



Bihar Medical Services & Infrastructure Corporation Limited, 2nd & 3rd Floor, Swasthya Bhawan, Behind IGIMS, Sheikhpura, Adjacent to State Health Society, Patna 800023, Phone/Fax: +91612 2283287,+ 91612 2283288

Corrigendum-I

Bihar Medical Services and Infrastructure Corporation Limited (BMSICL) had invited E-Bids from the interested parties for the procurement, rate contract and the supply of medical equipment for different Govt. Institutions of Bihar vide Notice Inviting Tender No.- BMSICL/2025-26/ME-433. During and after Pre-bid meeting various suggestions were received from different prospective bidders regarding amendment in technical specification of equipment which were discussed and deliberated on by the experts, who after due deliberation recommended certain amendments in the technical specification of the equipment, which are annexed as Annexure-I of this corrigendum. In order to facilitate maximum participation of bidders the tender schedule is being revised as follows:-

Tender Reference No.	BMSICL/2025-26/ME-433
Last date and time of submission of online bids	11th September 2025 till 17:00 Hrs.
Last date and time of submission of original documents of EMD, Tender Fee and Document	12th September 2025 till 14:00 Hrs.
Date, Time and Place of opening of Technical Bid	12th September 2025 (at 15:00 Hrs.) on the website of https://eproc2.bihar.gov.in in the office of BMSICL
Date and time of opening of financial Bids	To be announced later on https://eproc2.bihar.gov.in

Note:-

- 1. Bidders are advised to refer to the Annexure-I of this corrigendum before submission of bid.**
- 2. Those who have submitted their bids are requested to re-submit their bids in accordance with this corrigendum.**

Annexed:- as above

**Sd/-
GM (Procurement)
BMSICL**

Annexure-I		
Amendment Request from Prospective Bidder for tender No-BMSICL/2025-26/ME-433		
Name of Equipment :- Automated Blood Culture Machine		
SI. No	Technical Specification as per tender	Final Amendment
1	The system should be a fully automated, walk away system capable of culture and detection of bacteria, mycobacteria and fungi from blood and sterile body fluids.	No Change
2	Should have capacity to hold at least 40 bottles at a time. The capacity should be upgradable.	Should have capacity to hold at least 100 bottles at a time. The capacity should be upgradable.
3	The system should continuously monitor the samples for growth and report it as and when it occurs.	No Change
4	The culture media provided should have sufficient mechanism to neutralize the Inhibitory effect of antibiotics and other substances in blood.	No Change
5	a) Culture media should be available for detecting bacteria and fungi, including fastidious organisms.	No Change
	b) The culture bottles should be unbreakable in normal conditions.	No Change
	c) The culture system should be suitable for processing blood and sterile body fluids.	No Change
6	Should be capable of processing both adult and the pediatric samples.	No Change
7	The system should be maintenance free without any need for regular calibrations, controls or standards run by the user.	No Change
8	The system should use leak proof and on non-invasive system to avoid contamination of equipment and environment.	No Change
9	The culture bottles should have high stability and (4-6) months shelf life.	No Change
10	The system should have all the facilities for data management and storage and quality control.	No Change
11	The system should be supplied in a complete system with all accessories, hardware like computer, printer, on line UPS (30 minutes battery backup) etc and required software.	No Change
12	Any software or database updates should be done free of cost by the firm, during the WARRANTY & CMC of equipment, as and when it is released by the manufacturer.	No Change
13	Any calibration, routine maintenance, and replacement of the parts like sensors, lamps etc required during the warranty and	No Change

	CMC and which is not covered by the same, should be declared and the respective costs quoted.	
14	Should give 200 Culture bottles (Aerobic/Paediatric) as Free of Cost along with the instrument.	No Change
15	The system should have Provision bi-directional LIS interface.	No Change
16	USFDA/European CE (Issued by Notified body) approved model should be offered.	No Change