(BMSIC/DRUGS/15-05)		
S.N.	***Technical Eligibility Criteria as per NIT***	***Name of Bidder-M/S Modern Laboratories*** ***Address: Unit: Village-Bhud, NH 21A (Near Engg. College Hostel), Baddi, Distt-Solan (H.P.)- 173 205*** ***45-47, Sector D-2, Sanwer Road, Indore- 452015*** ***"Chart has been prepared on the basis of number of downloaded Pg. received (i.e.-1-159)"***
1	Constitution of the Bidding Company/Firm such as Memorandum of Association and Article of Association with complete address. As per Clause 3(c).	Partnership deed Submitted (Pg. No. 31-39)
2	Power of Attorney or Resolution of Board by which the authorised signatory has been authorised by bidder firm to sign the documents. As per Clause 3 (e) of the NIT.	General power of attorney submitted by Shri Anil kharia(Partner) in the name of Arun Kharia (pg28-30)
3	List of item Quoted in prescribed format as Annexure III as per Clause 3 (o)	List of items quoted are NIT S.No. 6, 7, 9,10,11, 16,17,18,19,20,21,22,23,29,35,36,40,41,42,44,45,46,47,48,49,55,56,58,61,62,63,64, 65,81,82,83,87,97,98,106,107,122,124,128,129,130,131,132,134,143, 150,151,152,154,159,160,165,176,179,181,198,233. **(Total 62 items)** (Pg. No. 108-112).
4	• Minimum three years old valid manufacturing licence of the product quoted with latest license renewal certificate. As per clause 3(f). • Approved product list as per the license issued for quoted drugs for minimum three years. As per clause 3(f). • Manufacturing License along with approved product list must be valid till the last date of the submission of tender. As per clause 3(f). Bidders shall submit self attested copies of required manufacturing license and approved product list in support of above mentioned condition and they are required to specify the quoted product in their approved product list by highlighting it. As per Clause 3(f)	Mfg. License in Form 28, License No. 28/17/79, date of issue 09/05/1979 (Address: 45, Sector D-2, Sanwer road, Industrial Estate, Indore), and same License No., date of issue 06/09/2001 (Address: 45, Sector D-2, Sanwer road, Industrial Estate, Indore), in which "due to change in constitution" is mentioned on top of Page, Submitted (Pg. No. 87-88). License in Form 25, License No. 25/31/89, date of issue 06/09/2001 (Address- 45-47, Sector D-2, Sanwer Road, Industrial Area, Indore, Madhya Pradesh) Submitted (Pg. No. 86) . Certificate of Renewal of Licence No. 28/17/79 granted on 06/09/2001, in Form-26 Submitted (Pg. No. 85), Renewed from 01-01-2012 to 31-12-2016. Another Certificate of Renewal of Licence No. 25/31/89 in Form-26 Submitted (Pg. No. 84), Renewed from 01-01-2012 to 31-12-2016 (Address- 45-47, Sector D-2, Sanwer Road, Industrial Area, Indore, Madhya Pradesh) Product approval :- **Details in enclosure**
5	In case of Importer, the bidder (importer) firm must have minimum three years old valid import License of the quoted product. All Quoted products should be accompanied by their invoices, statement and import License showing that the quoted product are being imported and sold in India by the bidder (Importer) firm minimum for Last three years. Import license must be valid on the last date of submission of tender.As per Clause 3(g)	NA
6	Self attested copies of Market standing certificate (in India) of minimum three years, issued by the concerned Licensing Authority from Drugs Control Department for the quoted product. As per Clause 3(h)	Manufacturing and Marketing Certificate No. 5320 (Pg. No. 107), 5335 (Pg. No. 100), 7361 (Pg. No. 94), issued by Licensing authority FDA, M.P. on dated 08-09-2015 , 08-09-2015 and 05-11-2014 repectively Submitted (Pg. No. 101-107, 95-100, 89-94), stating that "the following products are also being marketed for the last three year and have not been cancelled during last three years". (Pg. No.89-107) **Details in Enclosure**

7	Self attested copy of Non Conviction Certificate (NCC), issued by the concerned Licensing Authority from Drug Control Administration of the state for last three years. It should be not more than one year old. As per Clause 3(i).	i)NCC Certificate No. 5331, issued on 08-09-2015, issued by Licensing authority, FDA, M.P. , Submitted, stating "As per the record available with this office and also on the basis of affidavit filed by shri arun kharia s/o late shri P.C. kharia parter of the firm, it is further certified, that siad licence has not been convicted since last three years i.e. 2012-13 , 2013-14 and 2014-15 in the state of madhya pradesh by a court of law under the said Act & rules " (Pg. No. 46) ii) NCC Certificate No. 1581,8106,8865,1501 issued on 18-03-2015,06-12-2014,14-11-2013,03-02-2012 respectively, all issued by Licensing authority, FDA, M.P. , Submitted (Pg. No. 42-45).
8	Self attested copies of WHO-GMP/GMP as per revised Schedule- 'M'/COPP Certificate of the manufacturing unit issued by the Licensing Authority / Drugs Control Department. The GMP certificate must not be older than one year form the last date of submission of tender. As per clause 3(j).	i) A validity certificate of pharmaceutical products issued by Licensing authority, FDA, M.P. vide Letter No. 4884 dated 14-08-2015 in which it is mentioned that "Since applications for renewal is under consideration the validity of certificate of pharmaceutical products granted shall be deemed to be valid 12-02-2016". (Pg. No. 118) Another Certificate No.2/2013 valid upto 12.8.15 The certificate is only for Tablet dosage form. (Pg no.119) ii) GMP Certificate no. M-808/2013, issued on 29-10-2013, by Licensing authority , FDA, M.P. Submitted. The GMP certificate is older than one year from the last date of submission of tender. Certificate is valid upto 29-10-2013 It is mentioned "The firm is following GMP for small volume parentral preparations liquids and dry injectable (Beta-Lactum and Non Beta-Lactum both), Eye / Ear drops, tablets (Beta-Lactum and Non Beta-Lactum both), capsules (Beta-Lactum and Non Beta-Lactum both), Syrup - Liquid and dry both, external liquids, powders - oral and external both , ointments and repacking section only) It is mentioned ".This certificate is valid up to one year from the date of issue." (pg no.122) iv) An order vide no.6964 dt 15.10.2014 of the Office of Commisioner Food Safety & Controller FDA MP, enclosed, wherein it has been stated that the valdity of the GMP Certificate shall be 5 yrs from the date of issue or shall be valid upto validity of the licence, whichever is earljer .(pg no.120)
9	Self attested copies of Maximum Production Capacity Certificate (section wise) issued by concerned Licensing Authority form Drugs Control Department highlighting the quoted product section. As per Clause 3(k).	i) Maximum production capacity certificate , issued by Chartered Accountant, and Submitted (Pg. No. 124). ii) Production capacity certificate, issued by Licensing Authority, FDA, M.P., vide Letter No. 758 dated 29/01/2014 Submitted. (Pg. No. 123) The firm is having installed capacity as mentioned below as per the acknowledgment given by the D.I.C Indore. The annual capacity on per shift basis mentioned are as follow:- Injection Liquid Vials (All Types) of Add New Products- 120 Lac Nos. Injection Ampoules - 240 Lac Nos. Dry Injection Vials- 120 Crore Tablets - 120 Crore Liquid Orals preparations (Bottle)- 135 Lacs Capsules - 40 Crore ORS Powder- 5 Crore Dry Syrup Oral (Bottle)- 90 Lacs Opthalmic Preparation (Eye / Ear Drops)- 90 Lacs External Preparation- 60 Lacs Ointment Preparation - 60 Lacs It is mentioned on the bottom of the page that "This certificate is being issued on the basis of acknowledgement given by DIC , Indore and certificate issued by C.A."
10	An affidavit sworn before first class magistrate/Notary stating that the firm & its quoted product is not black listed currently (as on the last date of submission of the tender) by Central Government/ Central Government Agencies/any state government/any of the state government agencies/any Drug Procurement Agencies or by BMSICL as per Annexure II (clause 3(n) of NIT).	An affidavit sworn before Notary stating that the firm & its quoted product is not black listed currently (as on the date of submission of the tender) by Central Government/ Central Government Agencies/any state government/any of the state government agencies/any Drug Procurement Agencies or by BMSICL as per Annexure II (clause 3(n) of NIT) submitted (Pg. No. 125-128)

11	EMD details (DD number/BG number and date with issuing bank) as per Clause 3(b).	Rs. 5 Lacs, 08/09/2015, 17/2015, Canara Bank (Pg. no. 16)
12	Tender Fee Rs 10,000/- in form of DD as per Clause 3(a).	Rs. 10000/-, DD No. 165656, Canara Bank
13	Self attested copies of the Audited Annual Balance Sheet and Profit and Loss statement showing details of their annual average turnover not less than 25 crores for any three of the last four consecutive financial years. (Auditor/ C.A. Certificate of turnover will not be accepted). As per Clause 3(l)	F.Y. 11-12 Pg. 25 Rs. 24.14 Cr. F.Y. 12-13 Pg. 22 Rs. 28.94 Cr. F.Y. 13-14 Pg. 14 Rs. 55.97 Cr.
14	Self attested copy of Income Tax Return for any three of last four consecutive Assessment years. As per Clause 3(m).	A.Y. 11-12 Pg. 116 A.Y. 12-13 Pg. 115 A.Y. 13-14 Pg. 114
15	Self attested copy of PAN Card of the Bidder Company. As per Clause 3(p)	Pan No. - AACFM5920B, Pg. 40
16	Self attested copy of Certificate of valid Sales tax/VAT registration of the bidder company. As per Clause 3(q)	23811100276, Pg. 27
17	Affidavit declaration regarding acceptance of tender conditions to be submitted by the bidding firm as per Annexure IV (Clause 5(k) of NIT).	Declaration in form of affidavit regarding acceptance of tender submitted by the bidding firm as per Annexure IV (Clause 5(k) of NIT) Submitted (Pg. No. 129)

The facts in clause No-1 to 10 and 17 of the provided sheet has been compiled with due deligence and care, on the basis of document provided by BMSICL, in compliance of letter No 515(15) Dt-21.08.2015 of Health Department, Govt of Bihar and letter no BMSIC/40010/2-2014/1678 Dt 18.09.2015 of GM (Supply Chain), BMSICL, Patna. Inspite, some inadvertent dicrepencies could have been crept in. Humble request to all concerned to bring to notice in due time, if any discrepencies is observed, for rectification.

Kumar 31/12/15

The facts in pages mentioned against the clause No-1 to 10 and 17 of this chart has been evaluated & verified

31.12.15

Modern Laboratories

PRODUCT APPROVAL & MARKET STANDING STATUS

Sl.no.	NITS No	Product Name	Product Approval Date/Validity (Pg. No. / S.No.)	Remarks	Market Standing of more than three years (Pg. No. / S.No.)	Remarks
1	6	Albendazole Oral Suspension IP 200 mg / 5 ml - 10 ml bottle	18.02.2015/31.12.2016 (48/05)	(i) 3 yrs of manufacturing/ date of product approval cannot be ascertained (ii) Pack size not specified	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (107/01)	Pack size not specified
2	7	Albendazole Tab. IP 400 mg	18.02.2015/31.12.2016 (56/23)	3 yrs of manufacturing/ date of product approval cannot be ascertained	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (107/04)	
3	9	Amikacine Inj. IP 100 mg / 2 ml - 2 ml vial	18.02.2015/31.12.2016 (81/22)	(i) 3 yrs of manufacturing/ date of product approval cannot be ascertained (ii) Pack size not specified	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (107/05)	Pack size not specified
4	10	Amikacin Injection IP 250 mg / 2 ml - 2 ml vial	18.02.2015/31.12.2016 (81/23)	(i) 3 yrs of manufacturing/ date of product approval cannot be ascertained (ii) Pack size not specified	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (107/06)	Pack size not specified
5	11	Aminophylline Inj. IP 25 mg / ml - 10 ml. Amp.	18.02.2015/31.12.2016 (81/26)	(i) 3 yrs of manufacturing/ date of product approval cannot be ascertained (ii) Pack size not specified	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (107/08)	Pack size not specified
6	16	Amoxycillin Capsule IP 500 mg	18.02.2015/31.12.2016 (64/04)	3 yrs of manufacturing/ date of product approval cannot be ascertained	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (106/11)	
7	17	Amoxycillin Capsule IP 250 mg	18.02.2015/31.12.2016 (64/03)	3 yrs of manufacturing/ date of product approval cannot be ascertained	Vide Letter No. 7361, dated 05.11.2014 , issued by Licencing authorty F&DA, M.P. (94/16)	
8	18	Amoxycillin Dispersible Tab. IP 125 mg	18.02.2015/31.12.2016 (59/01)	3 yrs of manufacturing/ date of product approval cannot be ascertained	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (106/12)	

9	19	Amoxycillin Oral Suspension IP 125 mg / 5 ml -60ml	18.02.2015/31.12.2016 (62/01)	(i) 3 yrs of manufacturing/ date of product approval cannot be ascertained (ii) Pack size not specified	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (106/13)	Pack size not specified
10	20	Ampicillin & Cloxacillin Injection (Each Vial contains : Ampicillin IP - 500 mg & Cloxacillin IP 500 mg)	18.02.2015/31.12.2016 (79/52)	3 yrs of manufacturing/ date of product approval cannot be ascertained	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (106/15)	
11	21	Ampicillin and Cloxacillin Injection (Each vial contains : Ampicillin - 250 mg & Cloxacillin - 250 mg)	18.02.2015/31.12.2016 (79/49)	3 yrs of manufacturing/ date of product approval cannot be ascertained	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (106/14)	
12	22	Ampicillin Injection IP 500 mg	18.02.2015/31.12.2016 (80/42)	3 yrs of manufacturing/ date of product approval cannot be ascertained	Vide Letter No. 5335, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (99/11)	
13	23	Antacid Tablets (Tablet Containing : Dried Aluminium Hydroxide Gel - 250 mg, Activated Dimethicon 50 mg	18.02.2015/31.12.2016 (55/38)	3 yrs of manufacturing/ date of product approval cannot be ascertained	Vide Letter No. 5335, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (98/40)	
14	29	Azithromycin Oral Suspension IP 200 mg / 5 ml - 15 ml bottle	18.02.2015/31.12.2016 (62/09)	(i) 3 yrs of manufacturing/ date of product approval cannot be ascertained (ii) Pack size not specified	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (106/18)	Pack size not specified
15	35	Calcium Gluconate Inection IP 100 mg / ml - 10 ml. vial	18.02.2015/31.12.2016 (78/80)	(i) 3 yrs of manufacturing/ date of product approval cannot be ascertained (ii) Pack size not specified	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (106/22)	Pack size not specified
16	36	Calcium Tablets (Each tablet Contains : Calcium Carbonate equivalent to 500 mg elemental calcium and Vit D3 - 250 I.U.)	18.02.2015/31.12.2016 (58/17)	3 yrs of manufacturing/ date of product approval cannot be ascertained	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (106/23)	
17	40	Cefadroxil Oral Suspension IP 125 mg / 5 ml - 30 ml bottle	18.02.2015/31.12.2016 (61/17)	(i) 3 yrs of manufacturing/ date of product approval cannot be ascertained (ii) Pack size not specified	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (106/24)	Pack size not specified
18	41	Cefadroxil Tablets IP 500 mg	18.02.2015/31.12.2016 (58/21)	3 yrs of manufacturing/ date of product approval cannot be ascertained	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (105/25)	

19	42	Cefixime Tablets IP 50 mg	18.02.2015/31.12.2016 (58/22)	3 yrs of manufacturing/ date of product approval cannot be ascertained	**Not submitted**	
20	44	Cefixime Tablets IP 100 mg	18.02.2015/31.12.2016 (58/23)	3 yrs of manufacturing/ date of product approval cannot be ascertained	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (105/26)	
21	45	Cefixime Tablets 400 mg	18.02.2015/31.12.2016 (58/27)	3 yrs of manufacturing/ date of product approval cannot be ascertained	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (105/28)	
22	46	Cefotaxime Sodium Injection IP 1 gm vial	18.02.2015/31.12.2016 (76/112)	3 yrs of manufacturing/ date of product approval cannot be ascertained	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (105/32)	
23	47	Cefotaxime Sodium Injection IP 250 mg vial	18.02.2015/31.12.2016 (76/108)	3 yrs of manufacturing/ date of product approval cannot be ascertained	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (105/30)	
24	48	Cefotaxime Sodium Injection IP 500 mg vial	18.02.2015/31.12.2016 (76/110)	3 yrs of manufacturing/ date of product approval cannot be ascertained	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (105/31)	
25	49	Cefotaxime Sodium Injection IP 125 mg vial	18.02.2015/31.12.2016 (77/106)	3 yrs of manufacturing/ date of product approval cannot be ascertained	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (105/29)	
26	55	Ceftriaxone Injection IP 250 mg vial	18.02.2015/31.12.2016 (75/127)	3 yrs of manufacturing/ date of product approval cannot be ascertained	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (105/33)	
27	56	Ceftriaxone Injection IP 500 mg vial	18.02.2015/31.12.2016 (75/130)	3 yrs of manufacturing/ date of product approval cannot be ascertained	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (105/34)	
28	58	Cetrizine Hydrochloride Tablets IP 10 mg	18.02.2015/31.12.2016 (55/48)	3 yrs of manufacturing/ date of product approval cannot be ascertained	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (105/36)	
29	61	Chloroquine Phosphate Suspension IP 50 mg / 5 ml - 60 ml bottle	18.02.2015/31.12.2016 (48/07)	(i) 3 yrs of manufacturing/ date of product approval cannot be ascertained (ii) Pack size not specified	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (105/37)	Pack size not specified
30	62	Chlorpheniramine Injection IP 10 mg / ml - 10 ml vial	18.02.2015/31.12.2016 (74/161)	(i) 3 yrs of manufacturing/ date of product approval cannot be ascertained (ii) Pack size not specified	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (105/39)	Pack size not specified

31	63	Chlorpheniramine Tablets IP 4 mg	18.02.2015/31.12.2016 (54/53)	3 yrs of manufacturing/ date of product approval cannot be ascertained	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (104/40)	
32	64	Ciprofloxacin Eye Drops IP 0.3% w/v - 5ml Pack	18.02.2015/31.12.2016 (67/06)	(i) 3 yrs of manufacturing/ date of product approval cannot be ascertained (ii) Pack size not specified (iii) Composition not as per NIT	Vide Letter No. 5335, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (98/29)	(i) Pack size not specified (ii) Composition not as per NIT
33	65	Ciprofloxacin Hydrochloride Tablets IP 250 mg	18.02.2015/31.12.2016 (54/59)	3 yrs of manufacturing/ date of product approval cannot be ascertained	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (104/41)	
34	81	Dicyclomine HCL and Paracetamol (Each tab contains : Dicyclomine HCL 20 mg & Paracetamol 325 mg)	18.02.2015/31.12.2016 (53/77)	3 yrs of manufacturing/ date of product approval cannot be ascertained	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (104/47)	
35	82	Dicylomine hydrochloride Tablets IP 20 mg	18.02.2015/31.12.2016 (53/78)	3 yrs of manufacturing/ date of product approval cannot be ascertained	Vide Letter No. 7361 dated 05.11.2014 , issued by Licencing authorty F&DA, M.P. (91/76)	
36	83	Dicyclomine Hydrochloride Injection IP 10 mg / ml - 2 ml. Amp.	18.02.2015/31.12.2016 (83/05)	(i) 3 yrs of manufacturing/ date of product approval cannot be ascertained (ii) Pack size not mentioned	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (104/48)	Pack size not specified
37	87	Doxycycline Capsules IP 100 mg	18.02.2015/31.12.2016 (63/23)	3 yrs of manufacturing/ date of product approval cannot be ascertained	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (104/49)	
38	97	Gentamicin Eye Drops IP 0.3% w/v of gentamicin - 5 ml Pack	18.02.2015/31.12.2016 (66/16)	(i) 3 yrs of manufacturing/ date of product approval cannot be ascertained (ii) Pack size not mentioned	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (104/54)	Pack size not specified
39	98	Gentamicin Injection IP 40 mg / ml - 2 ml vial	18.02.2015/31.12.2016 (83/06)	(i) 3 yrs of manufacturing/ date of product approval cannot be ascertained (ii) Pack size not mentioned	Vide Letter No. 7361, dated 05.11.2014 , issued by Licencing authorty F&DA, M.P. (91/85)	Pack size not specified
40	106	Iron and Folic Acid Syrup IP (Each 5 ml contains Ferrous sulphate equivalent to 100 mg of elemental ferrous iron & 0.5 mg of Folic Acid. (To be	18.02.2015/31.12.2016 (60/33)	(i) 3 yrs of manufacturing/ date of product approval cannot be ascertained (ii) Pack size not mentioned	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (104/50)	Pack size not specified

41	107	Iron and Folic Acid Tab. (Each Tablet contains ferrous sulphate equivalent to 20 mg of ferrous elemental Iron and 100 µg of Folic Acid (to be labelled as per Schedule V))	18.02.2015/31.12.2016 (57/49)	(i) 3 yrs of manufacturing/ date of product approval cannot be ascertained (ii) Composition not mentioned in the approval	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (104/51)	Composition not mentioned in the certificate
42	122	Lignocaine Hydrochloride Injection IP 2% w/v - 30 ml vial	18.02.2015/31.12.2016 (73/213)	(i) 3 yrs of manufacturing/ date of product approval cannot be ascertained (ii) Pack size not mentioned	Vide Letter No. 5335, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (97/57)	Pack size not specified
43	124	Magnesium Sulphate Injection IP 500 mg / ml - 5 ml. Amp.	18.02.2015/31.12.2016 (71/230)	(i) 3 yrs of manufacturing/ date of product approval cannot be ascertained (ii) Pack size not mentioned	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (103/58)	Pack size not specified
44	128	Meropenem Injection IP 500 mg	18.02.2015/31.12.2016 (71/232)	3 yrs of manufacturing/ date of product approval cannot be ascertained	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (103/59)	
45	129	Meropenem Injection IP 1000 mg	18.02.2015/31.12.2016 (71/233)	3 yrs of manufacturing/ date of product approval cannot be ascertained	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (103/60)	
46	130	Metoclopramide Injection IP 5 mg / ml - 2 ml. Amp.	18.02.2015/31.12.2016 (71/237)	(i) 3 yrs of manufacturing/ date of product approval cannot be ascertained (ii) Pack size not mentioned	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (103/61)	Pack size not specified
47	131	Methylergometrine Injection IP 0.2 mg / ml - 1 ml. Amp.	18.02.2015/31.12.2016 (71/235)	(i) 3 yrs of manufacturing/ date of product approval cannot be ascertained (ii) Pack size not mentioned	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (103/62)	Pack size not specified
48	132	Methylergometrine Tablets IP 0.125 gm	18.02.2015/31.12.2016 (52/119)	3 yrs of manufacturing/ date of product approval cannot be ascertained	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (103/63)	
49	134	Metoclopramide Tab. IP 10 mg	18.02.2015/31.12.2016 (52/121)	3 yrs of manufacturing/ date of product approval cannot be ascertained	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (103/64)	
50	143	Moxifloxacin Eye Drops IP 0.5% w/v - 5 ml Pack	18.02.2015/31.12.2016 (66/27)	(i) 3 yrs of manufacturing/ date of product approval cannot be ascertained (ii) Pack size not mentioned (iii) Not as per NIT specification	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (103/66)	(i) Pack size not specified (ii) Not as per NIT specification

51	150	Ofloxacin Ophthalmic Solution IP 0.3% w/v - 10 ml Pack	18.02.2015/31.12.2016 (65/30)	(i) 3 yrs of manufacturing/ date of product approval cannot be ascertained (ii) Pack size not mentioned	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (103/68)	Pack size not specified
52	151	Ofloxacin Oral Suspension IP 50 mg / ml - 30 ml	18.02.2015/31.12.2016 (72/47)	(i) 3 yrs of manufacturing/ date of product approval cannot be ascertained (ii) Pack size not mentioned	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (103/69)	Pack size not specified
53	152	Ondansetron Hydrochloride Injection 2 mg / ml - 2 ml. Amp.	18.02.2015/31.12.2016 (83/08)	(i) 3 yrs of manufacturing/ date of product approval cannot be ascertained (ii) Pack size not mentioned	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (103/70)	Pack size not specified
54	154	Ondansetron Tablets IP 4 mg	18.02.2015/31.12.2016 (51/145)	3 yrs of manufacturing/ date of product approval cannot be ascertained	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (103/71)	
55	159	Paracetamol Syrup IP 125 mg / 5 ml - 60 ml	18.02.2015/31.12.2016 (72/56)	(i) 3 yrs of manufacturing/ date of product approval cannot be ascertained (ii) Pack size not mentioned	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (102/73)	Pack size not specified
56	160	Pheniramine Injection IP 22.75 mg / ml - 2 ml. Amp.	18.02.2015/31.12.2016 (83/09)	(i) Composition not as per NIT (ii) 3 yrs of manufacturing/ date of product approval cannot be ascertained (iii) Pack size not mentioned	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (102/76)	(i) Composition not as per NIT (ii)Pack size not specified
57	165	Piperacillin & Tazobactum for Injection (Piperacillin 4 gm & Tazobactum 500 mg) - 4.5 gm vial	18.02.2015/31.12.2016 (70/273)	3 yrs of manufacturing/ date of product approval cannot be ascertained	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (102/79)	
58	176	Rabeprazole Sodium for Injection 20 mg	18.02.2015/31.12.2016 (69/288)	3 yrs of manufacturing/ date of product approval cannot be ascertained	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (101/84)	
59	179	Ranitidine Tablets IP 150 mg	18.02.2015/31.12.2016 (50/166)	3 yrs of manufacturing/ date of product approval cannot be ascertained	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (101/85)	
60	181	Salbutamol Tablets IP 4 mg	18.02.2015/31.12.2016 (49/170)	3 yrs of manufacturing/ date of product approval cannot be ascertained	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (101/88)	

61	198	Tramadol Hydrochloride Injection 50 mg / ml - 2 ml. Amp.	18.02.2015/31.12.2016 (68/305)	(i) 3 yrs of manufacturing/ date of product approval cannot be ascertained (ii) Pack size not mentioned (ii) Composition as per approval is 50 mg / 2 ml whereas NIT specification is 50 mg / ml	Vide Letter No. 5320, dated 08.09.2015 , issued by Licencing authorty F&DA, M.P. (101/89)	Pack size not specified
62	233	Menadione Sodium Bisulphite Injection 10 mg - 1 ml. Amp.	18.02.2015/31.12.2016 (82/16)	(i) 3 yrs of manufacturing/ date of product approval cannot be ascertained (ii) Pack size not specified	**Not submitted**	