

File No.- BMSIC/40030/878-2025/ 9768

दिनांक:- 11 / 12 / 2025

बिहार चिकित्सा सेवाएं एवं आधारभूत संरचना निगम लि०, बिहार पटना

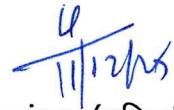
BMSICL की निविदा संख्या-BMSIC/MEDICAL DEVICES/CONSUMABLES/25-12 से संबंधित प्राप्त ई-निविदाओं के तकनीकी मूल्यांकन विवरणी पर साक्ष्य सहित दावा/आपत्ति की माँग संबंधी सूचना

बिहार चिकित्सा सेवाएं एवं आधारभूत संरचना निगम लि०, द्वारा Medical Devices/Consumables की अधिप्राप्ति हेतु ई-निविदा (निविदा संख्या-BMSIC/MEDICAL DEVICES/CONSUMABLES/25-12) आमंत्रित की गई थी, जिसके आलोक में निर्धारित समय सीमा के अन्तर्गत कुल 06 (छः) ई-निविदा प्राप्त हुई। उक्त 06 (छः) 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 18/12/2025 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


महाप्रबंधक (अधिप्राप्ति)

ज्ञापांक:- BMSIC/40030/878-2025/ 9768

दिनांक:- 11 / 12 / 2025

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/MEDICAL DEVICES/CONSUMABLE/25-12

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

Company Name: Genetix Biotech Asia Ltd. Total Number of Pages Submitted in bid documents: 01 to 344.

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	334	Yes	
2	3.(b)	EMD shall be submitted ONLY in form of Bank Guarantee (Offline mode), drawn in favour 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	277-281	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	282-310	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	38-105	Yes	
5	3.(l)	Copy of Income Tax Return for any three of last four consecutive Assessment years should be submitted (Self Attested).	Yes	165-168	Yes	
6	3.(o)	Copy of PAN Card of the bidder company should be submitted (self-attested).	Yes	156	Yes	
7	3.(p)	Copy of certificate of valid GST registration of the bidder company should be submitted (self-attested).	Yes	153-155	Yes	

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TENDER NO. BMSIC/MEDICAL DEVICES/CONSUMABLE/25-12

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

Company Name: - M/s Genetix Biotech Asia Ltd. Total Number of Pages Submitted in bid documents:- 344

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	106	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	107	No	Notarized power of attorney not submitted.
3	3.(f)	Bidders must have:- (I) Valid Manufacturing License of the quoted product.	Yes	110 & 175-177	No	(i) Not submitted approval for self sample collection kit & HPV Detection kit. (ii) Retention not submitted for DNA extraction kit.
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	111-112	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	110 & 175-177	No	(i) Not submitted approval for self sample collection kit & HPV Detection kit. (ii) Retention not submitted for DNA extraction kit.

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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. 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.	NA	NA	NA	
11	3.(g)	Bidder must have Valid Market Standing Certificate 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	118-120	No	Not submitted for self sample collection kit & HPV detection kit.
12	3.(h)	Non-Conviction Certificate (NCC) issued by the concerned Licensing Authority from Drugs Control Administration of the state should be submitted (wherever applicable). It should be not more than one year old. Self-attested copies are to be submitted.	Yes	121-122	No	Not submitted for Manufacturing unit.
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	317	No	ISO 13485: 2016 submitted valid upto 01.11.2025
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	123	No	Not issued by concerned authority.
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	108-109	Yes	

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16	3.(n)	List of items quoted in prescribed format as per Annexure-III duly signed.	Yes	37	Yes	
17	5.(k)	Affidavit declaration regarding acceptance of tender conditions to be submitted by the bidding firm as per Annexure-IV .	Yes	148-149	Yes	
18	5.(l)	Filled check list as per given Annexure-VI to be submitted at the time of uploading the bid.	Yes	124-125	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	107	No	Notarized power of attorney not submitted.
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	126-127	No	Non Judicial stamp paper less than 100/-
22	—	Performance Statement- Annexure- IX	Yes	131-146	Yes	
23	—	AFFIDAVIT FOR NON DRUG ITEM(S) ANNEXURE-VII	NA	NA	NA	

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TENDER NO. BMSIC/MEDICAL DEVICES/CONSUMABLE/25-12										
Check list - III: Evaluation sheet to be used for preliminary evaluation of technical bids										
Company Name: - M/s Genetix Biotech Asia Ltd. Total Number of Pages Submitted in bid documents:- 344										
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			Valid Market Standing Certificate issued by the concerned Licensing Authority from Drugs Control Department/Concerned
		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	6	Pap smear (DNA extraction kit)	Not submitted	Technical Specification of Rapid Antigen detection test for Bacterial meningitis (Meningococci) :- 1. The kit should contain reagents and controls 2. Result $\leq$ 5 minutes 3. Shelf Life 24-36 Months 4. Certification BIS/CE/US FDA 5. Sensitivity $\geq$ 90% 6. Specificity $\geq$ 95% Technical Specification of Pap Smear (DNA extraction Kit for diagnosis of cervix cancer) :- ☐ Self-Sample Collection Kit for community level • Sterile device for self-collection, validated for home use, Kit should be easy to use for self-collection by women. • Comes with a sterile swab and a viral inactivation & DNA stabilizing medium. • Sample should remain stable for 14 days at room temperature. • Shelf life of the collection kit: Minimum 24 months. • Clinically validated on minimum 2,000 samples. • Pack size -50 sample in one Box. 2. RTPCR lab at the district hospital level for confirmatory test	Not submitted		Not submitted	Not submitted	Not submitted	Not submitted
			Nucleic Acid Extraction kit	2.1 DNA Extraction Kit • High nucleic acid binding and recovery. • Compatible with various sample media, including HPV preservation medium. • Quick extraction protocol, completed within 30 minutes. • Suitable for low DNA input and degraded samples. 2.2 HPV Detection Kit (RT-PCR) • Based on hydrolysis fluorescence probe principle (e.g., TaqMan). • Detects 14 high-risk HPV genotypes, with separate detection for HPV 16 & 18. • Multiplexed, two-tube/single-tube format for each test. • Includes internal control (Beta-globin) in a separate fluorescent dye. • Sensitivity: Detects as low as 1000 copies/ • Shelf life: Minimum 12 months. • Reaction setup must be user-friendly, allowing reaction preparation in $\leq$ 3- 4steps. • Simple protocol: 3 steps, total test time $\leq$ 2 hours.	Nucleic Acid Extraction kit	Medical Device	19.08.2020	18.08.2025 (Retention not submitted)	Generic	Submitted (page no. 118)
			Not submitted		Not submitted	Not submitted	Not submitted	Not submitted	Not submitted	Not submitted

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TENDER NO. BMSIC/MEDICAL DEVICES/CONSUMABLE/25-12

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

Company Name: - M/s Genetix Biotech Asia Ltd. Total Number of Pages Submitted in bid documents:- 344

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	6	Pap smear (DNA extraction kit)	(i) Not submitted	Not submitted	Not submitted	Not submitted	Not submitted	Not submitted	Not submitted	Not submitted
			MD- 9	MFG/IVD/2020/000112	NA	NA	19.08.2020	18.08.2025 (Retention not submitted)	20.10.2023	01.11.2025
			(iii) Not submitted	Not submitted	Not submitted	Not submitted	Not submitted	Not submitted	Not submitted	Not submitted

Note:- Assisted in technical evaluation in reference to letter no. BMSIC/40030/878-2025/9470 dt 05.12.2025 on the basis of documents provided by BMSICL as check list II, III & IV. Provided checklist compiled with due diligence and care. In spite, 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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