

**Meeting Notice**

**Subject: Opening of Financial Bid(s) for Procurement of "Ultraportable Handheld X-Ray with Computer Aided Detection (CAD) Technology and Artificial Intelligence (AI) Machine"- reg.**

In pursuance of the approval of the Competent Authority and in continuation of the procurement process under NIT No.JKMSCL/M&E/2025/670 dated 19.08.2025, it is hereby notified that the financial bid in respect of the item "Ultraportable Handheld X-Ray with Computer Aided Detection (CAD) Technology and Artificial Intelligence (AI) Machine", falling under the category Machinery and Equipment, shall be opened as per the details given below:

Date: 14.01.2026

Time: 02:00 PM

Venue: Conference Hall, Corporate Head Office/ Srinagar, J&K Medical Supplies Corporation Limited, Jammu.

Accordingly, the following officers are requested to kindly attend the meeting on the scheduled date, time and venue:

1. Managing Director, JKMSCL
2. Financial Advisor/ Chief Accounts Officer, JKMSCL
3. General Manager (K), JKMSCL
4. General Manager (Adm), JKMSCL

**Issued with the approval of the Competent Authority.**

10.01.26

**(M A Choudhary) JKAS**  
General Manager (Adm)  
J&K Medical Supplies Corporation Limited

No.: JKMSCL/GM/Adm/2025-26/4165-68

Dated: 10.01.2026.

**Copy for the information to:**

1. Managing Director, J&K Medical Supplies Corporation Limited.
2. Financial Advisor & CAO, J&K Medical Supplies Corporation Limited.
3. General Manager (K), J&K Medical Supplies Corporation Limited.
4. Assistant Programmer for necessary action.
5. Office Copy



# 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

Corporate Office: Corporate Office Kashmir, Near Haj House, Bemina (190018), Srinagar

email: gmjkkmscl@gmail.com, jkmsclpm@gmail.com; website: www.jkmscl.nic.in

JKMSCL

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## MINUTES OF MEETING

**Subject: Meeting of the Technical Evaluation/Advisory Committee held on 29.12.2025 for the acceptance for demonstration report for the finalization of Rate Contract for the procurement of "Ultraportable Handheld X-Ray with Computer Aided Detection (CAD) Technology and Artificial Intelligence (AI) Machine" under NIT No. JKMSCL/Mach/2025/670 dated 19.08.2025.**

1. E- bid for the finalization of Rate Contract for the procurement of "Ultraportable Handheld X-Ray with Computer Aided Detection (CAD) Technology and Artificial Intelligence (AI) Machine" under Group Machinery & Equipments was uploaded vide no NIT No: NIT/JKMSCL/Mach/2025/670 Dated 19.08.2025 and was opened on 07<sup>th</sup> November 2025 at Corporate Head Office, Jammu and Srinagar at 11.00 A.M. The aforesaid tender was opened in the presence of technical advisory/evaluation committee constituted for the purpose by Board of Directors, JKMSCL during 3<sup>rd</sup> board meeting for the purpose.
2. With reference to clause 2.1.9 Chapter II of the Standard Procurement Procedure approved during 2<sup>nd</sup> Board Meeting, it was decided by the technical advisory/evaluation committee that the Chairman, Technical Advisory/Evaluation Committee shall get the preliminary evaluation done by the sub-committee and then shall get the minor infirmities asked from the respective bidders, if any. After downloading the minor infirmities submitted by the bidders, the same shall be re-evaluated by the Sub Committees followed by technical evaluation by the technical experts and the final report shall be placed before the Technical Advisory Committee for further decision/recommendation.
3. Further, sub committees constituted to assist the Technical Evaluation Committee of JKMSCL completed the compilation and scrutiny/evaluation process and thereafter the observations made by the sub- committee were placed before Technical Advisory Committee constituted for the purpose by Board of Director, JKMSCL for final scrutiny/evaluation of bid documents so as to arrive at qualifying bidders for further evaluation by means of power point presentation/ demonstration and catalogue evaluation by the technical experts :

S No.	Name of bidder	Model	Infirmitiess/shortcomings	Status	Remarks
<b>01. Ultraportable Handheld Digital X-Ray KIT (with AI Technology Based Software for TB Screening)</b>					
1/5	<b>Bidder:</b> M/s Fidoc Ventures Private Limited Gurugram Haryana <b>Manufacturer:</b> M/s Prognosys Medical Systems Pvt. Ltd. Bangalore and Deeptek medical Imaging Pvt Ltd -AS Pre-Annexure I	<b>Prorad Atlas Ultra-Portable X ray System</b>	<b>To Submit:</b> 1. Compliance Sheet of the quoted equipment (self-attested) (As per corrigendum). 2. Technical Bid Submission Sheet, duly filled. (The submitted sheet is without page numbering.). 3. Client Base of the quoted model/make with reference of the supply orders, for any of three financial years in Last five years along with satisfactory performance certificate of	Submitted Submitted Submitted	"Recommended to the Technical Advisory/Evaluation Committee for further necessary action."

*W Nov 4 2025 JKMSCL*

		<p>minimum one purchase order and installation report of Govt, Institution (Copies of reference supply orders and satisfactory performance certificate need to be attached)</p> <p>4. To submit minimum of 5 years' experience in the healthcare sector and also submit documentary proof of having supplied at least 10 units of the same quoted model in Government institutions for TB screening.</p> <p>5. Average Annual Turnover Statement for the last 3 financial years (2021-22, 2022-23, and 2023-24) of the Principal Manufacturer, issued by a Chartered Accountant/competent authority with UDIN. <i>(The bidder has submitted its own turnover statement, which is not acceptable in this case.)</i></p> <p>6. Copies of audited Balance Sheet and Profit &amp; Loss Account for the last three financial years (2021-22, 2022-23, and 2023-24) of the Principal Manufacturer, certified by a Chartered Accountant with UDIN. <i>(The bidder has submitted its own audit balance sheets, which is not acceptable in this case.)</i></p> <p>7. For X-ray Device Certification submit BIS standard IS 7620 (Part 1):1986 or shall have valid Electrical Safety Approval or US FDA / CE certification, as applicable</p> <p>8. For ICMR Approval submit ICMR field approval</p> <p>9. For NAAT Approval submit NAAT approval for use in TB screening applications</p> <p>10. For AERB Approval of the principal manufacturer/bidder submit a valid AERB approval certificate for the quoted equipment</p> <p>11. For AI Software Certification submit CE or US FDA approved Artificial Intelligence (AI) software.</p> <p>12. For AI Software Licensing submit a valid CDSCO license</p> <p>13. submit HIPAA and/or EU GDPR standards for data protection and privacy</p> <p>14. Submit ISO 27001 (Information Security Management System) and ISO 13485:2016 (Quality Management System for Medical Devices).</p> <p>15. <b>Clarification - Deepak Medical Imaging Pvt. Ltd.:</b> The bidder has submitted the US FDA certificate, CDSCO registration, and other certificates pertaining to M/s Deepak Medical Imaging Pvt. Ltd. The bidder is requested to clarify their role/association with M/s Deepak Medical Imaging Pvt. Ltd. and provide all relevant documentary evidence in support of the same as per NIT. It is also noted that the same documents were found in the technical bid of M/s SS Agency, Jammu, and</p>	Submitted	
			Submitted and clarified	

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			<i>clarification is required regarding this duplication.</i>		
2/5	<p><b>Bidder:</b> M/s SS Agencies Jammu  <b>Importer:</b> M/s Lab India Healthcare Pvt. Ltd  <b>Manufacturer:</b> OTOM Korea</p>	<b>Mine 2</b>	<p><b>To Submit:</b></p> <ol style="list-style-type: none"> <li>1. Compliance Sheet of the quoted equipment (self-attested) (As per corrigendum).</li> <li>2. Submit a satisfactory performance certificate for at least one purchase order, along with the corresponding installation report from a government institution, related to the reference supply orders submitted in the technical bid.</li> <li>3. To submit minimum of 5 years' experience in the healthcare sector and also submit documentary proof of having supplied at least 10 units of the same quoted model in Government institutions for TB screening.</li> <li>4. As per the catalogue submitted, both M/s Lab India Healthcare and OTOM are shown as associated entities. The bidder is required to clarify the relationship between the Principal Manufacturer and the Importer (VizM/s Lab India Healthcare and OTOM) and submit all supporting documentary evidence establishing this relationship.</li> <li>5. Submit Average Annual Turnover Statement for the last 3 financial years (2021-22, 2022-23, and 2023-24) of the importer, issued by a Chartered Accountant/competent authority with UDIN. (Submitted is without UDIN)</li> <li>6. Self-attested photo copy of IEC certificate and Permission/Authorization for sale for sale from the foreign principal manufacturer (in case of imported product)</li> <li>7. For X-ray Device Certification resubmit BIS standard IS 7620 (Part 1):1986 or shall have valid Electrical Safety Approval or US FDA / CE certification, as applicable</li> <li>8. For X-ray Device Certification submit BIS standard IS 7620 (Part 1):1986 or shall have valid Electrical Safety Approval or US FDA / CE certification, as applicable.</li> <li>9. For ICMR Approval submit ICMR field approval</li> <li>10. For NAAT Approval submit NAAT approval for use in TB screening applications</li> <li>11. For AERB Approval of the principal manufacturer/bidder submit a valid AERB approval certificate for the quoted equipment.</li> <li>12. For AI Software Certification submit CE or US FDA approved Artificial Intelligence (AI) software.</li> </ol>	<p><b>Submitted</b></p>	<p>Not Recommended to the Technical Advisory/Evaluation Committee</p>

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			<p>13. For AI Software Licensing submit a valid CDSCO license.</p> <p>14. Submit <i>HIPAA and/or EU GDPR standards for data protection and privacy</i></p> <p>15. Submit ISO 27001 (Information Security Management System) and ISO 13485:2016 (Quality Management System for Medical Devices).</p> <p>16. Clarification - Deepak Medical Imaging Pvt. Ltd.: The bidder has submitted the US FDA certificate, catalogue, CDSCO registration, and other certificates pertaining to M/s Deepak Medical Imaging Pvt. Ltd. The bidder is requested to clarify their role/association with M/s Deepak Medical Imaging Pvt. Ltd. and provide all relevant documentary evidence in support of the same as per NIT. It is also noted that the same documents were found in the technical bid of M/s Findoc Ventures Private Limited Gurugram Haryana, and clarification is required regarding this duplication.</p> <p>17. Clarification - LG Electronics India Pvt. Ltd.: The bidder has submitted the US FDA certificate, catalogue, CDSCO registration, and other certificates of M/s LG Electronics India Pvt. Ltd. The bidder is requested to clarify their role/association with M/s LG Electronics India Pvt. Ltd. and submit all relevant documentary evidence in support of NIT.</p>	<p>Submitted of M/s Deep tech medical imaging Pvt. Ltd.</p> <p>Submitted M/s Deep tech medical imaging Pvt. Ltd.</p> <p><b>Submitted</b></p> <p><b>Not submitted as per requirement</b></p> <p><b>Not submitted as per requirement</b></p>	
3/5	<p>M/s Genuine Medica Private Limited Delhi <b>(Bidder)</b></p> <p>M/s Edusoft Healthcare Pvt. Ltd New Delhi <b>(Manufacturer)</b></p>	ERAY SMART 5HS	<p><b>To Submit:</b></p> <ol style="list-style-type: none"> <li>1. Compliance Sheet of the quoted equipment (self-attested) (As per corrigendum).</li> <li>2. Technical Bid Submission Sheet, duly filled. (The submitted sheet is without page numbering).</li> <li>3. Declaration for Latest Non Conviction, non blacklisting on non judicial Stamp paper of Rs 100 furnished by the Principal Manufacturer as per proforma duly notarised.</li> <li>4. Declaration for Latest Non Conviction, non blacklisting on non judicial Stamp paper of Rs 100 furnished by the bidder as per proforma duly notarised.</li> <li>5. Submit a satisfactory performance certificate of the quoted model for at least one purchase order, along with the corresponding installation report from a government institution, related to the reference</li> </ol>	<p><b>Submitted</b></p> <p><b>Submitted</b></p> <p><b>Submitted</b></p> <p><b>Submitted</b></p> <p><b>Submitted</b></p>	<p>Not Recommended to the Technical Advisory/ Evaluation Committee</p>

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		supply orders submitted in the technical bid.	
6.	To submit minimum of 5 years' experience in the healthcare sector and also submit documentary proof of having supplied at least 10 units of the same quoted model in Government institutions for TB screening.	Submitted for 2022-23, 2023-24 and 2024-25 only.	
7.	Average Annual Turnover Statement for the last 3 financial years (2021-22, 2022-23, and 2023-24) of the Principal Manufacturer, issued by a Chartered Accountant/competent authority with UDIN. <i>(The bidder has submitted its own turnover statement, which is not acceptable in this case.)</i>	Submitted	
8.	Copies of audited Balance Sheet and Profit & Loss Account for the last three financial years (2021-22, 2022-23, and 2023-24) of the Principal Manufacturer, certified by a Chartered Accountant with UDIN. <i>(The bidder has submitted its own audited Balance Sheet, which is not acceptable in this case.)</i>	Submitted	
9.	For X-ray Device Certification submit BIS standard IS 7620 (Part 1):1986 or shall have valid Electrical Safety Approval or US FDA / CE certification, as applicable	Submitted	
10.	For ICMR Approval submit ICMR field approval	Submitted	
11.	For NAAT Approval submit NAAT approval for use in TB screening applications	Submitted	
12.	For AERB Approval of the principal manufacturer/bidder submit a valid AERB approval certificate for the quoted equipment.	Submitted	
13.	For AI Software Certification submit CE or US FDA approved Artificial Intelligence (AI) software.	Submitted of Qure.ai Technologies Pvt Ltd	
14.	For AI Software Licensing submit a valid CDSCO license.	Submitted Qure.ai Technologies Pvt Ltd	
15.	Submit HIPAA and/or EU GDPR standards for data protection and privacy.	Submitted of Qure.ai Technologies Pvt Ltd	
16.	Submit ISO 27001 (Information Security Management System) and ISO 13485:2016 (Quality Management System for Medical Devices).	Submitted	
17.	<b>Clarification:</b> <i>The bidder has submitted the Certificates pertaining to Qure.ai Technologies Pvt Ltd. The bidder is requested to clarify their role/association with Qure.ai Technologies Pvt Ltd. and provide all relevant documentary evidence in</i>	Not submitted as per requirement	

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Signature

			<p>support of the same as per NIT.</p> <p><b>18. Clarification:</b> The bidder has submitted the Certificates pertaining to <b>Inno Care Optoelectronics Corp</b> . The bidder is requested to clarify their role/association with <b>Inno Care Optoelectronics Corp</b> . and provide all relevant documentary evidence in support of the same as per NIT.</p> <p><b>19. Name, photograph &amp; specimen signature of the designated officer/ representative of the Bidder who is authorized to make correspondence with the JKM SCL</b></p>	<p>Not submitted as per requirement</p> <p>Submitted</p>	
4/5	<p><b>Bidder:</b> M/s Surgident India <b>Manufacturer:</b> Starnuke Consulting Engineering Services Pvt. Ltd., Jaipur</p>	<p><b>Mine 2 in</b></p>	<p><b>To Submit:</b></p> <ol style="list-style-type: none"> <li>1. Submit Proof of payment towards the cost of tender and tender processing fee has not been submitted.</li> <li>2. Submit Compliance Sheet of the quoted equipment (self-attested) (As per corrigendum).</li> <li>3. Technical Bid Submission Sheet, duly filled. (The submitted sheet is without page numbering.).</li> <li>4. Declaration for Latest Non Conviction, non blacklisting on non judicial Stamp paper of Rs 100 furnished by the bidder as per proforma duly notarised.</li> <li>5. Declaration for Latest Non Conviction, non blacklisting on non judicial Stamp paper of Rs 100 furnished by the Principal Manufacturer as per proforma duly notarised.</li> <li>6. Resubmit Client Base of the quoted model/ make with reference of the supply orders, for any of three financial years in Last five years along with satisfactory performance certificate of minimum one purchase order and installation report of Govt, Institution (Copies of reference supply orders and satisfactory performance certificate need to be attached)</li> <li>7. To submit minimum of 5 years' experience in the healthcare sector and also submit documentary proof of having supplied at least 10 units of the same quoted model in Government institutions for TB screening.</li> <li>8. Average Annual Turnover Statement for the last 3 financial years (2021-22, 2022-23, and 2023-24) of the Principal Manufacturer, issued by a Chartered Accountant/competent authority with UDIN. (<i>The bidder has submitted its own turnover statement, which is not acceptable in this case.</i>)</li> <li>9. Copies of audited Balance Sheet and Profit &amp; Loss Account for the last three financial years (2021-22,</li> </ol>	<p>Submitted</p> <p>Submitted</p> <p>Submitted</p> <p>Submitted</p> <p>Submitted</p> <p>Submitted</p> <p>Submitted</p> <p>Submitted</p> <p>Submitted</p>	<p>Recommended to the Technical Advisory/ Evaluation Committee for further necessary action.</p>

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			<p>2022-23, and 2023-24) of the Principal Manufacturer, certified by a Chartered Accountant with UDIN. (The bidder has submitted its own audit balance Sheets, which is not acceptable in this case.)</p> <p>10. For X-ray Device Certification submit BIS standard IS 7620 (Part 1):1986 or shall have valid Electrical Safety Approval or US FDA / CE certification, as applicable</p> <p>11. For ICMR Approval submit ICMR field approval</p> <p>12. For NAAT Approval submit NAAT approval for use in TB screening applications</p> <p>13. For AERB Approval of the principal manufacturer/bidder submit a valid AERB approval certificate for the quoted equipment.</p> <p>14. For AI Software Certification submit CE or US FDA approved Artificial Intelligence (AI) software.</p> <p>15. Submit CDSO registration/License for the quoted item.</p> <p>16. For AI Software Licensing submit a valid CDSCO license.</p> <p>17. submit HIPAA and/or EU GDPR standards for data protection and privacy</p> <p>18. Submit ISO 27001 (Information Security Management System) and ISO 13485:2016 (Quality Management System for Medical Devices).</p> <p>19. Authorization from principal manufacturer (On the letterhead of Principal manufacturer) is to be submitted (strictly as per annexure VI)</p> <p>20. Nature of the Firm/Public Company / Private Company/ Partnership/ Proprietorship/any other with Documentary proof.</p>	Submitted	
5/5	<p>M/s G.T. Traders Narayana Nagar Jalandhar, Punjab <b>(Bidder)</b></p> <p>M/s Rohit Jafa Ventures India Private, Limited. <b>(Manufacturer)</b></p>	RX- 706	<p><b>To Submit:</b></p> <ol style="list-style-type: none"> <li>1. Tender processing fee and tender fee not submitted, Hence cant not be considered <i>(Submit Proof of payment towards the cost of tender and tender processing fee if submitted, then accordingly submit the below mentioned documents.)</i></li> <li>2. Compliance Sheet of the quoted equipment (self-attested) (As per corrigendum).</li> <li>3. Technical Bid Submission Sheet, duly filled. (The submitted sheet is without page numbering.).</li> <li>4. Declaration for Latest Non Conviction, non blacklisting on non judicial Stamp paper of Rs 100 furnished by the bidder as per proforma duly notarised.</li> </ol>	<p>Minor Infirmities not submitted</p>	<p>Not Recommend ed to the Technical Advisory/ Evaluation Committee</p>

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5. Declaration for Latest Non Conviction, non blacklisting on non judicial Stamp paper of Rs 100 furnished by the Principal Manufacturer as per proforma duly notarised.
6. Resubmit Client Base of the quoted model/ make with reference of the supply orders, for any of three financial years in Last five years along with satisfactory performance certificate of minimum one purchase order and installation report of Govt, Institution (Copies of reference supply orders and satisfactory performance certificate need to be attached)
7. To submit minimum of 5 years' experience in the healthcare sector and also submit documentary proof of having supplied at least 10 units of the same quoted model in Government institutions for TB screening.
8. Average Annual Turnover Statement for the last 3 financial years (2021-22, 2022-23, and 2023-24) of the Principal Manufacturer, issued by a Chartered Accountant/competent authority with UDIN. (The bidder has submitted its own turnover statement, which is not acceptable in this case.)
9. Copies of audited Balance Sheet and Profit & Loss Account for the last three financial years (2021-22, 2022-23, and 2023-24) of the Principal Manufacturer, certified by a Chartered Accountant with UDIN. (The bidder has submitted its own audit balance sheets which is not acceptable in this case.)
10. For X-ray Device Certification submit BIS standard IS 7620 (Part 1):1986 or shall have valid Electrical Safety Approval or US FDA / CE certification, as applicable
11. For ICMR Approval submit ICMR field approval
12. For NAAT Approval submit NAAT approval for use in TB screening applications
13. For AERB Approval of the principal manufacturer/bidder submit a valid AERB approval certificate for the quoted equipment
14. For AI Software Certification submit CE or US FDA approved Artificial Intelligence (AI) software.
15. For AI Software Licensing submit a valid CDSCO license
16. submit HIPAA and/or EU GDPR standards for data protection and privacy
17. Submit ISO 27001 (Information Security Management System) and ISO 13485:2016 (Quality Management System for Medical Devices).

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It was apprised that the Power Point Presentation before the experts was held on 19.12.2025, wherein the technical experts were of the opinion that PPT demonstration of both the bidders was taken in which certain points were not cleared with ambiguity regarding tube current, drop test including physical demonstration and to have fair selection of the system it was decided to have re-demonstration & physical drop test.

Accordingly, the Physical Demonstration of the equipments was conducted 23.12.2025 by the technical experts and on the basis of Demonstration, catalogue evaluation & certifications as per the conditions of the NIT, the following bidders were shortlisted and declared "**qualified for opening of financial bid**" by the technical experts with the remarks that "After detailed evaluation, we find product of Findoc Ventures Private Limited qualifies the technical Evaluation. The product of Surgident India does not qualify due to fixed tube current, fixed KVP and poor spatial resolution.

S. No	Name of the item	Bidder qualified for PPT	Bidders qualified by the technical experts	Bidders Dis-qualified by the technical experts
1.	Ultraportable Handheld Digital X-Ray KIT (with AI Technology Based Software for TB Screening	M/s Findoc Ventures Private Limited Gurugram Haryana	M/s Findoc Ventures Private Limited Gurugram Haryana	M/s Surgident India
		M/s Surgident India		

**Decision:** Accordingly, the Technical Advisory /Evaluation Committee accepted the report of the technical expert committee and recommended opening of "**financial bid of the qualified bidders**" after observing all codal formalities for final approval.

After the opening of financial bid of single bid qualified items, the committee decided that before placing such equipments in the Purchase Committee, JKSMCL, price justification/rate reasonability with other premier institutes be also done so that price justification is ensured.

S. No.	Name of the Officer	Designation	Signatures
1.	Mohd. Ashraf Choudhary,	General Manager (Adm)	✓
2.	Dr Upali Vagno	mo QC	upali
3.	Dr. Raji Ramnani	Dy. General Manager (Phs) Korba	✓
4.	Chander kiran.	Asst, Manager	Chander

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5.	Karan Singh	AAO	o
6.	Dr Pankaj koul	Consultant. Radiologist Govt. Hospital Srinagar	pankaj
7.	Vishal Mahajan	MHD JEN. JAMMU.	Mahajan
8.	Ambar Kalra.	FM, DJe	Ambar
9.	Dr. Archad. Bhat.	Consultant Radiologist- SMGS. Hospital Jammu	Archad
10	Sharminder Singh	Asst. o, 09999999999	Sharminder

(Mohd. Ashraf Choudhary) JKAS  
General Manager (Adm.),

J&K Medical Supplies Corporation Ltd

Dated: 03/12/2025

No.: JKMSCL/GM (Adm.)/2025/ 3967-76

Copy for information to the:-

1. Principal, Govt. Medical College, Srinagar (Kashmir).
2. Principal, Government Medical College, Jammu.
3. Principal, Government Dental College, Srinagar.
4. Principal, Government Dental College, Jammu.
5. Managing Director, Jammu and Kashmir Medical Supplies Corporation Limited, Jammu
6. Director, Health Services, Jammu.
7. Director, Health Services, Kashmir.
8. Controller, Drug & Food Control Organization, J&K.
9. F.A/Chief Accounts Officer, J&K Medical Supplies Corporation Limited.
10. Dy. General Manager (Tendering), J&K Medical Supplies Corporation Limited
11. Office Record.