TENDER NO. BMSIC/DRUGS/18-04

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

Company Name _RELIANCE LIFE SCIENCES PVT. LTD._____ Total Number of Pages Submitted in bid documents__1 TO 185_____

Sl. No	SBD clause serial No	Descrption 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	"01"	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 follows:- Upto 5 drugs- Rs1,00,000/- (One Lakh only), for 6 to 10 drugs- Rs 2,00,000/- (Two Lakh only), for 11 to 20 drugs- Rs 3,00,000/-(Three Lakh only) and for More than 20 drugs Rs 5,00,000/- (Five Lakh only)	yes	"89-85"	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	"68"-"24"	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	"144"	yes	
5	3.(e)	**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. **Format to be used.**	yes	"16"-"15"	yes	
6	3.(l)	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	"124"&"118" &"112"	yes	

Archana Singh
11/7/2018.

7	3.(m)	Copy of **Income Tax Return for any three of last four Consecutive Assessment years** should be submitted **(self-attested)**.	yes	"75"&"74"& "73"	yes	
8	3.(p)	**Copy of PAN Card of the bidder company should be submitted (self-attested).**	yes	"17"	yes	
9	3.(q)	**Copy of certificate of valid Sales Tax/VAT and GST registration of the bidder company should be submitted (self-attested).**	yes	"81"	yes	

Archanasingh
14/7/2018

TENDER NO. BMSIC/DRUGS/18-04						
Check list - II: Evaluation sheet to be used for preliminary evaluation of technical bids						
Company Name: Reliance life sciences Pvt Ltd.				**Total Number of Pages Submitted in bid documents: 185**		
Sl. No	**SBD Clause serial No**	**Descrption of the Clause/ Technical Eligibility Criteria as per SBD, as mentioned in Check List**	**Document submission status (Yes/ No)**	**Page Number as numbered at 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**	**144**	**Yes**	
2	**3.(e)**	**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. **Format to be used.**	**Yes**	**14 to 16**	**Yes**	
3	**3.(f)**	Minimum **three years old valid Manufacturing License of the product quoted with latest license renewal certificate**.	**Yes**	**94 104 110**	**Yes**	**Manufacturing Licence in Form 28E of plant-4A, 4B for the licence no.28E-KD/8 is not submitted**
4		**Approved product list as per the license issued for quoted drugs for minimum three years.**	**Yes**	**91 to 110**	**Yes**	
5		**Manufacturing License** along with approved product list must be **valid till the last date of the submission of tender.**	**Plant-I Yes**	**91 to 110**	**Yes**	**Manufacturing Licence in Form 28E of plant-II for licence no.28E-KD/8 is not submitted**
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.	-	-	-	
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).	**Yes**	**69 to 72**	**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.	-	-	-	

Rajeev Kumar Ranjan

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.	-	-	-	
10		FFS (Flow Fill & Seal Process) Technology will be accepted wherever applicable	-	-	-	
		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.	-	-	-	
11	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.	-	-	-	
12	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.	-	-	-	
13	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**	**18 to 20**	**Yes**	
14	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.	**Plant-I** **Yes** **Plant-II**	**134** **(127 to 138)** **(82 to 84)** **138**	**Yes**	

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14/03/2018

15	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**	**5 to 13**	**Yes**	Maximum production capacity certificate for the product at NIT S.N. 102 is not submitted.
16	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**	**21 to 22**	**Yes**	**Whether Blacklisted or not** **Not Blacklisted**
17	3.(o)	**List of item quoted in prescribed format as per Annexure-III duly signed.**	**Yes**	**4**	**Yes**	
18	5.(k)	An Affidavit (with stamp) regarding acceptance of tender conditions to be submitted by the bidding firm as per **Annexure-IV.**	**Yes**	**145 to 146 & 2 to 3**	**Yes**	
19	5.(l)	**Filled check list as per given Annexure-VI to be submitted at the time of uploading the bid.**	**Yes**	**147**	**Yes**	
20	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**	**14 to 16**	**Yes**	
21	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.**	-	-	-	
22	Corrigendum point	**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**	**142 to 143**	**Yes**	

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14/07/2018

TENDER NO. BMSIC/DRUGS/18-04

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

Company name : Reliance Life sciences Pvt ltd. **Total number of pages submitted in bid documents 185**

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	66	Dried Human Anti Haemophilic Fraction(Freeze Dried Human Coagulation Fractor 250 IU)	Dried Factor VIII Fraction IP 250 IU/Lyophilised	Nil 250 IU	IP 250 IU	Vial	Vial	Dry powder for injection	Lyophilised	28 E-KD/8 02.01.2015	1/1/2020	Generic Name	3yrs marketing completed(page no.72)
2	67	Dried Human Anti Haemophilic 500 IU Fraction(Freeze Dried Human Coagulation Factor VIII)	Dried Factor VIII Fraction IP 500 IU/Lyophilised	Nil 500 IU	IP 500 IU	Vial	Vial	Dry powder for injection	Lyophilised	28 E-KD/8 02.01.2015	1/1/2020	Generic Name	3yrs marketing completed(page no.72)
3	80	Erythropoietin injection IP 2000 IU/ml	Erythropoietin injection IP 2000 IU per **0.5 ml**	IP 2,000 IU	IP 2,000 IU	1ml Vial	**0.5ml Prefilled syringe**	Injection	Prefilled syringe	28D-KD/7 11.07.2014	4/28/2019	*Generic Name	3yrs marketing completed(page no.71)
4	81	Erythropoietin injection IP 10000 IU/ml	Erythropoietin injection IP 10000 IU/ml	IP 10,000 IU	IP 10,000 IU	1ml Vial	1 ml Prefilled syringe	Injection	Prefilled syringe	28D-KD/7 11.07.2014	4/28/2019	Generic Name	3yrs marketing completed(page no.71)
5	102	Hepatitis B Immunoglobulin Injection 100 IU/ml	Hepatitis B Immunoglobulin IP(For Intramuscular Administration)	IP 100 IU	IP 100 IU	1ml Vial -	1ml Vial	Injection	Injection	28 E-Plant I 28E-3 05.03.2015	11/1/2019	Generic Name	Not submitted

See – Page No. 102

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TENDER NO. BMSIC/DRUGS/18-04

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

Company name: Reliance life sciences Pvt Ltd. **Total number of pages submitted in bid documents: 185**

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	66	Dried Human Anti Haemophilic Fraction(Freeze Dried Human Coagulation Factor 250 IU)	28E	KD/8(Plant -II)	NA	NA	4/29/2014	4/28/2019	Plant-I 24.05.18 06.03.2017 18.05.18 Plant-II 14.03.17 27.12.17 16.05.18	23.05.19 03.03.19 - 03 17.05.19 - KD7 18.0518 - KD7 26.12.19 - KD8 15.05.19 - KD/8
2	67	Dried Human Anti Haemophilic 500 IU Fraction(Freeze Dried Human Coagulation Fractor VIII)	28E	KD/8(Plant -II)	NA	NA	4/29/2014	4/28/2019	Plant-I 24.05.18 06.03.2017 18.05.18 Plant-II 14.03.17 27.12.17 16.05.18	23.05.19 03.03.19 - 03 17.05.19 - KD7 18.0518 - KD7 26.12.19 - KD8 15.05.19 - KD/8

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3	80	Erythropoietin injection IP 2000 IU/ml	28D	KD/7	NA	NA	4/29/2014	4/28/2019	Plant-I 24.05.18 06.03.2017 18.05.18 Plant-II 14.03.17 27.12.17 16.05.18	 23.05.19 03.03.19 - 03 17.05.19 - KD7 18.0518 - KD7 26.12.19 - KD8 15.05.19 - KD/8
4	81	Erythropoietin injection IP 10000 IU/ml	28D	KD/7	NA	NA	4/29/2014	4/28/2019	Plant-I 24.05.18 06.03.2017 18.05.18 Plant-II 14.03.17 27.12.17 16.05.18	 23.05.19 03.03.19 - 03 17.05.19 - KD7 18.0518 - KD7 26.12.19 - KD8 15.05.19 - KD/8
5	102	Hepatitis B Immunoglobulin Injection 100 IU/ml	28E	28E-Plant -I 28E-3	NA	NA	02.11.14	01.11.19	Plant-I 24.05.18 06.03.2017 18.05.18 Plant-II 14.03.17 27.12.17 16.05.18	 23.05.19 03.03.19 - 03 17.05.19 - KD7 18.0518 - KD7 26.12.19 - KD8 15.05.19 - KD/8

Note: - Assisted in technical evaluation in reference to letter No-559(15), dated 26.6.2018 on the basis of documents provided by BMSICL in checklist-II, III & IV. Provided checklist compiled with due deligence and care. Inspite, some inadvertent discrepancies could have been crept in. Humble request to all concerned to bring to the notice in due time, if any discrepancies are observed, for rectification.

Rajeev Kumar Rautray
14/07/2018