Tender Reference no.-BMSIC/DRUG/16-07

S.N	Technical Eligibility Criteria as per NIT	Firm Name - M/s Celon Laboratories Pvt. Ltd. Corporate Address: Plot No. 264 Patrika Nagar, Madhapur Hyderabad-500081 Manufacturing Unit Address:-Plot No.-2, Aleap Industrial Estate, Gajularamaram R.R. District-500090
1	Constitution of the Bidding Company/Firm such as Memorandum of Association and Article of Association with complete address. As per Clause 3(c).	(a) Self attested copy of coverpage of Memorandum of Association & Articles of Association of Celon Laboratories Private Ltd. is submitted (Pg. No. - 138 & 343 & also 366) (b) Self attested copy of memorandum of Association of Celon Laboratories Private Ltd. is submitted (Pg. No. - 144 to 150) and 371 to 375) (c) Self attested copy of Articles of Association of Celon Laboratories Private Ltd. is submitted (Pg. No. - 151 to 208) (d) Self attested copy of certificate of incorporation of **Medvin Labs Private Ltd.** on 23rd Jan 2001 is submitted (Pg. No. - 143 & 348) and also 370 to 449). (e) Self attested copy of certificate of incorporation for the change of company name from **Medvin Labs Private Ltd.** to **Galen Labs Private Ltd**. on 19th October 2002 is submitted (Pg. No. - 142 & 347) and also 368) (f) Self attested copy of certificate of incorporation for the change of company name from **Galen Labs Private Ltd.** to **Celon Laboratries Private Ltd**. on 29th may 2007 is submitted (Pg. No. - 141 & 346 and also 369) (g) Self attested copy of certificate of incorporation for the change of Company name from **Celon Laboratories Private Ltd. to Celon Laboratories Ltd.** on 15th June 2007 is submitted (Pg. No. - 140 and 345 and also 367) (h) Self attested copy of certificate of incorporation for the change of company name from Celon Laboratories Ltd. to Celon Laboratories Private Ltd. on 18th August 2015 is submitted. (Pg. No. - 139 and 344) (i) Copy of Bidder information/Bidder details as per Annexure V of NIT is submitted (Pg. No. - 29)
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.	(a) Self attested copy of Power of Attorney is submitted where in it is stated that Mr. Yogesh Kumar Chopra to be its true and lawful constituted Attorney for and on behalf of the compnay and execute and perform to sign, submit and follow up the matters relating to tenders filed with any government authority or any other authority in india. (Pg. No. - 350 to 351)
3	List of item Quoted in prescribed format as Annexure III as per Clause 3 (o)	(a) Copy of List of items quoted is submitted as per Annexure - II of NIT. No. of quoted items - 17 (seventeen)

Sl.No.	NIT sl. No.	Name of Drug	Specification/strength
1	17	Cyclophosphamide Inj.	500mg
2	18	Cyclophosphamide Inj. IP	200mg
3.	21	Dopamine Inj. IP	40mg/ml, 5 ml vial
Note:- At sl. no. 3 of list of quoted items the product is mentioned as Dopamine injection IP 40mg/ml and specification column as USP.			
4.	27	Heparin Sodium Injection IP	5000 Units/ml, 5 ml vial
5.	57	Propofol Injection IP	10mg/ml, 20 ml vial
6.	74	Tranexamic Acid Injection IP	100mg/ml, 5 ml ampoule
7.	77	Vancomycin intravenous infusion IP	1000mg
8.	78	Vancomycin intravenous infusion IP	500mg
9.	79	Vasopressin Injection IP	20 IU/ml, 1 ml ampoule
10	80	vincristine Injection IP	1mg/ml
11.	84	Fluorouracil Injection IP	250mg/5ml, 5 ml
12.	94	Heparin Sodium Injection IP	5000 IU/5ml, 5 ml vial
13.	101	Teicoplanin Injection	200mg
14.	102	Teicoplanin Injection	400mg.
15.	103	Vecuronium Bromide Injection	4mg/2ml, 2 ml ampoule
16.	105	Methotrexate Tab. IP	2.5mg., 10x10
17	106	Methotrexate Injection IP	50mg/2ml, 2 ml ampoule

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). • In Case of those drugs which are notified first time in IP 2014 then Manufacturing and Marketing license should be in any official compendium (USP/BP/IP) along with current valid License in IP in continuation. • Market standing certificate & Manufacturing certificate issued by the Licensing Authority as a Manufacturer for each drug quoted for the last 3 years (Certificate should be enclosed with list of items) except for the drugs falling under the category of 'New Drug' as defined by CDSCO (Central Drugs Standard Control Organization). If permission in Form 46 from DCGI has been obtained, then the 3 years Manufacturing & Market standing clause will be relaxed. • For all regulated products, the bidder should have at least two years of manufacturing and marketing experience of the particular items as a manufacturer for each regulated product quoted in the tender. However, this would not apply to regulated products which have been licensed by DCGI less than two years ago. A certificate from DCG (I) shall be required for all new regulated products to this effect. • FFS (Flow Fill & Seal Process) Technology will be accepted wherever applicable. Bidders shall submit self attested copies of required manufacturing license and approved product list in support of above mentioned	(a) Self attested copy of Licence to manufacture in form 25 of **M/s CELON LABORATORIES LTD. (No. 25/HD/AP/2003/F/CC** dated 27-05-2003) issued by Licensing Authority , Hyderabad is submitted. Wherein it is stated that the license will be in force from **05-06-2007 to 04-06-2012.** (Pg. No. - 331 to 332) (b) Self attested copy of certificate of Renewal in form 26 of Licence No. 25HD/AP/2003/F/R date 27/05/2003 of **M/s CELON LABORATORIES LTD.** issued by Director, Drugs control Administration, Hyderabed is submitted where in it is stated that the Licence has been renewed from **05-06-2012 to 04-06-2017**. (Pg. No. - 335 to 336) (c) Self attested copy of Licence to manufacture in form 28 (No. 14/RR/AP/2008/F/G date 24/04/2008) of M/s CELON LABORATORIES LTD. issued by Director , Drug control Administraion , Hyderabed is submitted where in it is stated that the licence shall be in force from 24/04/2008 to 23/04/2013 (Pg. No. - 341) (d) Self attested copy of a letter (No. L. Dis . No. 01144/P & B/2015 dt 26-03-15 regarding application for renewal of drug licence in form - 28 for period 24/04/2013 to 23/04/2018 is received in the office on 20/04/2013 issued by Deputy Director & Certifying Authority, Drugs Control Administration, Government of Telangana is submitted. (Pg. No. - 342) (e) Self attested copy of letter (L. Dis. No. - 19119/E(M)/TS/2015 dated 21-03-2016) issued by licensing & controlling Authority, Drug control Administration, Govt. of Telangana is submitted. Wherein it is stated that the Drug manufacturing Licence in **Form- 28** bearing No. 14/RR/AP/2008/F/CC dt. 24/04/2008 duly renewed from 24/04/2013 to 23/04/2018 and **effectig change of constitution of the firm from M/s Celon laboratories Ltd. to M/s Celon laboratories Private Ltd. w.e.f. 18/08/2015 for the products mentioned in the list enclosed. The licence is valid from 18/08/2015 to 17/08/2020. (Pg. No. - 327 to 328)** **Note - New Drug Mfg. Licence in Form 28 after change of constitution not submitted.** (f) Self attested copy of letter (L. Dis. No. 19118/E(M)/TS/2015, dated 21.03.2016 issued by Licensing & Controlling Authority, Drug Control Administration, Govt. of Telangana is submitted where in it is stated that the Drug Manufacturing License in Form 25 bearing No. 25/HD/AP/2003/F/CC, dated 27.05.2003 **effecting change of constitution of the firm M/s Celon Laboratories Limited to M/s Celon Laboratories Private Limited w.e.f. 18.08.2015 for the products mentioned in the list enclosed. The license is valid from 18.08.2015 to 17.08.2020. (Page No. 329-330)** **Note - New Drug Mfg. License in Form 25 change of constitution not submitted.** (g) Approved product list under L. Dis. No. 19119/E(M)/TS/2015. Renewal & change of constitution of M/s Celon Laboratories Pvt. Ltd. in Form 28 bearing No. 14/RR/AP/2008/F/CC, dated 24.04.2008. NIT S.N. Page No. 17----------266, 18----------266 21----------254 (Mentioned as Dopamine Hydrochloride Injection **USP 200 mg/5 ml), 27----------263** (Mentioned as Heparin Injection IP **25000 IU/5 ml)** 57----------259 (Mentioned as Propopal Injection IP **200 mg/20 ml),** 74----------257 (Mentioned as Tranexamin Acid Injection IP **500 mg/5 ml)** 77----------265 (Mentioned as Vancomycin **Hydrochloride** for Intravenous Infusion IP 1000 mg) 78----------264 (Mentioned as Vancomycin **Hydrochloride** for Intravenous Infusion IP 500 mg), 79----------261 80----------258 (Mentioned as Vincristine Sulphate Injection IP 1 mg), 84----------260 94----------262 (Mentioned as **Heparin Injection** IP 5000 IU/5 ml instead of **Heparin Sodium Injection** IP 5000 IU/5 ml) 101---------256, 102---------257, 103---------265 (Mentioned as **Vecuronium Bromide for Injection 4 gm instead of Vecuronium Bromide for Injection 4 mg/2 ml)** 105---------269, 106---------255 Documentary evidence in support of additional item to M/s Celon Laboratories Ltd. in Form 28 bearing No. 14/RR/AP/2008/F/G valid up to 2013 for various products submitted. (Page No. 271 to 303) (h) Self attested copy of letter no. 7618/BIM/2007, dated 14.06.2007 issued by Director, Drug Control Administration, Andhra Pradesh to M/s Celon Laboratories bearing license no. 25/HD/AP2003/F/CC granted on 27.05.2003 in From 25 with change of constitution w.e.f. 05.06.2007 and it is valid up to 04.06.2010 and change of name of your firm from M/s Galen Labs. Pvt. Ltd. to M/s Celon Laboratories Limited w.e.f. 05.06.2007 for the products mentioned in the list enclosed. Letter also state that the products earlier approved to M/s Galen Labs Pvt. Ltd. under Drug License 28/HDG/AP/2003/F/G granted on 27.05.2003 are hereby deleted from the approved list. Further Drug license in Form 28 bearing no. 25/HD/AP/2003/F/G is here by cancelled. (Page No. 305-306) (i) Self attested copy of Manufacturing & Marketing Certificate for the quoted product is submitted (Except the two products). (Page No. 309-319) 17----------315, 18----------, 21----------318, 27----------312, 57----------318, 74----------310, 77----------317, 78----------319, 79----------318, 80----------316, 84----------312, 94----------318, 101---------309, 102---------317, 103---------309, 105---------312, 106---------
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)	NA
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).	Self attested copy of Non Conviction Certificate (L. Dis. No. NC/RLA/DI/MDL/HYD/29/04/2016, dated 30.04.2016 issued by Licensing & Controlling Authority, DCA, Govt. of Telengana is submitted where in it is stated that M/s Celon Laboratories Private Limited holding drug manufacturing license in From 25 bearing no. 25/HD/AP/2003/F/CC, dated 27.05.2003 valid up to 17.08.2020 & in Form 28 bearing no. 14/RR/AP/2008/F/CC, dated 24.04.2008 valid up to 17.08.2020 has not been convicted at any instance during the last three years under Drugs & Cosmetics Act, 1940 and rules there under. (Page No. 321)

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(i).	Self attested copy of G.M.P. certificate (L. Dis. No. 5185/E(M)/TS/2016, dated 3.04.2016) issued by Licensing & Controlling Authority, Drugs Control Administration, Government of Telangana is submitted where in it is stated that this certificate is valid up to 08.12.2016. (Page No. 326)
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.In case of Importer An affidavit (With Stamp) sworn before first class magistrate/Notary stating the batch production capacity of the firm and also that said production (Importing)capacity shall be adequate for requirement laid in NIT. Importers will have also to submit Invoices/Evidence of import in items of said product with quantity details. As per Clause 3(k).	Self attested copy of Production Capacity certificate (L. Dis. No. 08789/E(M)/TS/2016, dated 11.08.2016) issued by Joint Director & Licensing Authority, Drug Control Administration, Government of Telengana is submitted. (Page No. 352)
10	An affidavit (with stamp)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).	Self attested copy of notarised affidavit (with stamp) for Non-Blacklisting is submitted as per Annexure II of NIT. (Page No. 322-323) **Note - Affidavit submitted by Authorized signatory, Mr. Yogesh Kumar Chopra.**
11	EMD details (DD number/BG number and date with issuing bank) as per Clause 3(b) .	"DD NO.":-504552" (ICIC Bank) `3,00,000/- Page No:-325
12	Tender Fee Rs 10,000/- in form of DD as per Clause 3(a).	"DD No":-504553" (ICIC Bank) `10,000/- Page No:-365
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)	**FY-2012-13 `4761.97/-(in Lacs)- Page No :-70** **FY-2013-14 `3951.58/- (in Lacs)- Page No :-70** **FY-2014-15 `5820.47/-(Lacs)- Page No:-37** **Note: -Revenue from operation net taken as "turnover"**
14	Self attested copy of Income Tax Return for any three of last four consecutive Assessment years. As per Clause 3(m).	FY-2015-16 Page No:-359 Fy-2013-14 Page No-360 Fy-2014-15 Page No:-358
15	Self attested copy of PAN Card of the Bidder Company. As per Clause 3(p)	PAN NO:- "AACCG1654P" Page No:-361
16	Self attested copy of Certificate of valid Sales tax/VAT registration of the bidder company. As per Clause 3(q)	TIN No:- "36210111802" Page No:-362
17	Affidavit (with stamp)declaration regarding acceptance of tender conditions to be submitted by the bidding firm as per Annexure IV (Clause 5(k) of NIT).	Self attested copy of notarised affidavit (with stamp) regarding acceptance of tender conditions is submitted as per Annexure IV of NIT. (Page No. 353-354)

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 1146(15) Dt-05.11.2016 of Health Department, Govt of Bihar . Inspite, some inadvertent discrepencies could have been crept in. Humble request to all concerned to bring to notice in due time if any discrepencies is observed for rectification.

Hari