

TENDER NO. BMSIC/DRUGS/20-16

Check list - I: Evaluation sheet to be used for preliminary evaluation of technical bids

Name of the Bidder : - EUROLIFE HEALTHCARE PVT. LTD. Total Number of Pages Submitted in bid document: - 01 TO 156

Sl. No	SBD clause serial No	Description of the Clause/ Technical Eligibility Criteria as per SBD as mentioned in the Check List	Document/ fees submission status (Yes/ No)	Page number	Do the provided documents meet the eligibility criteria ? (Yes/ No)	Remarks
1	3. (a)	Tender Fee (Non -Refundable) of Rs 10,000/- in form of Demand Draft drawn in Favor of "Managing Director, Bihar Medical Services and Infrastructure Corporation Limited" payable at Patna. This fee is payable only once for one tender irrespective of items contained therein.	YES	75	YES	-
2	3.(b)	Bidder are required to submit Earnest Money Deposit in the form of Demand Draft / Bank Guarantee drawn in favor of Managing Director, Bihar Medical Services and Infrastructure Corporation Limited from any Scheduled/Nationalized bank payable at Patna as per following table :- No. of drugs quoted Upto 5 drugs:- Rs1,00,000/- (One Lakh only) For 6 to 10 Drugs:- Rs 2,00,000/- (Two Lakh only) For 11 to 15 Drugs:- Rs 3,00,000/- (Three Lakh only) For 16 to 20 Drugs:- Rs 4,00,000/- (Four Lakh only) More than 20 Drugs:- Rs 5,00,000/- (Five Lakh only)	YES	71-74	YES	-
3	3.(c)	Documentary evidence of the constitution of the company/firm/Proprietorship such as Memorandum and Articles of Association, Partnership Deed etc. should be submitted with details of the Name, Address, Telephone Number, Fax Number, e-mail address of the firm and of the Managing Director / Partners/ Proprietor should be submitted.	YES	20-25	YES	-
4	3.(d)	The details of Bidder Name, Address, Telephone Number, Fax. Number, e-mail address of the bidder and of the Managing Director / Partners / Proprietor should be submitted in Annexure-V.	YES	153	YES	
5	3.(e)	Power of Attorney or Resolution of the Board by which the authorized signatory has been authorized by the bidder firm to sign the documents should be submitted.	YES	26	YES	
6	3.(f)	Copies of the Audited Balance Sheet and Profit and Loss statement showing details of their annual average turnover not less than 25 Crores (Twenty-Five) for any three of the last four consecutive financial years (Auditor/CA certificate of turnover will not be accepted). Self-attested copies are to be submitted.	YES	79 & 81 & 83	YES	-
7	3.(m)	Copy of Income Tax Return for any three of last four Consecutive Assessment years should be submitted (self-attested).	YES	84-87	YES	-
8	3.(p)	Copy of PAN Card of the bidder company should be submitted (self-attested).	YES	88	YES	-
9	3.(q)	Copy of certificate of valid GST registration of the bidder company should be submitted (self-attested).	YES	90	YES	-

Arundhati Singh

TENDER NO. BMSIC/DRUGS/20-16

Check list - II: Evaluation sheet to be used for preliminary evaluation of technical bids

Company Name:- M/s Eurolife Healthcare Pvt. Ltd, Addredd- Khasra No. 520 Village Bhagwanpur , Roorkee, Distt Haridwar
Total Number of Pages Submitted in bid documents:- 156

Sl. No	SBD Clause serial No	Description of the Clause/ Technical Eligibility Criteria as per SBD, as mentioned in Check List	Document/ fees submission status (Yes/ No)	Page Number as numbered by BMSICL	Do the provided documents meet the eligibility criteria ? (Yes/ No)	Remarks
1	3.(d)	The details of Bidder Name, Address, Telephone Number, Fax, Number, e-mail address of the bidder and of the Managing Director / Partners / Proprietor should be submitted in Annexure-V.	Yes	153	Yes	
2	3.(e)	Power of Attorney or Resolution of the Board by which the authorized signatory has been authorized by the bidder firm to sign the documents should be submitted.	Yes	26	Yes	
3		Minimum three years old valid Manufacturing License of the product quoted with latest license renewal certificate.	Yes	28-31	Yes	
4		Approved product list as per the license issued for quoted drugs for minimum three years.	Yes	37-38	Yes	
5		Manufacturing License along with approved product list must be valid till the last date of the submission of tender.	Yes	28-38	Yes	
6		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.	NA	NA	NA	
7		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 Organisation). If permission in Form 46 from DCGI has been obtained, then the 3 years Manufacturing & Market standing clause will be relaxed. The provisions of Rule 122 E of Drugs and Cosmetics Act rule 1945 shall be applicable.	Yes	46-48	Yes	
8	3.(f)	Explanation- In case of Importers Permission in Form 45 from DCGI is required as Per Drugs & Cosmetics Act 1940 & Rules 1945. For all new 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 DCGI (I) shall be required for all new regulated products to this effect.	NA	NA	NA	

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9		FFS (Flow Fill & Seal Process) Technology will be accepted wherever applicable.	Yes	50-51	Yes	
10		Valid Pollution Control Clearance Certificate in accordance with Water [Prevention and control of Pollution] Act, 1974 & Air [Prevention and control of Pollution] Act, 1981 and Hazardous Wastes (Management, Handling & Trans Boundary Movement) Rules 2008 (Self Attested Copy of Certificate to be enclosed) (as applicable).	Yes	52-56	Yes	
11		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.	Yes	28-38	Yes	
12	3.(g)	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.	NA	NA	NA	
13	3.(h)	Bidder must have Market Standing Certificate (in India) of minimum three years issued by the concerned Licensing Authority from Drugs Control Department for the quoted product. Self-attested copies are to be submitted.	NA	NA	NA	
14	3.(i)	Non-Conviction Certificate (NCC) issued by the concerned Licensing Authority from Drugs Control Administration of the state for last three years should be submitted. It should be not more than one year old. Self-attested copies are to be submitted.	Yes	62	Yes	
15	3.(j)	WHO-GMP/GMP (Good Manufacturing Practice) 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 from the last date of submission of tender. Self-attested copies are to be submitted. Explanation- Generally the GMP Certificate issued for one-year validity. Hence the provision that it should not be older than one year from the last date of submission of tender implies mutatis mutandis that the GMP certificate should remain valid till the last date of submission of tender.	Yes	63	Yes	
16	3.(k)	Maximum Production Capacity Certificate (section wise) issued by concerned Licensing Authority form Drugs Control Department highlighting the quoted product section. Self-attested copies are to be submitted. 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. Importer will have also to submit Invoices/Evidence of import in items of said product with quantity details.	Yes	64-68	Yes	

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17	3.(n)	The tenderer should give 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 date of submission of the tender) by Central Government / Central Government agencies/any state government or any of the state government agencies / or any Drug procurement agencies or by BMSICL as per Annexure-II.	Yes	69-70	Yes	
18	3 (o)	List of item quoted in prescribed format as per Annexure-III duly signed.	Yes	27	Yes	
19	5.(k)	An Affidavit (with stamp) regarding acceptance of tender conditions to be submitted by the bidding firm as per Annexure-IV.	Yes	91	Yes	
20	5.(l)	Filled check list as per given Annexure-VI to be submitted at the time of uploading the bid.	Yes	154-156	Yes	
21	7.(c)	Power of Attorney or Resolution of the Board by which the authorized signatory has been authorized by the bidder firm should sign the documents in cases where person other than the Managing Director/Managing Partner or sole Proprietor signs the document.	Yes	26	Yes	
22	10.(c)	If a particular document/Certificate to be uploaded as specified in bid, is not applicable for a bidder, the bidder shall attach a scanned copy of declaration in the letter head stating that the specific document is not applicable/exempted for the bidder in connection to this tender.	NA	NA	NA	
23	2(d) Explanation	The price quoted by the bidders must not exceed the ceiling price as fixed by NPPA (National Pharmaceutical Pricing Authority) as per the provisions of "Drugs Price Control Order" and the quoted rate should be at least 20% less than its MRP. In extraordinary case the Managing Director has discretion to take decision. Explanation- In order to ensure procurement of the tendered drugs at the most appropriate and competitive rate, the bidders are directed to quote their lowest price as compared to the Rates provided to their respective Distributors/Dealers/Wholesalers/Carrying and Forwarding Agents/Authorized depot sales point in the State of Bihar. A Notarized affidavit to this effect on a Rs 100/- Non-Judicial Stamp Paper should be submitted with the Bid in the prescribed format as mentioned in Annexure-X.	Yes	19	Yes	
24	5 (j)	PERFORMANCE STATEMENT as per Annexure-IX submitted with the bid document should be self-attested by the authorized signatory of the firm with official seal, wherever required.	Yes	92-94	Yes	
25	-	Production Capacity Statement (Self Declaration) Annexure- VIII	Yes	18	Yes	


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Check list - III: Evaluation sheet to be used for preliminary evaluation of technical bids

Name- M/s Eurofife Healthcare Pvt. Ltd, Address- Khasra No. 520 Village Bhagwanpur, Roorkee, Distt Haridwar

Total Number of Pages Submitted in bid documents- 156

Sheet to be used for verification of product approval and market standing

		Name of the Quoted Drug		Pharmacopoeial Specification/ Strength		Pack Size		Dosage Form		Approval Details			Manufacturing Certificate	
Sl. No	NIT Sl. No	As per NIT	As per Approval	As per NIT	As per Approval	As per NIT	As per Approval	As per NIT	As per Approval	First Approval	Approved Upto	Approved in Brand /Generic Name		
	1	19	Linezolid Injection	Linezolid I.V. Injection 2mg/ml	2mg/ml	Each 100 ml contains: Linezolid IP 200 mg Dextrose (Anhydrous) IP 5% w/v water for Injection IP q.s (page no. 34)	300 ml bottle	Not mentioned	Infusion	Injection	24.01.2018	16.10.2022	Generic	Submitted (page no. 47 & 48)
	2	28	Multiple Electrolyte & Dextrose Injection Type I [Form, Fill & Seal process]	Multiple Electrolyte & Dextrose Injection Type I IP 500 ml	Strength as Per IP	Each 100 ml contains: Dextrose (Anhydrous) IP 5,000 gm Potassium Chloride IP 0.130 gm Dibasic Potassium Phosphate USP 0.026 gm Sodium Acetate IP 0.320 gm Magnesium Chloride IP 0.031 gm Water for Injection IP q.s (page no. 38)	500ml Bottle	500 ml	Infusion	Injection	24.01.2018	16.10.2022	Generic	Submitted (page no. 47 & 48)
	3	29	Multiple Electrolyte & Dextrose Injection Type III [Form, Fill & Seal process]	Multiple Electrolyte & Dextrose Injection Type III IP 500 ml	Strength as Per IP	Each 100 ml contains: Dextrose (Anhydrous) IP 5.0 gm Sodium Chloride IP 0.10 gm Dibasic Potassium Phosphate USP 0.13 gm Sodium Acetate IP 0.28 gm Potassium Chloride IP 0.15 gm Water for Injection IP q.s (page no. 38)	500ml Bottle	500 ml	Infusion	Injection	24.01.2018	16.10.2022	Generic	Submitted (page no. 47 & 48)

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Check list - IV: Evaluation sheet to be used for preliminary evaluation of technical bids

ame- M/s Eurolife Healthcare Pvt. Ltd, Addredd- Khasra No. 520 Village Bhagwanpur , Roorkee Distt Hardwar

Total Number of Pages Submitted in bid documents:- 156

Sheet for verification of licence details

Sl. No.	NIT Sl. No	Name of the Drug as per NIT	Manufacturer		Importer		Validity		GMP/ WHO GMP/ COPP (Import)	
			Forms Number	Manufacturing license Number	Forms Number	Import license Number	From	To	From	To
1	19	Linezolid Injection	28- D	119/UA/SC/P (LVP)/2007	-	-	24.01.2018	16.10.2022	24.02.2020	24.02.2023
2	28	Multiple Electrolyte & Dextrose Injection Type I [Form, Fill & Seal process]	28- D	119/UA/SC/P (LVP)/2007	-	-	24.01.2018	16.10.2022	24.02.2020	24.02.2023
3	29	Multiple Electrolyte & Dextrose Injection Type III [Form, Fill & Seal process]	28- D	119/UA/SC/P (LVP)/2007	-	-	24.01.2018	16.10.2022	24.02.2020	24.02.2023

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