

TENDER NO. BMSIC/DRUGS/19-09

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

Company Name : Cadila Pharmaceuticals Ltd.

Total Number of Pages 241 Submitted in bid documents:

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	15-14	Yes	
2	3.(b)	A bidder is required to submit Earnest Money Deposit in the form of Demand Draft/ Bank Guarantee of Rs. 1,00,000/- (one Lakh) drawn in favor of Managing Director, Bihar Medical Services and Infrastructure Corporation Limited from any Scheduled/Nationalized bank payable at Patna.	Yes	20-16	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	57-21	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	58	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	61	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 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	191-188& 147-144	Yes	

Abdullah Khan
10/12/12

12/12/12

314

7	3.(m)	Copy of Income Tax Return for any three of last four Consecutive Assessment years should be submitted (self-attested).	Yes	223-221	Yes	
8	3.(p)	Copy of PAN Card of the bidder company should be submitted (self-attested).	Yes	232	Yes	
9	3.(q)	Copy of certificate of valid GST registration of the bidder company should be submitted (self-attested).	Yes	229-228	Yes	

Chandra Kumar
10/14/19

Sant
12/12/19

TENDER NO. BMSIC/DRUGS/19-09

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

Company Name:- Cadila Pharmaceutical Ltd. Industrial Growth Centre, J & K SIDCO, Samba - 184121 (J & K)

Total Number of Pages Submitted in bid documents:- 241

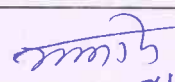
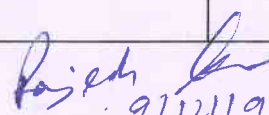
Sl. No	SBD Clause serial No	Description of the Clause/ Technical Eligibility Criteria as per SBD, as mentioned in Check List	Document/ fees submission status (Yes/ No)	Page Number as numbered by BMSICL	Do the provided documents meet the eligibility criteria ? (Yes/ No)	Remarks
1	3.(d)	The details of 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	58	Yes	
2	3.(e)	Power of Attorney or Resolution of Board by which the authorised signatory has been authorized by the bidder firm to sign the documents. Should be submitted.	-	-		
3		Minimum three years old valid Manufacturing License of the product quoted with latest license renewal certificate.	Yes	63 to 70	Yes	
4		Approved product list as per the license issued for quoted drugs for minimum three years.	Yes	71 to 79	Yes	
5		Manufacturing License along with approved product list must be valid till the last date of the submission of tender.	Yes	63 to 79	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			

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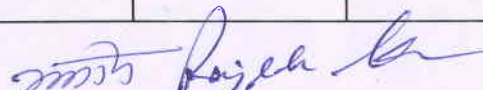
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7	3.(f)	Market standing certificate & Manufacturing certificate issued by the Licensing Authority as a Manufacturer for each drug quoted for the last 3 years (Certificate should be enclosed with list of items) except for the drugs falling under the 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	80	Yes	
8		If permission in Form 46 (For manufacturers) / Form 45 (For Importers) from DCGI has been obtained, then the 3 years manufacturing/ Import & Market standing clause will be relaxed . The provisions of Rule 122 E of Drugs and Cosmetics Act rule 1945 shall be applicable.	NA			
9		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.	NA			
10		FFS (Flow Fill & Seal Process) Technology will be accepted wherever applicable	NA			
11		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).	Yes	82 to 86	Yes	
12		NOTE: 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.	NA			
13	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			

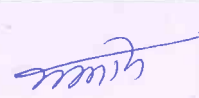
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14	3.(h)	Bidder must have Market Standing Certificate (in India) of minimum three years issued by the concerned Licensing Authority from Drugs Control Department for the quoted product. Self-attested copies are to be submitted.	NA			
15	3.(i)	Non Conviction 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. For Surgicals In case of Non-Drug item "the bidder shall submit an affidavit on Non-Judicial stamp paper stating that the quoted product is not covered under Drugs & Cosmetics Act."	Yes	87-94	Yes	
16	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	95-100	Yes	
17	3.(k)	Maximum Production Capacity Certificate (section wise) issued by concerned Licensing Authority from 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	101	Yes	
18	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	237, 238	Yes	
19	3.(o)	List of item quoted in prescribed format as per Annexure-III duly signed.	Yes	62	Yes	


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20	5.(k)	An Affidavit (with stamp) regarding acceptance of tender conditions to be submitted by the bidding firm as per Annexure-IV .	Yes	231	Yes	
21	5.(l)	Filled check list as per given Annexure-VI to be submitted at the time of uploading the bid.	Yes	10 to 13	Yes	
22	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	61	Yes	
23	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	-	-	
24	2(d)	A Notarized affidavit on a Rs 100/- Non-Judicial Stamp Paper representing that the offered price is the 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.	Yes	234	Yes	




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TENDER NO. BMSIC/DRUGS/19-09

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

Company Name:- Cadila Pharmaceutical Ltd. Industrial Growth Centre, J & K SIDCO, Samba - 184121 (J & K)

Total Number of Pages Submitted in bid documents:- 241

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

Sl. No	NIT Sl. No	Name of the Quoted Drug		Pharmacopoeial Specification/ Strength		Pack Size		Dosage Form		Approval Details			Last 3yrs market standing certificate/ New Drug
		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	8	Ethambutol	Ethambutol Tablet IP 400 mg	400 mg	Each uncoated Tablet contains: Ethambutol Hydrochloride IP 400 mg (page no. 71)	10x10	Not mentioned	Tablet	Tablet	20.04.2007	28.07.2021	Generic	Yes (Page No. 80)
2	9	Ethambutol	Ethambutol Tablet IP 800 mg	800 mg	Each uncoated Tablet contains: Ethambutol Hydrochloride IP 800 mg	10x10	Not mentioned	Tablet	Tablet	20.04.2007	28.07.2021	Generic	

Rajesh K

Sams
9/11/18

Smt

TENDER NO. BMSIC/DRUGS/19-09

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

Company Name:- Cadila Pharmaceutical Ltd. Industrial Growth Centre, J & K SIDCO, Samba - 184121 (J & K)

Total Number of Pages Submitted in bid documents:- 241

Sheet for verification of licence details

	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	8	Ethambutol 400 mg	25	JK/01/06-07/110	-	-	29.07.06	28.07.2021	10.01.2018	13.12.2019
2	9	Ethambutol 800 mg	25	JK/01/06-07/110	-	-	29.07.06	28.07.2021	10.01.2018	13.12.2019

Note:- Assisted in technical evaluation in reference to letter no.BMSIC/40020/09-2016/5031 dt 26.11.2019 on the basis of documents provided by BMSICL as check list II,III & IV. Provided checklist compiled with due diligence and care.Inspite, some inadvertent discrepencies could have been crept in. Humble request to all concerned to bring to the notice in due time if any discrepencies are observed for rectification.

[Signature]
9/12/19

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9/12/19