

File No.- BMSIC/40030/1044-2026/2988

दिनांक:-21/07/2026

बिहार चिकित्सा सेवाएं एवं आधारभूत संरचना निगम लि०, बिहार पटना
BMSICL की निविदा संख्या- **BMSIC/DRUGS/26-01** से संबंधित प्राप्त ई-निविदाओं के तकनीकी
मूल्यांकन विवरणी पर साक्ष्य सहित दावा/आपत्ति की माँग संबंधी सूचना

बिहार चिकित्सा सेवाएं एवं आधारभूत संरचना निगम लि०, द्वारा औषधियों की अधिप्राप्ति हेतु ई-निविदा (निविदा संख्या- **BMSIC/DRUGS/26-01**) आमंत्रित की गई थी, जिसके आलोक में निर्धारित समय सीमा के अन्तर्गत कुल 42 (बयालीस) ई-निविदा प्राप्त हुई। उक्त 42 (बयालीस) ई-निविदा के दस्तावेजों के प्राथमिक तकनीकी मूल्यांकन के उपरान्त तैयार की गई तकनीकी मूल्यांकन विवरणी के दावा/आपत्ति के आमंत्रण हेतु BMSICL के **website (www.bmsicl.gov.in)** एवं e-proc 2.0 के **website (www.eproc2.bihar.gov.in)** पर अपलोड की गई है।

निविदा में भाग लेने वाले निर्माता संस्थान अथवा अन्य आपूर्तिकर्ता अपलोड किये गए उपर्युक्त तकनीकी मूल्यांकन प्रपत्र में किसी प्रकार की विसंगति पाये जाने की स्थिति में, साक्ष्य सहित अपना दावा/आपत्ति दिनांक 28/07/2026 के अपराह्न 05:00 बजे तक केवल e-proc 2.0 के **website (www.eproc2.bihar.gov.in)** के माध्यम से कर सकते हैं।

e-proc 2.0 के **website (www.eproc2.bihar.gov.in)** के अलावा अन्य किसी माध्यम से प्राप्त किसी प्रकार के दावा/आपत्ति पर कोई विचार नहीं किया जायेगा।

अनुलग्नक- यथोक्त


महाप्रबंधक (अधिप्राप्ति)

ज्ञापांक:- BMSIC/40030/1044-2026/2988

दिनांक:-21/07/2026

प्रतिलिपि :- सहायक प्रबंधक (सिस्टम) एवं प्रबंधक (औषधि) को सूचनार्थ प्रेषित। निदेश है कि सूचना का अनुलग्नक सहित अविलंब BMSICL के **website (www.bmsicl.gov.in)** एवं e-proc2 (**www.eproc2.bihar.gov.in**) पर अपलोड करना सुनिश्चित करें।


महाप्रबंधक (अधिप्राप्ति)



TENDER NO. BMSIC/DRUGS/26-01

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

Company Name :-Modern Laboratories. Total Number of Pages Submitted in bid documents: - 1 to 140.

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/- + GST @18%, Total Rs 11,800 shall be submitted online (On E-Proc-2). This fee is payable only once for one tender irrespective of items contained therein.	Yes	28	Yes	-
2	3.(b)	EMD shall be submitted ONLY in form of Bank Guarantee (Offline mode), drawn in favor of Managing Director, Bihar Medical Services and Infrastructure Corporation Limited from any Scheduled/Nationalized bank payable at Patna, as mentioned in Clause 2, as per following table:- S.N.No. of drugs quoted EMD Amount 1 Upto 5 drugs Rs 1,00,000/- (One Lakh only) 2 For 6 to 10 Drugs Rs 2,00,000/- (Two Lakh only) 3 For 11 to 15 Drugs Rs 3,00,000/- (Three Lakh only) 4 For 16 to 20 Drugs Rs 4,00,000/- (Four Lakh only) 5 More than 20 Drugs Rs 5,00,000/- (Five Lakh only)	Yes	44-49	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	13-23	Yes	-
4	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 (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. It's mandatory for the firm to provide a document for the Unique Document Identification Number (UDIN) to verify the documents.	Yes	36-42	Yes	-
5	3.(m)	Copy of Income Tax Return for any three of last four Consecutive Assessment years should be submitted (self-attested).	Yes	33-35	Yes	-
6	3.(p)	Copy of PAN Card of the bidder company should be submitted (self-attested).	Yes	29	Yes	-
7	3.(q)	Copy of certificate of valid GST registration of the bidder company should be submitted (self-attested).	Yes	30-32	Yes	-

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TENDER NO. BMSIC/DRUGS/26-01

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

Company Name:- M/s Modern Laboratories. Total Number of Pages Submitted in bid documents:- 140

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	23 53-54	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	8-10	Yes	
3	3.(f)	Minimum three years old valid Manufacturing License of the quoted product.	Yes	83-87	Yes	
4		Manufacturing License along with approved product list must be valid till the last date of the submission of tender.	Yes	83-87 88-104	Yes	
5		Market standing certificate & Manufacturing certificate issued by the Licensing Authority as a Manufacturer for each drug quoted for the last 3 years. The Market Standing Certificate must have been issued within preceding one year from the last date of submission of tender (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. Explanation- In case of Importers Permission in Form 45 from DCGI is required as Per Drugs & Cosmetics Act 1940 & Rules 1945. For drugs with a patent that expired less than 2 years ago from the last date of bid submission: Non-patent holding bidders should submit a valid Market Standing Certificate and valid drug license, issued by the State Drug Licensing Authority in support of the claim. Bidders will be eligible only if one batch of new drug has been successfully manufactured/ imported for commercial use at the time of bidding. Bidders should submit document regarding patent expired less than 2 years ago from the last date of bid submission.	Yes	106-128	Yes	
6		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.	Yes	59-81	Yes	

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7		FFS (Flow Fill & Seal Process) Technology will be accepted wherever applicable.	No	No	No	FFS technology related document required...
8		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	129-134	Yes	
9		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	82-104	Yes	
10	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 is 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	
11	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. The Market Standing Certificate must have been issued within preceding one year from the last date of submission of tender. Self-attested copies are to be submitted.	Yes	106-128	Yes	
12	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 (wherever applicable). It should be not more than one year old. Self-attested copies are to be submitted.	Yes	11	Yes	
13	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, (wherever applicable). The GMP certificate must not be older than one year from the date of publication of tender. Self-attested copies are to be submitted. Explanation- Generally the GMP Certificate is issued for one-year validity. Hence the provision that it should not be older than one year from the date of publication of tender implies mutatis mutandis that the GMP certificate should remain valid till the date of submission of tender.	Yes	24-25	Yes	
14	3.(k)	Maximum Production Capacity Certificate (Section wise) issued by concerned Licensing Authority from Drugs Control Department highlighting the quoted product section (wherever applicable). 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	12	Yes	

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15	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 /Debarred 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	4-5 26-27	Yes	
16	3. (o)	List of items quoted in prescribed format as per Annexure-III duly signed.	Yes	2-3	Yes	
17	5.(k)	An Affidavit (with stamp) regarding acceptance of tender conditions to be submitted by the bidding firm as per Annexure-IV .	Yes	43	Yes	
18	5.(l)	Filled check list as per given Annexure-VI or VI-A to be submitted at the time of uploading the bid.	Yes	50-52	Yes	
19	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	8-10	Yes	
20	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	
21	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 provision of "Drugs Price Control Order" (wherever applicable) and the quoted rate should be at least 20% less than its MRP. In extraordinary cases the Managing Director, BMSICL 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 Rate 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-VIII .	Yes	57	Yes	

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

Company Name:- M/s Modern Laboratories. Total Number of Pages Submitted in bid documents:- 140

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		Dosage Form		Pack Size		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	Approval From	Approved Upto	Approved in Brand /Generic Name	
1	1	Amikacin	Amikacin Sulphate Injection IP	250 mg	Each 2 ml contains: Amikacin sulphate IP Eq. to Amikacin 250 mg	Injection	Injection	2 ml Vial	Not mentioned	01.04.2022	31.12.2026	Generic	Yes (page no. 110)
2	4	Amoxycillin	Amoxycillin Oral Suspension IP Dry Syrups	125 mg/5 ml	Each 5 ml Reconstituted Suspension Contains: Amoxycillin Trihydrate IP Eq. to Amoxycillin 125 mg	Oral liquid	Oral Suspension	30ml Bottle	Not mentioned	01.04.2022	31.12.2026	Generic	Yes (page no. 119)
3	5	Atropine Sulphate	Atropine Eye Drops BP	1% w/v	Each ml contains: Atropine IP 1% w/v	Eye drops	Eye drops	5 ml bottle	Not mentioned	01.04.2022	31.12.2026	Generic	Yes (page no. 118)
4	7	Calcium carbonate + Vitamin D3	Calcium and Vitamin D3 Tablet IP	500mg Elemental Ca + 250 IU D3	Each uncoated Tablet contains: Calcium carbonate IP 1250 mg Eq. to Elemental Calcium 500 mg Vitamin D3 IP 250 I.U	Tablet	Tablet	10x10	Not mentioned	01.04.2022	31.12.2026	Generic	Yes (page no. 127)
5	8	Chloroquine Phosphate IP	Chloroquine Phosphate IP	250 mg (250 mg of Chloroquine Phosphate is approximately equivalent to 155 mg of Chloroquine)	Each film coated tablet contains: Chloroquine Phosphate IP 250 mg Eq. to Chloroquine base 155 mg.	Tablet	Tablet	10 x 10	Not mentioned	01.04.2022	31.12.2026	Generic	Yes (page no. 112)

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6	10	Dicyclomine HCL	Dicyclomine Oral solution IP	10mg/ml	Each 5 ml contains: Dicyclomine Hydrochloride IP 10 mg	Drop	Oral liquid Syrups	15 ml Bottle (Palatable with dropper and plastic Container/ glass bottle as per I.P	Not mentioned	01.04.2022	31.12.2026	Generic	Yes (page no. 123)
7	11	Diethylcarbamazine	Diethylcarbamazine Tablet IP	100 mg	Each uncoated tablet contains: Diethylcarbamazine Citrate IP 100 mg	Tablet	Tablet	10 x 10	Not mentioned	01.04.2022	31.12.2026	Generic	Yes (page no. 113)
8	12	Folic acid	Folic acid Tablet IP	5 mg	Each uncoated tablet contains: Folic acid IP 5 mg	Tablet	Tablet	10x10	Not mentioned	01.04.2022	31.12.2026	Generic	Yes (page no. 114)
9	14	Ibuprofen	Ibuprofen Tablet IP	200 mg	Each film coated tablet contains: Ibuprofen IP 200 mg	Tablet	Tablet	10x10's	Not mentioned	01.04.2022	31.12.2026	Generic	Yes (page no. 123)
10	15	Iron sucrose	Iron sucrose Injection USP	20 mg/ml	Each 5 ml contains: Iron (as Iron Sucrose) IH 100 mg	Injection	Injection	5 ml Ampoule	Not mentioned	01.04.2022	31.12.2026	Generic	Yes (page no. 106)
11	17	Lignocaine	Lignocaine Injection IP	2% w/v	Each ml contains: Lignocaine Hydrochloride IP 20 mg	Injection	Injection	30 ml Vial	Not mentioned	01.04.2022	31.12.2026	Generic	Yes (page no. 106)

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12	20	Ondansetron	Ondansetron Tablet IP	4 mg	Each uncoated tablet contains Ondansetron Hydrochloride IP Eq. to Ondansetron 4 mg	Tablet	Tablet	10x10	Not mentioned	01.04.2022	31.12.2026	Generic	Yes (page no. 116)
13	21	Pantoprazole	Pantoprazole Tablet IP	40 mg	Each Gastro Resistant Enteric coated tablet contains: Pantoprazole sodium IP Eq. to Pantoprazole 40 mg	Tablet	Tablet	10x10	Not mentioned	01.04.2022	31.12.2026	Generic	Yes (page no. 124)
14	24	Piperacillin (A) + Tazobactam (B)	Piperacillin and Tazobactam Injection IP	1 g (A) + 125 mg (B)	Each vial contains: Piperacillin sodium IP Eq. to Piperacillin 1 gm Tazobactam sodium IP Eq. to Tazobactam 125 mg	Powder for Injection	Powder for Injection	Vial	Not mentioned	01.04.2022	31.12.2026	Generic	Yes (page no. 121)
15	25	Prednisolone	Prednisolone Tablet IP	20 mg	Each uncoated tablet contains: Prednisolone IP 20 mg	Tablet	Tablet	10x10	Not mentioned	01.04.2022	31.12.2026	Generic	Yes (page no. 123)
16	28	Sodium bicarbonate	Sodium bicarbonate Injection IP	7.5% w/v	Each ml contains: Sodium bicarbonate IP 7.5% w/v	Injection	Injection	Ampoule	Not mentioned	01.04.2022	31.12.2026	Generic	Yes (page no. 122)

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

Company Name:- M/s Modern Laboratories. Total Number of Pages Submitted in bid documents:- 140

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	1	Amikacin	25 & 28	25/31/89 & 28/17/79	NA	NA	Form- 25 08.08.1989 & Form- 28 09.05.1979	31.12.2026	25.08.2025	24.08.2028
2	4	Amoxycillin	25 & 28	25/31/89 & 28/17/79	NA	NA	Form- 25 08.08.1989 & Form- 28 09.05.1979	31.12.2026	25.08.2025	24.08.2028
3	5	Atropine Sulphate	25 & 28	25/31/89 & 28/17/79	NA	NA	Form- 25 08.08.1989 & Form- 28 09.05.1979	31.12.2026	25.08.2025	24.08.2028
4	7	Calcium carbonate + Vitamin D3	25 & 28	25/31/89 & 28/17/79	NA	NA	Form- 25 08.08.1989 & Form- 28 09.05.1979	31.12.2026	25.08.2025	24.08.2028
5	8	Chloroquine Phosphate IP	25 & 28	25/31/89 & 28/17/79	NA	NA	Form- 25 08.08.1989 & Form- 28 09.05.1979	31.12.2026	25.08.2025	24.08.2028
6	10	Dicyclomine HCL	25 & 28	25/31/89 & 28/17/79	NA	NA	Form- 25 08.08.1989 & Form- 28 09.05.1979	31.12.2026	25.08.2025	24.08.2028
7	11	Diethylcarbamazine	25 & 28	25/31/89 & 28/17/79	NA	NA	Form- 25 08.08.1989 & Form- 28 09.05.1979	31.12.2026	25.08.2025	24.08.2028

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8	12	Folic acid	25 & 28	25/31/89 & 28/17/79	NA	NA	Form- 25 08.08.1989 & Form- 28 09.05.1979	31.12.2026	25.08.2025	24.08.2028
9	14	Ibuprofen	25 & 28	25/31/89 & 28/17/79	NA	NA	Form- 25 08.08.1989 & Form- 28 09.05.1979	31.12.2026	25.08.2025	24.08.2028
10	15	Iron sucrose	25 & 28	25/31/89 & 28/17/79	NA	NA	Form- 25 08.08.1989 & Form- 28 09.05.1979	31.12.2026	25.08.2025	24.08.2028
11	17	Lignocaine	25 & 28	25/31/89 & 28/17/79	NA	NA	Form- 25 08.08.1989 & Form- 28 09.05.1979	31.12.2026	25.08.2025	24.08.2028
12	20	Ondansetron	25 & 28	25/31/89 & 28/17/79	NA	NA	Form- 25 08.08.1989 & Form- 28 09.05.1979	31.12.2026	25.08.2025	24.08.2028
13	21	Pantoprazole	25 & 28	25/31/89 & 28/17/79	NA	NA	Form- 25 08.08.1989 & Form- 28 09.05.1979	31.12.2026	25.08.2025	24.08.2028
14	24	Piperacillin (A) + Tazobactam (B)	25 & 28	25/31/89 & 28/17/79	NA	NA	Form- 25 08.08.1989 & Form- 28 09.05.1979	31.12.2026	25.08.2025	24.08.2028
15	25	Prednisolone	25 & 28	25/31/89 & 28/17/79	NA	NA	Form- 25 08.08.1989 & Form- 28 09.05.1979	31.12.2026	25.08.2025	24.08.2028
16	28	Sodium bicarbonate	25 & 28	25/31/89 & 28/17/79	NA	NA	Form- 25 08.08.1989 & Form- 28 09.05.1979	31.12.2026	25.08.2025	24.08.2028

Note:- Assisted in technical evaluation in reference to letter no. BMSIC/40030/1044-2026/71870 dt 11.06.2026 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 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.

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