect of Tang-kuei. If you are using any anti-coagulant or blood thinning medication such as warfarin or aspirin, consult your health care professional prior to using this product.		1. Jouranl of Ethnopharmacology. 2. Hirata, Janie D.; Swiersz, Lillian M.; Zell, Bonnie; Small, Rebecca; Ettinger, Bruce. Does Dong Quai Have Estrogenic Effects in Postmenopausal Women? A Double-Blind, Placebo-Controlled Trial. Obstetrical & Gynecological Survey. 53(5):295-296, May 1998.	No Approved Reference	bvgÄyi Kiv nj|
14.	Total Herbal & Nutraceuticals	Capsule Bee Pollen Each capsule contains 500 mg Bee Pollen	1) Bee pollen may have antioxidant and anti-inflammatory activity. 2) Renew skin. 3) Boost immunity. 4) Decrease allergy symptoms.	Bee pollen is contraindicated in people with a known history of atopy or allergy to pollen or plant products because of the risk of hypersensitivity.	New	The Complite Commission E Monograph-Page No:187	Abyjgv`b Kiv †h‡Z cv‡i|	Abyjgv`b 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	‡UKwbK¨vj mve KwgwUi wm×vš	mfvi wm×vš
15.	Total Herbal & Nutraceuticals	Capsule Boswellia Extract (Aflapin) + Chicken Collagen type II (UC-II) Each capsule contains Boswellia Serrata Standardized Extract (Aflapin) 100mg + Chicken Collagen type II (UC-II) 40mg	1) Joint Pain 2) Cartilage formation. 3) To support the structure and function of their body's joints	*No data are available*		1) PDR for Herbal Medicine, Fourth Edition, Page-319-320 and 129. *3) Indian Herbal Pharmacopeia., Page-1409-1410.* 2)Who Monograph of Sleeted Medicinal Plant –Volume - 3-Page 56-68	Abygv`b Kiv †h‡Z cv‡i	Abygv`b Kiv nj	
16.	Total Herbal & Nutraceuticals	Tablet Melatonin Each Tablet contains 5 mg Melatonin	(a) Jet lag, (b) Sleeping difficulties, (c) Cancer prevention, (d) 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	1. British National Formulary (BNF) Page no: 422. 2. USP–Dietary Supplement Compendium. USP32-NF27 Page1556	Abygv`b Kiv †h‡Z cv‡i	Abygv`b 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	‡UKwbK¨vj mve KwgwUi wm×vš	mfvi wm×vš
17.	Total Herbal & Nutraceuticals	Tablet BIOTIN 5 mg Each Tablet Contains 5mg Biotin	Hair loss, brittle nails, skin rash in infants (seborrheic dermatitis), diabetes, Diabetic nerve pain and mild depression.	It is not known if this product interacts with any medicines.	New	Japanese Pharmacopeia (Herbal) Page no: 458 USP – Dietary Supplement Compendium, USP32- NF27, Page -447-448 British Pharmacopeia-2009, Page-714-715	Ncessary Documents not Supplied	bvgÄyi Kiv nj	
18.	Total Herbal & Nutraceuticals	Tablet BIOTIN 10 mg Each Tablet Contains 10 mg Biotin	Hair loss, brittle nails, skin rash in infants (seborrheic dermatitis), diabetes, Diabetic nerve pain and mild depression.	It is not known if this product interacts with any medicines.	New	Japanese Pharmacopeia (Herbal) Page no: 458 USP – Dietary Supplement Compendium, USP32- NF27, Page -447-448 British Pharmacopeia-2009, Page-714-715	Not Accepted deu to but no supplyed Reference Pharmacitucal item	bvgÄyi Kiv nj	
19.	Total Herbal & Nutraceuticals	Capsule Artichoke + L orthonine L aspartame Each capsule contains Artichoke 150mg + L orthonine L aspertam 250 mg	1)Acute & Chronic Hepatitis: Alcoholic liver Damage 2) Fatty Liver: Adjunct to Hepatoxic Drug 3) Liver Cirrhosis: Hepatic Encephalopathy 4) Post Hepatitis Convalescence: Jaundice	No adequate investigations of the use of this drug in children are available. Therefore, it should not be used in children under the age of 12 years.		1) Who Monograph of Selected Medicinal Plant. Volume 4, 2) PDR for Herbal Medicine, P.44-46, 362,339	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b 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	‡UKwbK¨vj mve Kwgwdi wm×vš	mfvi wm×vš
20.	Total Herbal & Nutraceuticals	Tablet L orthonine + aspartame Each Tablet Contains L orthonine L aspertam 500 mg	1)Acute & Chronic Hepatitis: Alcoholic liver Damage 2) Fatty Liver: Adjunct to Hepatoxic Drug 3) Liver Cirrhosis: Hepatic Encephalopathy 4) Post Hepatitis Convalescence: Jaundice	No adequate investigations of the use of this drug in children are available. Therefore, it should not be used in children under the age of 12 years.		1) PDR for Herbal Medicine, P.44-46, 362,339	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
21.	Total Herbal & Nutraceuticals	Tablet Magnesium Oxide Each Tablet Contains Magnesium Oxide 400 mg	a) Osteoporosis. b) Osteoarthritis. c) Control Diabetes mellitus, d) Migraine. e) Premenstrual syndrome, f) Control Blood Pressure, g) Athletes muscle cramp.	Side effects include indigestion or nausea. Possible reaction if allergic to shellfish.	New	1.*Korean Pharmacopeia (Herbal & Supplement)* .Page-KPX 767 2. British National Formulary-BNF-70, Page-43-49 *2. USP-Dietary Supplement Compendium-DSC, Volume - 2 Page No-3752.*	Pharmacitucal item no authentic Documents supplied about reference	bvgÄyi Kiv nj
22.	Total Herbal & Nutraceuticals	Tablet Glucosamine + Chondroitin + MSM Each Tablet Contains Glucosamine 500mg + Chondroitin 200mg+ MSM 250mg Tablet	a) Osteoarthritis b) Knee pain c) Rheumatoid arthritis d) Back pain e) Glaucoma	**Drug interactions:** There have been no reports of significant drug interactions with antibiotics or antidepressants. **Contraindications:** Hypersensitivity reaction to any of its component.	New	1) British National Formulary 70, Page No: 893-894 2) *United States Pharmacopeia 29 and National*	Pharmacitucal item Already in the market as pharma producet	bvgÄyi 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	‡UKwbK¨vj mve KwgwUi wm×vš	mfvi wm×vš
23.	Total Herbal & Nutraceuticals	Tablet Boswellia Serrata +Collagen Type II+ Glucosamine + Chondroitin + MSM Each Tablet Contains Boswellia Cerrata 100mg + Collagen Type II 40mg +Glucosamine 500mg + Chondroitin 200mg+ MSM 250mg Tablet	a) Osteoarthritis b) Knee pain c) Rheumatoid arthritis d) Back pain e) Glaucoma	**rug interactions:** There have been no reports of significant drug interactions with antibiotics or antidepressants. **traindications:** Hypersensitivity reaction to any of its component.	New	1. British National Formulary 70, Page No: 893-894 2. *WHO Monograph of Selected Medicinal Plant, Volueme-4, Page 92-107* *3)Indian Herbal Pharmacopeia., Page-1409-1410.* 3. *United States Pharmacopeia –USP-DSC-2009-2010.* Page No:566-567	Combination Producet but No Reference Product mentioned	bvgÄyi 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	‡UKwbK¨vj mve KwgwUi wm×vš	mfvi wm×vš
24.	Total Herbal & Nutraceuticals	Capsule Zinc 25 Acetate Zinc Acetate 25 mg	a) Zinc Deficiency b) Wilson's disease	**Drug interactions:** There have been no reports of significant drug interactions. SIDE-EFFECTS ▸ Common or very common Gastric irritation (usually transient) common Leucopenia . sideroblastic anaemia	New	1. British National Formulary 70, Page No: 872 3. *United States Pharmacopeia – USP-DSC -2009-2010.* Page No:742-743	Included in BNF as Pharma product	bvgÄyKiv nj |
25.	Total Herbal & Nutraceuticals	Capsule Zinc 50 Acetate Zinc Acetate 50 mg	Zinc Deficiency Wilson's disease	**Drug interactions:** There have been no reports of significant drug interactions SIDE-EFFECTS ▸ Common or very common Gastric irritation (usually transient) mmon Leucopenia . sideroblastic anaemia	New	1. British National Formulary 70, Page No: 872 3. *United States Pharmacopeia – USP-DSC -2009-2010.* Page No:742-743	Included in BNF as Pharma product	bvgÄyKiv nj |
26.	Total Herbal & Nutraceuticals	Tablet Curcumin+Calcium + Magnesium + Vitamin D3+Zinc Curcumin 100mg+ Calcium 500mg + Magnesium 200mg + Vitamin D3-400mg+Zinc 7.5mg Tablet	a) Osteoporosis, b) Osteoarthritis c) Joint Health & Knee pain, d) Rheumatoid arthritis, e) Pregnancy & Lactation, f) Who is in lactose intolerance	**Drug interactions:** There have been no reports of significant drug interactions with antibiotics or antidepressants. **Contraindications:** Hypersensitivity reaction to any of its component.	New	*United States Pharmacopeia 29 and National Formulary 24 and Supplements*. Page:467-469 2. *British National*	Combination Producet but No Reference Product mentioned	bvgÄyKiv nj |

bs	c¯ZKvi‡Ki bvg	Jl‡ai bvg I †R‡bwiK bvg	wb‡`kbv	Contra-indication & Side effect	Status (New Molecule/ Existing)	Reference	‡UKwbK¨vj mve KwgwUi wm×vš	mfvi wm×vš
27.	Total Herbal & Nutraceuticals	**Capsule Chitosan** Each capsules contains 500mg Chitosan	1)Weight Loss 2) Obesity Management	**Pregnancy and breast-feeding** No problems have been reported, but weight loss should not be attempted during pregnancy. **Adverse effects** There are no long-term studies assessing the safety of chitosan. However, chitosan may reduce the absorption of fat-soluble vitamins (A, D, E and K). This has been shown in animals, but not in humans.	**New**	**i.**l. British Pharmacopeia, Part –IV, (Herbal), Page:3448 **i.**2.European Pharmacopeia 5.0, Page:1248 **i.**3. PDR for Nutritional Supplement: Page130-132.	Abyjgv`b Kiv †h‡Z cv‡i	Abyjgv`b Kiv nj
28.	Total Herbal & Nutraceuticals	Capsule BCAA Each Capsule Contains	Exercise performance Muscle building	**Adverse effects** None reported, but there are no long-term studies assessing the safety of BCAAs. Large doses of BCAAs (> 20 g) may increase plasma ammonia levels and may impair water absorption, causing gastrointestinal discomfort	**New**	**ɣ.**l. PDR for Nutritional Supplement: Page100-104. **ɣ.**2. Dietary Supplement, Third Edition, Pamela Mason, Page:28-30	No Valide reference attached	bvgÄyi 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	‡UKwbK¨vj mve KwgwUi wm×vš	mfvi wm×vš
29.	Total Herbal & Nutraceuticals	Capsule Vitex Extract +Myo-inositol + Folic Acid Each Capsule Contains Vitex Extract 50mg +Myo-inositol 500mg + Folic Acid 400mcg	1) Polycystic ovary syndrome (PCOS) 2) Women Infertility. 3) Premenstrual syndrome (PMS) 4) Menopausal complaints	CONTRAINDICATIONS The drug is contraindicated in pregnancy and in nursing Mothers.	**New**	1) PDR for Herbal Medicine- Page -176-177. 312-33 2. Japanese Pharmacopeia, JP XVI- Page Page:861	No Valide reference Document supplyed	bvgÄyi Kiv nj|