



GOVERNMENT OF JAMMU & KASHMIR  
JAMMU AND KASHMIR MEDICAL SUPPLIES CORPORATION LTD.  
(Public Sector Undertaking of the Government of Jammu and Kashmir)

Corporate Head Office: Corporate Head Office: Plot No. 58, Friends Colony Satyam Road Trikuta Nagar Jammu

Corporate Office: Kashmir, Near Haj House, Benina 190018, Srinagar

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**Name of the Project** : Jhelum and Tawi Flood Recovery Project.  
**Name of the Work** : Supply and Installation of Digital Hand Held X Ray System (80 Numbers) including Comprehensive Maintenance of Five Years for Strengthening of Healthcare Facilities through JTFRP  
**Tender Id No.** : JKMSCL/Criticalcare/2025-26/New/05.  
**Date of Publishing the Bidding Document** : 27.04.2026  
**Date of Pre Bid Meeting** : 04-05-2026  
**Mode of Pre Bid Meeting** : Hybrid.  
**Date of Issue of Corrigendum 1** : 26.05.2026  
**Date of Issue of Corrigendum 2** : 02.06.2026  
**Date of Issue of Corrigendum 3** : 08.06.2026  
**Date of Issue of Corrigendum 4** : 15.06.2026

Table- 1 dated 19.06.2026

(Response/Clarification to Minutes of Pre-bid Meeting dated 04.05.2026)

S. No.	Clause No. ITB/GCC/ SSC/Forms	Gist of the Query	Response of the Purchaser	Reference to Sl. No of Corrigendum [Table 2] wherever applicable	Remarks
1	Section III. Evaluation and Qualification Criteria 1.1 Qualification Criteria (ITB 32.1) A. If a bidder is manufacturer (b) Experience and Technical Capacity	Experience and Technical Capacity Request to consider Experience of FY 2025-26 also.	Accepted	1	Modified

2	<p>Section III. Evaluation and Qualification Criteria  1.1 Qualification Criteria (ITB 32.1)  A. If a bidder is manufacturer    (a) Financial Capability Average Annual Turnover for the last three financial years</p>	Request to consider FY 2025-26 for Average Annual Turn Over requirement.	Bid prevail. Conditions	NA	No Change
3	<p>Section III. Evaluation and Qualification Criteria  1.1 Qualification Criteria (ITB 32.1)  B. If the Bidder is not a Manufacturer</p>	Request to consider Supply of Medical Equipment in the Similar Experience requirement if the bidder is not a manufacturer.	Accepted	2	Modified
4	<p>Section VII, Schedule of Requirements, Part 3, Annexure 1  Mandatory Requirement.</p>	It is suggested that the requirement of integration with EMR, Nikshay and NAAT be removed.	Bid prevail. Conditions	NA	No Change
5	<p>Section VII, Schedule of Requirements, Part 3, Annexure 1  Mandatory Requirement.</p>	The provision of Make in India to be Mandatory may be removed.	Accepted	3	Modified.
6.	<p>Section VII, Schedule of Requirements, Part 3, Annexure 1  Mandatory Requirement.</p>	Request to revise the Technical Specifications	Accepted	3	Modified



JKMSCL

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Table – 2 dated 19.06.2026  
(Corrigendum 5: As a result of Pre-bid Meeting dated 04.05.2026)

S.No.	Clause No. ITB/GCC/SSC/ Forms	As existing	As Amended	Reference to Sl. No of Response [Table 1] wherever applicable	Remarks

<p><b>1</b></p> <p><b>Section III. Evaluation and Qualification Criteria</b></p> <p><b>1.2 Qualification Criteria (ITB 32.1)</b></p> <p><b>A. If a bidder is manufacturer</b></p> <p><b>(b) Experience and Technical Capacity</b></p>	<p><b>If the bidder is manufacturer</b></p> <p>Experience and Technical Capacity</p> <p>The bidder should have successfully completed or substantially completed within the period of last three years at least one Contract of similar product (i.e similar equipment of different variants/models shall be accepted) to the extent of <b>at least 50% of the quantity</b> indicated against each schedule under "Section –VII, Schedule of Requirements" in any one of the last three financial years (2024-2025, 2023-2024, 2022-23). They should not have any adverse report regarding supplies for at least three years on the date of bid opening. A declaration may be furnished.</p>	<p><b>If the bidder is manufacturer</b></p> <p>Experience and Technical Capacity</p> <p>The bidder should have successfully completed or substantially completed within the period of last three years at least one Contract of similar product (i.e similar equipment of different variants/models shall be accepted) to the extent of <b>at least 50% of the quantity</b> indicated against each schedule under "Section –VII, Schedule of Requirements" in any one of the last three financial years (2025-2026, 2024-2025, 2023-24). They should not have any adverse report regarding supplies for at least three years on the date of bid opening. A declaration may be furnished.</p>	<p><b>1</b></p>	<p><b>Amended</b></p>
<p><b>2.</b></p> <p><b>Section III. Evaluation and Qualification Criteria</b></p> <p><b>1.1 Qualification Criteria (ITB 32.1)</b></p> <p><b>B. If the Bidder is not a Manufacturer</b></p>	<p><b>If the bidder is not manufacturer:</b></p> <p>If a bidder is not a manufacturer, but is offering the goods on behalf of the Manufacturer under Manufacturer's Authorization Form (Section IV Bidding Forms) the manufacturer shall demonstrate the above qualifications as detailed above at (a) and (b) and the bidder shall demonstrate that it has successfully completed at least one contract of similar nature of value not less than <b>Rs 05 (Five) crore</b> in the past three financial years.</p>	<p><b>If the bidder is not manufacturer:</b></p> <p>If the bidder is not manufacturer, but is offering the goods on behalf of the Manufacturer under Manufacturer's Authorization Forms (Section IV bidding Forms) the Manufacturer shall demonstrate the above qualifications as detailed above at (a) and (b) and the bidder shall demonstrate that it has successfully completed at least one contract of similar nature (<b>Supply of Medical Equipment</b>) of value not less than <b>Rs 05 (Five) crore</b> in the past three financial years (2025-2026, 2024-2025, 2023-24).</p>	<p><b>3</b></p>	<p><b>Amended</b></p>
<p><b>3.</b></p> <p><b>Section VII</b></p> <p><b>Schedule of Requirements, Part 3</b></p>	<p><b>INTRODUCTION:</b></p> <p>Digital ultraportable Radiographic system to capture X-ray images of the</p>	<p><b>INTRODUCTION:</b></p> <p>Digital ultraportable Radiographic system to capture X-ray images of the chest, using</p>	<p><b>5 and 6</b></p>	<p><b>Amended</b></p>

<p><b>Annexure I Mandatory Requirement.</b></p>	<p>chest, using digital portable technology. An ultraportable X-ray machine with complete Generator source, detector, stand, acquisition system for chest X-ray should be available.</p> <p>The following are the mandatory conditions to be fulfilled by the firm for responding to the tender.</p> <p><b>MANDATORY REQUIREMENT:</b> ICMR Field Approval is mandatory: ICMR verify machine working in different geographical conditions and in the actual feel conditions must be fully integrated with EMR, Nikshay and NAAT: First point of Test is X-Ray to identify the probable TB patients and NAAT is only approved machine by government of India for confirmation of TB Patient. So the data should be able to integrate with NAAT machine. Make in India is mandatory: To avoid sub-standard product.</p> <ul style="list-style-type: none"> <li>• Must have deployed at least 10 units of same quoted model in Govt Setups for TB Screening.</li> </ul> <p><b>CERTIFICATION for the quoted product :</b></p> <ul style="list-style-type: none"> <li>• The quoted digital X-ray system should be an ultra-portable type and should have AERB Type approval certificate.</li> </ul>	<p>digital portable technology. An ultraportable X-ray machine with complete Generator source, detector, stand, acquisition system for chest X-ray should be available.</p> <p>The following are the mandatory conditions to be fulfilled by the firm for responding to the tender.</p> <p><b>MANDATORY REQUIREMENT:</b> ICMR Field Approval is mandatory: ICMR verify machine working in different geographical conditions and in the actual feel conditions must be fully integrated with EMR, Nikshay and NAAT: First point of Test is X-Ray to identify the probable TB patients and NAAT is only approved machine by government of India for confirmation of TB Patient. So the data should be able to integrate with NAAT machine.</p> <ul style="list-style-type: none"> <li>• <b>Must have deployed at least 10 units of same quoted model in Govt Setups for TB Screening.</b> "The OEM should have successfully supplied and installed a minimum of 10 units of the same quoted model or any higher/similar model from the same OEM in Government healthcare institutions, Government screening programs, Government hospitals, Government medical colleges, or public health programs. Documentary</li> </ul>	
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	<ul style="list-style-type: none"> <li>• X-Ray device should be BIS IS 7620 (Part 1): 1986 or Electrical Safety approval for the product or FDA/CE 8s applicable.</li> <li>• CDSCO registration required for the quoted product and /OR manufacturing License.</li> </ul> <p><b>SYSTEM DESCRIPTION:</b></p> <p><u>Digital X-Ray system</u></p> <ul style="list-style-type: none"> <li>• The System should be capable of taking X-Rays of maximum human anatomies with minimal radiation exposure to patient /operator/attendants.</li> <li>• The X-Ray device should be lightweight, less than or equal to 7 kg including batteries.</li> <li>• The Radiation exposure time should be settable between 2 sec or Less with varying setting provision to allow imaging of any anatomy with minimal radiation exposure. X-Ray Generator / tube voltage should be 40-110 kv (Variable or fixed , value within this range).</li> <li>• Tube current between 2 to 6 mAs or higher.</li> <li>• Focal spot: 0.4mm to 0.8 mm or more with target angle 12 or more.</li> </ul>	<p><b>evidence shall be submitted along with the bid.</b></p> <p><u>CERTIFICATION for the quoted product :</u></p> <ul style="list-style-type: none"> <li>• The offered handheld/portable X-ray system shall be duly approved by the Atomic Energy Regulatory Board (AERB), Government of India. The bidder/OEM shall submit a valid AERB Type Approval Certificate for the quoted model along with the bid.</li> <li>• The digital x-ray system should have AERB Type approval certificate</li> <li>• The offered product should have any one of the following valid certifications: US FDA Approval / European CE Certificate issued by a 4-Digit Notified Body / BIS Certification. Relevant documentary evidence shall be submitted along with the bid.</li> <li>• The offered product shall have a valid CDSCO Manufacturing License or CDSCO Import License, as applicable, under the provisions of Medical Device Rules 2017. Relevant documentary evidence shall be submitted along with the bid.</li> </ul> <p><b>SYSTEM DESCRIPTION:</b></p> <p><u>Digital X-Ray system</u></p> <ul style="list-style-type: none"> <li>• The System should be capable of taking X-Rays of maximum human anatomies with minimal radiation exposure to patient /operator/attendants.</li> </ul>	
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		<ul style="list-style-type: none"> <li>• Minimum heat capacity 4000-6000 Joule or more.</li> <li>• X-ray should have wireless remote control.</li> <li>• Device should have X-Ray.</li> <li>• The X-Ray device should have digital or touch screen display of size 2 inch or more with APR based or simpler exposure settings and should show critical parameters Like KV, ma, and exposure time, battery charge Indicator with Low battery exposure out off and error messages etc.</li> <li>• The system should come with a large detector of at least 350 X 400 mm or more mm for taking images of all body parts.</li> <li>• The detectors should have a minimum spatial resolution of 3.5 l p/mm or more.</li> <li>• The detector must be high resolution of at least 2500 X 3000 pixels and 140 microns or less pixel pitch with A to D conversion 16 bits or more.</li> <li>• The Detector must have DQE more than 75% and MTF more than 80%.</li> <li>• The detector must be solid, sealed, water resistant, dust resistant and drop to a certain extent with appropriate certifications for these parameters</li> </ul>		
		<ul style="list-style-type: none"> <li>• The X-Ray device should be lightweight, less than or equal to 7 kg including batteries.</li> <li>• Radiation exposure time should be adjustable up to 2 seconds or less.</li> <li>• The system should provide variable exposure settings suitable for imaging of different anatomical regions while ensuring minimal radiation exposure to the patient. <i>X-Ray Generator / Tube Voltage: Fixed or variable tube voltage system. The operating tube voltage should be within the range of 40 kV to 110 kV. Systems operating at any fixed value or adjustable value within this range shall be acceptable.</i></li> <li>• Tube current between 2 to 6 mAs or higher.</li> <li>• Focal Spot Size: 0.4 mm to 0.8 mm or more.</li> <li>• Target Angle: 12° or more. Minimum heat capacity 4000-6000 Joule or more</li> <li>• The X-ray system should have wired or wireless remote control. All cables, accessories, and components required for complete functioning of the system shall be supplied along with the equipment..</li> <li>• The offered system should include an X-ray source suitable for digital radiographic imaging..</li> <li>• The X-Ray device should have digital or touch screen display of size 2 inch or</li> </ul>		

	<ul style="list-style-type: none"> <li>• The DR Console should be offered on a laptop with 14" screen of reputed make. (Minimum specifications: 11th Gen processor or 12 gen processor with &gt;8GB RAM &amp; 1 TB SOD)</li> <li>• Suitable portable light weight Tripod Stand with vertical movement range (50-150cm), rotation of 90 degree around vertical axis.</li> <li>• Wired or wireless Remote control enabled X-Ray acquisition.</li> <li>• Solid carrying case for all components which should be suitable for carriage of complete system including source, detector, acquisition console, stands for both source and detector by one person 1 operator.</li> <li>• The system must offer at least 3 years standard warranty and training with adequate manpower should be provided</li> </ul> <p><u>Training:</u></p> <ul style="list-style-type: none"> <li>• Operator/set training to be provided to the Radiology operator and clinician/radiographer on the use of the hardware &amp; software.</li> <li>• Operator/user training on troubleshooting and first line maintenance.</li> <li>• Artificial Intelligence Software.</li> </ul>		
		<p>more with APR based or simpler exposure settings and should show critical parameters Like KV, ma, and exposure time, battery charge Indicator with Low battery exposure out off and error messages etc.</p> <ul style="list-style-type: none"> <li>• The system should come with a large digital detector of at least 350 × 400 mm or more for taking images of all body parts.</li> <li>• The detectors should have a minimum spatial resolution of 3.5 lp/mm or more..</li> <li>• The detector must be high resolution of at least 2500 X 3000 pixels and 140 microns or less pixel pitch with A to D conversion 16 bits or more.</li> <li>• The Detector must have DQE more than 75% and MTF more than 80%.</li> <li>• The detector must be solid, sealed, water resistant, dust resistant and capable of withstanding accidental drops to a reasonable extent. Documentary evidence in the form of manufacturer specifications, product literature, test reports, certification documents, or equivalent supporting documents shall be submitted along with the bid,</li> <li>• The DR Console shall be provided on a laptop with a minimum 14-inch display and shall be suitable for operation of the offered handheld digital X-ray system, detector, acquisition software, AI application and data management platform. The laptop shall have a</li> </ul>	

		<ul style="list-style-type: none"> <li>• The AI should be able to triage/report subjects as Normal, TB related abnormality, Other Abnormality.</li> <li>• This software should work offline and online both. System should be loaded with CE or FDA approved AI software.</li> <li>• AI software must have CDSCO license.</li> <li>• Data Management system should be HIPPA and/or EU GDPR.</li> <li>• Complaint. ISO 27001, ISO 13485:2016 QMS.</li> </ul> <p>Provides a report that can be exported in MS excel/CVS and PDF report. Chest Application with response time less than 75 Sec. Live Demonstration of complete software starting from patient entry to data auto population. AI Interpretation will be integral part of evaluation.</p>	
		<p>minimum Intel Core i5 11th Generation or higher/equivalent AMD processor, minimum 8 GB RAM, minimum 1 TB SSD storage, and a licensed operating system compatible with the offered software. The system shall support detector image acquisition and display, patient registration and demographic data entry, exposure parameter recording and storage, image zoom, pan, rotate, invert and annotation functions, image storage, retrieval and export. The laptop shall be capable of operating the offered AI software and facilitating data transfer and integration with EMR, Nikshay and related workflow applications, wherever applicable. It shall have integrated Wi-Fi connectivity and necessary USB ports/interfaces required for detector and peripheral connectivity. The laptop shall be supplied complete with all software licenses, cables, connectors, accessories and interfaces required for complete and smooth functioning of the offered system.</p> <ul style="list-style-type: none"> <li>• Suitable portable light weight Tripod Stand with vertical movement range (50-150cm), rotation of 90 degree around vertical axis.</li> <li>• The X-ray system should have wired or wireless remote control. All cables, accessories, and components required for complete functioning of the system</li> </ul>	

				<p>shall be supplied along with the equipment..</p> <ul style="list-style-type: none"> <li>• Solid carrying case for all components which should be suitable for carriage of complete system including source, detector, acquisition console, stands for both source and detector by one person I operator.</li> <li>• After-Sales Support: The supplier shall provide comprehensive after-sales support during the warranty period, including installation, commissioning, configuration, user training, preventive maintenance, software updates/upgrades (where applicable), troubleshooting, corrective maintenance, replacement of defective parts, technical support , detector calibration, AI software support, and ensuring continued operational functionality of the complete system. All labor, travel, spare parts and services required for maintenance of the system during the warranty period shall be provided without additional cost to the purchaser.</li> <li>• The successful bidder shall supply Operation Manuals, User Manuals and Technical/Service Manuals (hard copy and/or soft copy) along with the equipment at the time of installation and commissioning.</li> </ul>
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		<p><b><u>Training:</u></b></p> <ul style="list-style-type: none"> <li>• Operator/set training to be provided to the Radiology operator and clinician/radiographer on the use of the hardware &amp; software.</li> <li>• Operator/user training on troubleshooting and first line maintenance.</li> <li>• Artificial Intelligence Software : The AI software shall be offered as a separate software solution and shall be fully compatible with the quoted X-ray system. In case the AI software is not developed by the bidder, a tender-specific authorization from the AI software manufacturer/developer shall be submitted along with the bid. The authorization shall be issued exclusively in favour of one bidder for this tender and shall clearly authorize the bidder to quote, supply, install, support and maintain the offered AI software for the entire contract period.</li> <li>• The AI software shall be suitable for chest X-ray screening and TB screening workflows. At a minimum, it shall be capable of processing chest X-ray images and classifying/reporting subjects as Normal, TB-related Abnormality, or Other Abnormality. The AI software shall support generation and export of reports and shall comply with the regulatory, interoperability and performance</li> </ul>		
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			<p>requirements specified elsewhere in the tender document.</p> <ul style="list-style-type: none"> <li>• This software should work offline and online both.</li> <li>• <b>AI systems operational on secured hospital networks without mandating full standalone offline capability. Make CE/USFDA approval OPTIONAL for AI software; accept valid CDSCO license as fully compliant.</b></li> <li>• AI software must have CDSCO license.</li> <li>• Data Management system should be HIPPA and/or EU GDPR.</li> <li>• Complaint. ISO 27001, ISO 13485:2016 QMS.</li> <li>• The AI software shall be compatible with and capable of functioning as an AI-powered Computer-Aided Detection (CAD) solution for chest X-ray screening and TB screening applications.</li> <li>• The AI software shall support both online and offline operation. In offline mode, the software shall be capable of processing and reporting chest X-ray images locally without dependency on internet connectivity. In online mode, it shall support data synchronization, reporting, integration and other workflow functionalities, wherever applicable.</li> <li>• Provides a report that can be exported in MS excel/CVS and PDF report. Chest Application with response time less than</li> </ul>		
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		75 Sec. Live Demonstration of complete software starting from patient entry to data auto population, AI Interpretation will be integral part of evaluation.		
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In addition to the above, the following changes have been made to the provisions of the bid document by the Purchaser :

1.	Section VII of Schedule Requirements 1. List of Goods & Delivery Schedule, Final (Site) Destination as specified in BDS.	The Table in the bidding document as issued is replaced with the table attached to this corrigendum.	Modified by Purchaser	Modified
2	Section IX. Special Conditions of Contract GCC 16.1	<p><b>GCC 16.1</b></p> <p>Payment shall be made in Indian Rupees in the following manner:</p> <p>(a)</p> <p>(i) <b>Advance Payment:</b> 20% of the total contract price shall be paid within (30) days of signing of Contract and upon submission of claims/against a simple receipt and a bank guarantee for the equivalent amount valid until the goods are delivered and in the form provided in the bidding documents or another form acceptable to the Purchaser.</p> <p>(ii) <b>On Delivery:</b> Sixty (60) % of the contract price shall be paid on receipt of Goods and upon submission of the documents specified in Clause 13 of SCC; and</p>	<p><b>GCC 16.1</b></p> <p>Payment shall be made in Indian Rupees on a pro-rata basis of goods supplied, installed, tested, commissioned and training provided to the onsite operating staff in the following manner</p> <p>(a)</p> <p>i. <b>Advance Payment:</b> 20% of the total contract price shall be paid within (30) days of signing of Contract and upon submission of claims/against a simple receipt and a bank guarantee for the equivalent amount valid until the goods are delivered and in the form provided in the bidding documents or another form acceptable to the Purchaser.</p> <p>ii. <b>On Delivery:</b> Sixty (60) % of the contract price shall be paid on receipt of Goods at</p>	<p>Modified by Purchaser</p> <p>Modified</p>

	<p>(ii) <b>On Final Acceptance:</b> the remaining twenty (20) % of the Contract Price shall be paid after completion of the delivery of goods against an unconditional Bank Guarantee of equivalent Amount valid until successful installation of the goods which shall be released within thirty (30) days after the date of the Acceptance Certificate issued by the Purchaser's representative, successful installation &amp; verification of the equipment in the proforma given in Section VII - item 6.</p> <p>(b) Not applicable</p> <p>(c) (i) For all the payments to be made, against Bank guarantees, the bank guarantee shall be issued by a Scheduled Indian Bank or a foreign bank located in India in the format enclosed at Section X. The guarantees issued by other banks should be confirmed by a Scheduled Indian Bank or a foreign bank operating in India.</p> <p>(ii) Bank guarantees for advance payment shall be released not later than 30 days after the date of completion of supply of the goods at their final destination.</p> <p>Note: 1. No supplies shall be accepted beyond 31<sup>st</sup> of August 2026 unless agreed specifically by the Purchaser.</p>	<p>Regional Drug Warehouse Bemina and upon submission of the documents specified in Clause 13 of SCC; and</p> <p>iii. <b>On Final Acceptance:</b> The remaining twenty (20) % of the Contract Price shall be paid after successful installation, testing and training of the staff at site as per the Consignee List in Table 1 : List of Goods ad Delivery Schedule in Section VII, Schedule of Requirement. The payment shall be released after submission of the Acceptance Certificate issued by the Purchaser's representative at site as per the Consignee List.</p> <p>(b) Not applicable</p> <p>(c) (i) For all the payments to be made, against Bank guarantees, the bank guarantee shall be issued by a Scheduled Indian Bank or a foreign bank located in India in the format enclosed at Section X. The guarantees issued by other banks should be confirmed by a Scheduled Indian Bank or a foreign bank operating in India.</p> <p>(ii) Bank guarantees for advance payment shall be released not later than 30 days after the date of completion of supply of the goods at their final destination.</p> <p>Note: 1. No supplies shall be accepted beyond 31<sup>st</sup> of August 2026 unless agreed specifically by the Purchaser. 2. The supplier shall supply, install and provide training to user as per Consignee List in Table 1 : List of Goods and</p>		
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	<p>2. The delivery destination shall be made at Regional Drug ware House Bemina, Srinagar. The training shall be provided to the staff deployed at the installation site to be notified by the Purchaser to the Supplier during the warranty cum Comprehensive Maintenance period of Five Years.</p>	<p>Delivery Schedule in Section VII, Schedule of Requirements.</p>		
<p><b>3</b></p> <p><b>Section IX. Special Conditions of Contract GCC 28.3</b></p>	<p>The period of validity of the Warranty shall be Five Years.</p> <p>For purposes of the Warranty, the place(s) of final destination(s) shall be Regional Drug Ware House Bemina, Jammu and Kashmir Medical Supplies Corporation Limited Srinagar :</p> <p>i. The warranty period shall be sixty (60) months from date of acceptance of the Goods or sixty six (66) months from the date of shipment, whichever occurs earlier. The Supplier shall, in addition, comply with the performance and/or consumption guarantees specified under the Contract. If, for reasons attributable to the Supplier, these guarantees are not attained in whole or in part, the Supplier shall, pay liquidated damages to the Purchaser with respect to the failure to meet the contractual guarantees. The rate of these liquidated damages shall be 0.5% of the contract price for each equipment that fails to meet the contractual performance guarantees per week or part thereof.</p>	<p>The period of validity of the Warranty shall be Five Years.</p> <p>For purposes of the warranty, the place(s) of final destination(s) shall be the consignee sites as reflected in Table 1 : List of Goods and Delivery Schedule in Section VII, Schedule of Requirements.</p> <p>i. The warranty period shall be sixty (60) months from date of acceptance of the Goods. The Supplier shall, in addition, comply with the performance and/or consumption guarantees specified under the Contract. If, for reasons attributable to the Supplier, these guarantees are not attained in whole or in part, the Supplier shall, pay liquidated damages to the Purchaser with respect to the failure to meet the contractual guarantees. The rate of these liquidated damages shall be 0.5% of the contract price for each equipment that fails to meet the contractual performance guarantees per week or part thereof subject to a maximum of 10% of the contract price for each equipment.</p> <p>ii. The Installation and the commissioning, training to users of the equipment is included in the price schedule with warranty period of</p>	<p>Modified by Purchaser</p>	<p><b>Modified.</b></p>

	<p>ii. The Installation and the commissioning, training to users of the equipment is included in the price schedule with warranty period of 05 years. The warranty of the equipment covers comprehensive maintenance of the equipment with spares and preventive maintenance. The Supplier shall, during the warranty period appearing in the contract, replace the whole subject matter of procurement or part(s), if any, and remove the manufacturing defects, if found during the above period so as to make the machinery and equipment/ordered items operative.</p> <p>iii. The Supplier warrants that all the Goods are new, unused, and of the most recent or current models, and that they incorporate all recent improvements in design and materials, unless provided otherwise in the Contract.</p>	<p>05 years. The warranty of the equipment covers comprehensive maintenance of the equipment with spares and preventive maintenance. The Supplier shall, during the warranty period appearing in the contract, replace the whole subject matter of procurement or part(s), if any, and remove the manufacturing defects, if found during the above period so as to make the machinery and equipment/ordered items operative.</p> <p>iii. The Supplier warrants that all the Goods are new, unused, and of the most recent or current models, and that they incorporate all recent improvements in design and materials, unless provided otherwise in the Contract.</p> <p>iv. During the warranty period, the Supplier shall be fully responsible for the proper functioning and performance of the equipment and shall provide, at no additional cost to the Purchaser, all preventive maintenance services, corrective maintenance, repairs, replacement of defective or worn-out spare parts, accessories, consumables essential for operation, software updates/upgrades, calibration, safety checks, and labour required for maintaining the equipment in optimal operating condition in accordance with OEM standards. The Supplier shall ensure availability of qualified service engineers and shall attend breakdown complaints within the response time specified in the Contract, failing which</p>	
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		<p>suitable penalties may be imposed as per tender conditions. Any equipment or component found defective during the warranty period shall be repaired or replaced with new genuine OEM parts without any additional financial implication to the Purchaser. The warranty shall also cover all integrated hardware, software, networking components, and peripherals supplied under the Contract. In case the equipment remains non-functional beyond the permissible downtime (i.e minimum 95% uptime is required), the warranty period for the affected equipment shall be extended proportionately for the duration of such downtime. The Supplier shall maintain proper service records/logbooks for all preventive and breakdown maintenance visits and submit the same to the designated authority whenever required.</p>		
<p><b>4.</b> <b>Section II</b> <b>Bid Data Sheet</b> <b>ITB 14.8</b> <b>(a) (iii)</b></p>	<p>“Final destination (Project Site)” is Jammu and Kashmir Medical Supplies Corporation Limited, Regional Drug Ware House Bemina Srinagar, Jammu and Kashmir, 190018 India as per delivery schedule.</p>	<p>“Final destination (Project Site)” is Consignee List in Table 1 : List of Goods and Delivery Schedule in Section VII, Schedule of Requirements.</p>	<p>Modified by Purchaser</p>	<p><b>Modified.</b></p>
<p><b>5.</b> <b>Section IX Special</b> <b>Conditions of</b> <b>Contract GC 1.1</b> <b>(o)</b></p>	<p>The Project Site(s)/Final Destination(s) is : Jammu and Kashmir Medical Supplies Corporation Limited Regional Drug Ware House Bemina Srinagar Jammu and Kashmir 190018 India</p>	<p>The Project sites(s)/“Final destination : Consignee List in Table 1 : List of Goods and Delivery Schedule in Section VII, Schedule of Requirements.</p>	<p>Modified by Purchaser</p>	<p><b>Modified.</b></p>

6.	E. Procurement Notice Invitation of Bids Sr. No. 4 (e) and 4 (f)	Last date and time for submission of online bids: 20.06.2026 upto 11.00 AM. Time and Date for opening of bids -Technical Part : 20.06.2026 at 11.15 hrs	Last date and time for submission of online bids: 27.06.2026 upto 11.00 AM. Time and Date for opening of bids -Technical Part : 27.06.2026 at 11.15 hrs	Amended by the Purchaser	Modified
7.	Section II – Bid Data Sheet ITB – 22.1	The deadline for uploading of bids is : Date [20.06.2026] Time [11.00 a.m]	The deadline for uploading of bids is : Date [27.06.2026] Time [11.00 a.m]	Amended by the Purchaser	
8.	Section II – Bid Data Sheet ITB – 25.1	The online opening of the Technical Parts of Bids shall take place at: JAMMU AND KASHMIR MEDICAL SUPPLIES CORPORATION LTD. (Public Sector Undertaking of the Government of Jammu and Kashmir) Corporate Head Office: Plot No. 58, Friends Colony Satyam Road Trikuta Nagar Jammu and Kashmir 180020 Country: INDIA Date of opening of e-bid : 20.06.2026 Time of opening of e-bid : 11.15 A.M	The online opening of the Technical Parts of Bids shall take place at: JAMMU AND KASHMIR MEDICAL SUPPLIES CORPORATION LTD. (Public Sector Undertaking of the Government of Jammu and Kashmir) Corporate Head Office: Plot No. 58, Friends Colony Satyam Road Trikuta Nagar Jammu and Kashmir 180020 Country: INDIA Date of opening of e-bid : 27.06.2026 Time of opening of e-bid : 11.15 A.M	Amended by the Purchaser	

### 1. List of Goods and Delivery Schedule

Line Item No	Description of Goods	Quantity	Physical unit	Final (Site) Destination as specified in BDS	Delivery (as per Incoterms) Date
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						Latest Delivery Date	Bid Security in Indian Rupees <sup>1</sup>
	[Insert item No]	[Insert description of Goods]	[Insert quantity of item to be supplied]	[Insert physical unit for the quantity]	[Insert place of Delivery]	[Insert the number of days following the date of Contract signature]	
	01	Digital Hand Held X Ray System (80 Numbers) including Comprehensive Maintenance of Five Years for Strengthening of Healthcare Facilities through JTRP	80	Number	As per Consignee List attached.	60	INR 500,000.00

Bid security listed here must be the same as provided under ITB/BDS 19.1

Note : [i] There is two stage delivery as below

Stage 1 : Delivery shall be at the Regional Drug Ware House Bemina, Jammu and Kashmir Medical Supplies Corporation Limited, Srinagar at address : 2<sup>nd</sup> Floor Corporate Office Bemina Srinagar 190018. The purpose of delivery at this stage is to allow the Purchaser to make entries in Stock Register. The process will take approximately an hour.

Stage 2 : Last Mile Delivery shall be as per Consignee List attached as Annexure 1 of this Table, after completion of Stage 1 delivery.

[ii] If the Consignee is unable to take delivery at Stage 2 within 1 working day, these items will be delivered back to the above warehouse after which it will be the responsibility of the Purchaser to make these deliveries as per Consignee List at the Purchaser's risk and cost.

**Annexure 1 to Table 1 : List of Goods and Delivery Schedule, Section VII : Schedule of Requirements  
Consignee List**

Destination	No. Of Equipment	Remarks

Government Medical College Jammu	15	List of individual hospitals to be provided at the time of Contract signing.
Government Medical College Doda	5	
Government Medical College Rajouri	5	
Government Medical College Kathua	5	
Hospitals under Director Health Services Jammu	10	
Government Medical College Srinagar	15	
Government Medical College Anantnag	4	
Government Medical College Baramulla	4	
Government Medical College Handwara	3	
Sher-I Kashmir Institute of Medical Sciences Srinagar (Director SKIMS)	4	
Hospitals under Director Health Services Kashmir	10	List of individual hospitals to be provided at the time of Contract signing.

No. JKMSCL/2026/ 1695

Dated: 19.06.2026

  
 General Manager  
 J&K Medical Supplies Corporation Ltd