

File No.- BMSIC/40030/661-2025/3270

दिनांक:- 02/07/2025

बिहार चिकित्सा सेवाएं एवं आधारभूत संरचना निगम लि०, बिहार पटना

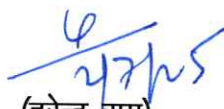
BMSICL की निविदा संख्या-BMSIC/MEDICAL DEVICES/CONSUMABLES/25-04 से संबंधित प्राप्त ई-निविदाओं के तकनीकी मूल्यांकन विवरणी पर साक्ष्य सहित दावा/आपत्ति की माँग संबंधी सूचना

बिहार चिकित्सा सेवाएं एवं आधारभूत संरचना निगम लि०, द्वारा Medical Devices/Consumables की अधिप्राप्ति हेतु ई-निविदा (निविदा संख्या-BMSIC/MEDICAL DEVICES/CONSUMABLES/25-04) आमंत्रित की गई थी, जिसके आलोक में निर्धारित समय सीमा के अन्तर्गत कुल 15 (पन्द्रह) ई-निविदा प्राप्त हुई। उक्त 15 (पन्द्रह) 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 09/07/2025 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


(हरेन्द्र राम)
महाप्रबंधक (अधिप्राप्ति)

ज्ञापांक:- BMSIC/40030/661-2025/3270

दिनांक:- 02/07/2025

प्रतिलिपि :- 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/MEDICAL DEVICES/CONSUMABLE/25-04

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

Company Name:-Biotrol Laboratories Private Limited. Total Number of Pages Submitted in bld documents:- 1 to 472

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	84	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 MEDICAL DEVICES/CONSUMABLE quoted EMD Amount 1 Upto 5 MEDICAL DEVICES/CONSUMABLE Rs 50,000/- (Fifty thousand only) 2 6 to 10 MEDICAL DEVICES/CONSUMABLE Rs 1,00,000 (One Lakhs only) 3 11 to 20 MEDICAL DEVICES/CONSUMABLE Rs 1,50,000 (One Lakhs fifty thousand only) 4 More than 20 MEDICAL DEVICES/CONSUMABLE Rs 2,50,000 (Two Lakhs fifty thousand only)	Yes	90-95	Yes	As per Bid document clause 3(b) BG Submitted for EMD must be Schedule/Nationalized Bank Payable at Patna But firm has not submitted the same.
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	01-24	Yes	—
4	3.(k)	Copies of the Audited Balance Sheet and Profit and Loss statement showing details of their annual average turnover not less than 5 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. It's mandatory for the firm to provide a document for the Unique Document Identification Number (UDIN) to verify the documents.	Yes	335-376	No	Profit & Loss Statement Submitted for F.Y-2022-2023 Only.
5	3.(l)	Copy of Income Tax Return for any three of last four consecutive Assessment years should be submitted (Self Attested).	Yes	87-89	Yes	—
6	3.(o)	Copy of PAN Card of the bidder company should be submitted (self-attested).	Yes	80	Yes	—
7	3.(p)	Copy of certificate of valid GST registration of the bidder company should be submitted (self-attested).	Yes	81-83	Yes	—

Anhan Singh

[Signature]

TENDER NO. BMSIC/MEDICAL DEVICES/CONSUMABLE/25-04

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

Company Name: - M/s Biotrol Laboratories Ltd. Total Number of Pages Submitted in bid documents: 472

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	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	217	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	36	Yes	
3	3.(f)	(I) Minimum three years old valid Manufacturing License of the quoted product.	Yes	40-49	No	Specification as per NIT are not mentioned for NIT sr. no. 1, 2, 3, 7 & 13.
4		In case of manufacturer, the bidder firm must have a valid manufacturing license or duly acknowledged renewal application with old license issued by the state licensing authority/Central licensing approving authority under Medical Devices Rules 2017 where ever applicable.				
5		Approved product list as per the license issued for quoted MEDICAL DEVICES/CONSUMABLE as Per Medical Devices Rules 2017.				
6		Manufacturing License along with approved product list must be valid till the last date of the submission of tender.				
7		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	50-51	Yes	
8		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	40-49	Yes	

[Signature]
28/06/25

[Signature]
28/06/25

[Signature]
27/6/25

[Signature]
30/06/25

9		(II) In case of Importer, the bidder (importer) firm must have a valid import License with product registration certificate issued by the Drugs controller General of India as per Medical Device Rule 2017. 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	
10		(III) In case of Non-drug item(s) where neither the D & C Act 1945 nor the Medical Device Rule 2017 is applicable the bidder must have a manufacturing license/import export certificate (IEC) with an undertaking/Self declaration in his letter pas as per Annexure-VII that the item(s) quoted by the bidder is/are non-drug item(s) i.e. neither covered under D & C Act nor Under Medical Device Rule 2017.	Yes	218-219	Yes	
11	3.(g)	Bidder must have Market Standing Certificate of minimum three years issued by the concerned Licensing Authority from Drugs Control Department/Concerned Government 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	52-59	No	(i) MSC for NIT sl. No. 1, 2, 3, 7 & 13 Submitted for only one year. (ii) Not submitted for three year.
12	3.(h)	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	68-79	Yes	
13	3.(i)	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	61-62	Yes	ISO 13485:2016 Certificate submitted (Page no.61 & 62)
14	3.(j)	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	52-59 63-67	No	Maximum Production Capacity Certificate not issued by concerned department.

[Signature]
22/06/25

[Signature]
26/06/25

[Signature]

[Signature]
23/06/25

15	3.(m)	The tenderer should give an affidavit 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	60	Yes	
16	3.(n)	List of items quoted in prescribed format as per Annexure-III duly signed.	Yes	37-39	Yes	
17	5.(k)	Affidavit declaration regarding acceptance of tender conditions to be submitted by the bidding firm as per Annexure-IV .	Yes	472	Yes	
18	5.(l)	Filled check list as per given Annexure-VI to be submitted at the time of uploading the bid.	Yes	214-215	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	36	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)	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-X .	Yes	219-220	Yes	
22	—	Performance Statement- Annexure- IX	Yes	98-213	Yes	
23	—	AFFIDAVIT FOR NON DRUG ITEM(S) ANNEXURE-VII	Yes	218-219	Yes	

[Signature]
28/06/25

[Signature]
26/06/25

[Signature]

[Signature]
30/06/25

TENDER NO. BMSIC/MEDICAL DEVICES/CONSUMABLE/25-04										
Check list - III: Evaluation sheet to be used for preliminary evaluation of technical bids										
Company Name: - M/s Biotrol Laboratories Ltd. Total Number of Pages Submitted in bid documents: 472										
Sheet to be used for verification of product approval and market standing										
Sl. No	NIT Sl. No	Name of the Quoted MEDICAL DEVICES/CONSUMABLE		Specification		Product category	Approval Details			Last 3yrs market standing certificate/ New Drug
		As per NIT	As per Approval	As per NIT	As per Approval		(Medical Device/Non Drug)	Product Approval	Approved Upto	Approved in Brand /Generic Name
1	1	Human Chorionic gonadotropin (HCG) Urine Test Kit for Pregnancy	HCG Rapid Test	Annexure-A Technical Specification of Human Chorionic gonadotropin (HCG) Urine Test Kit for Pregnancy 1. Should be rapid test for the qualitative detection of human chorionic gonadotropin in urine. 2. The test should have anti-hCG particles and anti-hCG coated on the membrane. 3. The HCG Pregnancy Rapid Test Cassette should have sensitivity of 25mIU/mL and should be capable of detecting pregnancy as early as 1 day after the first missed menses. 5. Detection Type: Qualitative 6. Testing Principle: Single Step, self-performing sandwiched immunoassay using colloidal gold and anti-hCG antibodies in lateral flow immunochromatography format. 8. Test result should be visible maximum within 5 minutes or lesser. 9. Specimen required for testing: Urine 10. Specificity: 100% (no cross reactivity with other gonadotropin hormones like LH, FSH etc) 11. Every test pack should have moisture indicating silica gel pouch. 12. Disposable dropper for urine specimen addition should be provided with each test pouch. 13. The manufacturer should have CDSCO Manufacturing License for the product. 14. The Manufacturer should have ISO 13485:2016 certification. 15. Stable at 2-40°C, allowing ease of storage and transport. 16. Shelf life in months from the date of manufacture: 24-36 month 17. Pack size: 25/50 Tests/Kit.	1. Yes 2. Not mentioned 3. Not mentioned 4. Not mentioned 5. Yes 6. Not mentioned 7. Not mentioned 8. Not mentioned 9. Not mentioned 10. Not mentioned 11. Not mentioned 12. Not mentioned 13. Yes 14. Yes 15. Not mentioned 16. Not mentioned 17. Not mentioned	Medical Device	17.11.2021 (page no. 44)	16.11.2026	Generic	(i) Submitted for only one year (ii) Not submitted for 3 year
2	2	Multiparameter urin strip (Urine test for Ph, Specific Gravity, Leucocyte esterase glucose, bilirubin, Ketone, Protin, Nitrate)	Urine Reagent strip for Leukocytes, Glucose, Ketone (Acetoacetic acid), Bilirubin, Blood, Specific Gravity, Protein, Urobilinogen, Nitrite and pH (10 Parameters).	Technical Specification of Multiparameter urin strip (Urine test for Ph, Specific Gravity, Leucocyte esterase glucose, bilirubin, Ketone, Protin, Nitrate) 1. Should be able to semi-quantitatively/qualitatively detect Glucose, Bilirubin, Ketone, Specific Gravity, Blood, pH, Protein, Urobilinogen, Nitrite and Leukocytes in urine. 2. The Strips should be made exclusively from medical-grade plastic, ensuring durability, uniform reaction surfaces, and resistance to warping or contamination. Paper based strips not eligible. 3. The urine test strips should be packed along with a drying agent in air tight plastic bottle. 4. Should be stable at room temperature between 15°-30°C. 5. Pack size: 25/50/100 Strips/Kit 6. Each strip should be stable up to 24-36 month from manufacture date when stored properly in a sealed container. 7. The manufacturer should have CDSCO Approved Manufacturing License for the product. 8. The Manufacturer should have ISO 13485:2016 certification.	1. Yes 2. Not mentioned 3. Not mentioned 4. Not mentioned 5. Not mentioned 6. Not mentioned 7. Yes 8. Yes	Medical Device	17.11.2021 (page no. 41)	16.11.2026	Generic	(i) Submitted for only one year (ii) Not submitted for 3 year

[Signature]
28/06/25

[Signature]
26/06/25

[Signature]

[Signature]
30/06/25

3	3	Dengue	Dengue Combo (NS1 Ag-IgG/ IgM) Rapid Test	Technical Specification of Dengue Rapid Card Test for NS1 antigen and IgM and IgG Antibodies 1. Should be a rapid chromatographic immunoassay for the qualitative detection of NS1 Antigen and IgG/IgM Antibodies of Dengue Virus in human serum and plasma. 2. Kit should be able to detect all the 4 serotypes of dengue viruses (DEN-1, DEN-2, DEN-3, and DEN-4) including early detection Day-1 Dengue fever. 3. The test should be able to differentially detect IgG and IgM Antibodies against all 4 serotypes of Dengue virus. 4. Test should use minimal sample volume (5ul) for testing IgG/IgM Dengue, and should not use buffer for testing Dengue NS1. 5. The control dot/band able to detect the presence of human immunoglobulin and not be just a procedural control or meant merely for checking the flow of reagents or integrity of antigens except in lateral flow technology. 6. Test should have no cross reactivity with other Flavivirus group mediated and mosquitoes-borne disease. 7. Sensitivity for Dengue NS1 Ag (%): $\geq 99\%$ 8. Specificity for Dengue NS1 Ag (%): $\geq 98\%$ 9. Sensitivity for Dengue IgG/IgM Antibody (%): $\geq 99\%$ 10. Specificity for Dengue IgG/IgM Antibody (%): $\geq 96\%$ 11. Stable at 2-40°C, allowing ease of storage and transport. 12. The manufacturer should have CDSCO Approved Manufacturing License for the product. 13. The Manufacturer should have ISO 13485:2016 certification. 14. Shelf life in months from the date of manufacture: 24-36 months 15. Pack size: 25/50 Tests/Kit.	1. Yes 2. Not mentioned 3. Not mentioned 4. Not mentioned 5. Not mentioned 6. Not mentioned 7. Not mentioned 8. Not mentioned 9. Not mentioned 10. Not mentioned 11. Not mentioned 12. Yes 13. Yes 14. 24 month 15. Not mentioned	Medical Device	05.05.2022 (page no. 47)	04.05.2027	Generic	(i) Submitted for only one year (ii) Not submitted for 3 year
4	7	Filariasis (endemic area only) - FST	Filariasis IgG/ IgM Rapid Test	Technical Specification of Filariasis (endemic area only) – FST Rapid Kit 1. Should be a rapid lateral flow immunoassay for the simultaneous detection and differentiation of IgG and IgM anti-lymphatic filarial parasites (W. Bancrofti and B. Malayi) in human serum and plasma. 2. The test kit should be stable for use and storage at room temperature. 3. Test Results should be read in 15 minutes. Positive results should be visible in as short as 1 minute. 4. Sensitivity: $\geq 99\%$ 5. Specificity: $\geq 96\%$ 6. The manufacturer should have CDSCO Approved Manufacturing License for the product. 7. Stable at 2-40°C, allowing ease of storage and transport. 8. The Manufacturer should have ISO 13485:2016 certification. 9. Shelf life in months from the date of manufacture: 24-36 months 10. Pack size: 25/50 Tests per Kit	1. Yes 2. Not mentioned 3. Not mentioned 4. Not mentioned 5. Not mentioned 6. Yes 7. Not mentioned 8. Yes 9. 24 month 10. Not mentioned	Medical Device	17.11.2021 (page no. 43)	16.11.2026	Generic	(i) Submitted for only one year (ii) Not submitted for 3 year
5	13	Malaria Test	Malaria Pf/ Pv Ag Rapid Test	Technical Specification of Malaria Test Rapid Card Test 1. Rapid, visual antigen-based test for detecting infection with P. falciparum & P. vivax malaria parasite in whole blood. 2. Sensitivity $\geq 98-99\%$ 3. Specificity $\geq 96\%$ or more 4. Easy result interpretation. 5. Minimum shelf life of 24-36 months at 4-30°C. 6. prominent colour bands 7. Results within 15-20 minutes.	1. Yes 2. Not mentioned 3. Not mentioned 4. Not mentioned 5. 24 month 6. Not mentioned 7. Not mentioned	Medical Device	05.05.2022 (page no. 48)	04.05.2027	Generic	(i) Submitted for only one year (ii) Not submitted for 3 year

[Signature]
28/06/25

[Signature]
26/06/25

[Signature]

[Signature]
30/06/25

TENDER NO. BMSIC/MEDICAL DEVICES/CONSUMABLE/25-04

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

Company Name: - M/s Biotrol Laboratories Ltd. Total Number of Pages Submitted in bid documents: 472

Sheet for verification of licence details

Sl. No	NIT Sl. No	Name of the MEDICAL DEVICES/CONSUMABLE as per NIT	Manufacturer		Importer		Validity		GMP/ WHO GMP/ COPP (Import)/BIS/ISI/QMS	
			Forms Number	Manufacturing license Number	Forms Number	Import license Number	From	To	From	To
1	1	Human Chorionic gonadotropin (HCG) Urine Test Kit for Pregnancy	MD- 5	MFG/IVD/2021/000109	NA	NA	17.11.2021	16.11.2026	ISO 13485: 2016 21.02.2023	20.02.2026 (page no. 61)
2	2	Multiparameter urin strip (Urine test for Ph, Specific Gravity, Leucocyte esterse glucose, bilirubin, Ketone, Protin, Nitrate)	MD- 9	MFG/IVD/2021/000109	NA	NA	17.11.2021	16.11.2026	ISO 13485: 2016 21.02.2023	20.02.2026 (page no. 61)
3	3	Dengue	MD- 9	MFG/IVD/2022/000056	NA	NA	05.05.2022 (page no. 49)	06.05.2027	ISO 13485: 2016 21.02.2023	20.02.2026 (page no. 61)
4	7	Filariasis (endemic area only) - FST	MD- 5	MFG/IVD/2021/000109	NA	NA	17.11.2021	16.11.2026	ISO 13485: 2016 21.02.2023	20.02.2026 (page no. 61)
5	13	Malaria Test	MD- 9	MFG/IVD/2022/000056	NA	NA	05.05.2022 (page no. 49)	06.05.2027	ISO 13485: 2016 21.02.2023	20.02.2026 (page no. 61)

Note:- Assisted in technical evaluation in reference to letter no. BMSIC/40030/661-2025/2932 dt 25.06.2025 on the basis of documents provided by BMSICL as check list II, III & IV. Provided checklist compiled with due deligence and care.Inspite, some inadvertent discreprencies could have been crept in. Humble request to all concerned to bring to the notice in due time if any discreprencies are observed for rectification.

Shalini
28/06/25

BMM
26/06/25

Sam

Spand
30/06/25