



JAMMU AND KASHMIR MEDICAL SUPPLIES CORPORATION LTD.

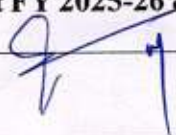
(Public Sector Undertaking of the Government of Jammu and Kashmir)
Corporate Head Office: Plot No. 58, Friends Colony Satyam Road Trikuta Nagar Jammu
Corporate Office: Opposite J&K Motor Garage Deptt near Hajj House Bemina Srinagar **Telefax:** 0194-2432008
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NOTICE

Subject: Evaluation of Technical bids for finalization of Rate Contract for Procurement of Cardiac Cath Lab and allied equipments under Machinery & Equipments No. NIT/JKMSCL/M&E/2026/688 dated: 13-04-2026.

With reference to the report of the Sub-committee, the evaluation of bid documents for the finalization of the rate contract for the procurement **Cardiac Cath Lab and allied equipments under Group Machinery & Equipments** with reference to the **No. NIT/JKMSCL/M&E/2026/688 dated: 13-04-2026.** the evaluation of the bid documents was done on the basis of uploaded documents and infirmities submitted by the bidder :-

Cardiac Cath Lab		
S.No	Name of firm/bidder	Shortcomings/Infirmities
1/2	M/s Erbis Engineering Co. Ltd., New Delhi (Bidder/ Importer) M/s Canon Medical Systems Corp./Canon Inc., Japan ? (Manufacturer) Model: Alphenix Core+ (INFX-8000V/Z1)	To Submit: 1. Compliance Sheet for each equipment (self-attested) (Bidder has to submit an affidavit that the quoted model is in compliance with all the specifications required as per NIT 688 of JKMSCL along with subsequent corrigendum with no deviation in any aspect.) 2. Declaration for Latest Non-Conviction, non-blacklisting on non-judicial Stamp paper of Rs 100 furnished by the Principal Manufacturer / Sole Importer/ Indian Subsidiary as per proforma duly notarised. (Annexure IV B) 3. Self-attested photocopy of IEC certificate and Permission/ Authorization for sale for sale from the foreign principal manufacturer (in case of imported product) Annexure A 4. Specify point of supply with full Address. Annexure G Note: As per list of quoted items & Authorization letter, Manufacturer is M/s Canon Inc. Japan whereas as per the CDSCO & quality certificates Manufacturer is M/s Canon Medical Systems Corporation, Japan. Clarify and submit all relevant documents accordingly
2/2	M/s S R Technomed, Jammu (Bidder) M/s Phillips India Ltd. Delhi (Importer) M/s Phillips Medical	To Submit: 1. Copy of Catalogue of the Quoted product (self-attested) (Submitted document is of Azurion 7 C12 & F12, whereas the quoted model is Azurion 7 M12) (Clarify & Resubmit) 2. Compliance Sheet for each equipment (self-attested) (Bidder has submitted document with deviation in compliance w.r.t Serial. No. B (03) & C (04) of the Technical specifications required for the quoted item as per NIT 688 of JKMSCL, which cannot be considered) (Note: Bidder has to submit an affidavit that the quoted model is in compliance with all the specifications required as per NIT 688 of JKMSCL along

	Systems, Netherland (Foreign Manufacturer) Model: Azurion7 M12	with subsequent corrigendum with no deviation in any aspect) Client Base on Letter Head of the Bidder / Manufacture/Indian Subsidiary of Principal Manufacturer with references of the supply orders, for any of the one financial years in last five financial years except FY 2023-24 & 2025-26 with at least one satisfactory performance of submitted Government Purchase Order for Cath Lab (Annexure V) - Authorization from principal manufacturer / Importer (On the letter head of Principal manufacturer / Sole Importer) In case authorization to the bidder is furnished by the Sole Importer/Indian Subsidiary, document confirming authorization from foreign Principal Manufacturer in favour of Indian Subsidiary / Sole Importer is to be submitted (strictly as per annexure VI) - Copies of Audited Balance sheet & profit loss account for last three financial years certified by Chartered Accountant of the Importer/ Indian Subsidiary for (2022-23, 2023-24 and 2024-25) with UDIN. In case of foreign manufacturer, the balance sheets of Indian Subsidiary/Sole Importer only shall be considered (Submitted document is without UDIN) - Self-attested photocopy of IEC certificate and Permission/ Authorization for sale for sale from the foreign principal manufacturer (in case of imported product) Annexure A - Quality Certifications on the products viz. ISI/CE/USFDA etc. whichever applicable. (Submitted CE Certificate has expired on 25.06.2025) - Valid MD-42 or equivalent document of the bidder Service escalation matrix (As asked at Serial 07 under Section VI B: - Special Conditions of Contract (SCC) on Page 60 of NIT 688 of JKMSCL) - Declaration of Availability of spare parts and consumables for the quoted equipment for at least 10 years/life of the item, from the date of installation.(As asked at 2.6 Section VI A: - General Conditions of Contract (GCC) on Page 42 of NIT 688 of JKMSCL) - Declaration of 5 years comprehensive warranty on labour and all supplied items (As asked at J.1 on Page 80 of NIT 688 of JKMSCL) Note: 1. Bidder has submitted letter under Cover-A (Technical Bid) with warranty & payment terms along with other conditions which are not in accordance with Terms & Conditions of the NIT 688 of JKMSCL. Clarify (T&C beyond the scope of NIT 688 of JKMSCL are not acceptable)
Rotablator		
1/1	M/S AI –Med Agencies, Srinagar (Bidder) M/s Boston Scientific	To submit: Client Base on Letter Head of the Bidder / Manufacture/Indian Subsidiary of Principal Manufacturer with references of the supply orders, for any of the one financial years in last five financial years except FY 2025-26 & 2026-27 with at least one 

	India (P) Ltd., Gurgaon (Importer) M/s Boston Scientific, USA (Foreign Manufacturer) Model: ROTAPRO Console Kit Assy Gen 230V ROTAGO™ Cart ASSY Power Cord	satisfactory performance of submitted Government Purchase Order for the quoted item 2. Authorization from principal manufacturer / Importer (On the letter head of Principal manufacturer / Sole Importer) In case authorization to the bidder is furnished by the Sole Importer/Indian Subsidiary, document confirming authorization from foreign Principal Manufacturer in favour of Indian Subsidiary / Sole Importer is to be submitted (strictly as per annexure VI) 3. Copies of Audited Balance sheet & profit loss account for last three financial years certified by Chartered Accountant of the Importer/ Indian Subsidiary for (2022-23, 2023-24 and 2024-25) with UDIN. In case of foreign manufacturer, the balance sheets of Indian Subsidiary/Sole Importer only shall be considered (Submitted balance sheets are without UDIN) 4. Self-attested photocopy of IEC certificate and Permission/ Authorization for sale for sale from the foreign principal manufacturer (in case of imported product) Annexure A 5. Valid MD-42 or equivalent document of the bidder
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Intra Aortic Balloon Pump

1/1	M/S S R Technomed, Jammu (Bidder) M/s Teleflex Medical (P) Ltd., Karnataka (Importer) M/s Arrow International LLC (Subsidiary of Teleflex Inc. USA) (Foreign Manufacturer) Model: AC3 Optimus	To Submit: 1. List of Items quoted by the Bidder mentioning name of manufacturer/ importer with make & model as per annexure. Annexure I (As per document submitted, manufacturer for the quoted item is M/s Teleflex Inc. USA whereas as per CDSCO Manufacturer is M/s Arrow International LLC) (Clarify & submit all relevant documents accordingly) 2. Compliance Sheet for each equipment (self-attested) (Bidder has submitted document with deviation in compliance w.r.t minimum battery backup of 2hrs & alarms, therapy status & history report under Basic Features & Clinical Functionality, respectively of the Technical specifications required for the quoted item as per NIT 688 of JKMSCL, which cannot be considered) (Note: Bidder has to submit an affidavit that the quoted model is in compliance with all the specifications required as per NIT 688 of JKMSCL along with subsequent corrigendum with no deviation in any aspect) 3. Declaration for Latest Non-Conviction, non blacklisting on non-judicial Stamp paper of Rs 100 furnished by the Sole Importer/ Indian Subsidiary as per proforma duly notarised. Annexure IV B (Clarify relation between M/s Teleflex Medical (P) Ltd., Karnataka & M/s Smartwork Co working space) 4. Copies of Audited Balance sheet & profit loss account for last three financial years certified by Chartered Accountant of the Importer/ Indian Subsidiary for (2022-23, 2023-24 and 2024-25) with UDIN. In case of foreign manufacturer, the balance sheets of Indian Subsidiary/Sole Importer only shall be considered (Submitted balance sheet of 2024-25 is without UDIN) 5. Self-attested photocopy of IEC certificate and Permission/ Authorization for sale for sale from the foreign principal manufacturer (in case of imported product) Annexure A
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		6. Copy of CDSCO Registration Certification, wherever applicable. (As per submitted MD 15, legal manufacturer is M/s Manufacturer is M/s Arrow International LLC USA for the quoted item, whereas as per list of quoted items Teleflex Inc. USA is the Principal Manufacturer for the quoted item. Clarify & submit all relevant documents accordingly) 7. Valid MD-42 or equivalent document of the bidder 8. Service escalation matrix (As asked at Serial 07 under Section VI B: - Special Conditions of Contract (SCC) on Page 60 of NIT 688 of JKMSCL) 9. Declaration of Availability of spare parts and consumables for the quoted equipment for at least 10 years/life of the item, from the date of installation. (As asked at 2.6 Section VI A: - General Conditions of Contract (GCC) on Page 42 of NIT 688 of JKMSCL) 10. Declaration of 5 years comprehensive warranty on labour and all supplied items (As asked at J.1 on Page 80 of NIT 688 of JKMSCL) Note: 1. Bidder has submitted letter under subject "Offered Equipment List for Cardiac Cath Lab dated 15.08.2026, wherein make & model for the quoted item is CARDIOSAVE HYBRID Automatic IABP-5Y, IABP of Balloon-Linear 34CC, IABP of Balloon-Linear 40CC whereas as per list of quoted items, make & model are Teleflex & AC3 Optimus respectively. Clarify and submit all relevant documents accordingly. 2. Bidder has submitted letter under Cover-A (Technical Bid) with warranty & payment terms along with other conditions which are not in accordance with Terms & Conditions of the NIT 688 of JKMSCL. Clarify (T&C beyond the scope of NIT 688 of JKMSCL are not acceptable)
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Intra Vesicular Ultrasound System (IVUS)

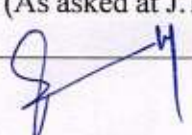
1/2	M/S AI –Med Agencies, Srinagar (Bidder) M/s Boston Scientific India (P) Ltd., Gurgaon (Importer) M/s Boston Scientific, USA (Foreign Manufacturer) Model: Avvigo+ Latest Generation AI enabled IVUS	To Submit: 1. Copy of Catalogue of the Quoted product (Submitted document is for Avvigo+ multi modality guidance system instead of Avvigo+ Latest Generation AI enabled IVUS & also not self attested) (Clarify & resubmit the self attested document) 2. Compliance Sheet for each equipment (self-attested) (Bidder has submitted document with deviation in compliance w.r.t Serial. No. 04 & 20 of the Technical specifications required for the quoted item as per NIT 688 of JKMSCL, which cannot be considered) (Note: Bidder has to submit an affidavit that the quoted model is in compliance with all the specifications required as per NIT 688 of JKMSCL along with subsequent corrigendum with no deviation in any aspect) 3. List of Items quoted by the Bidder mentioning name of manufacturer/ importer with make & model as per annexure. Annexure I (As per the document submitted the quoted model is Avvigo+ Latest Generation AI enabled IVUS whereas as per catalogue & USFDA certificate, the model is
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		Avvigo+ multi modality guidance system. Clarify & submit all the relevant documents) Client Base on Letter Head of the Bidder / Manufacture/Indian Subsidiary of Principal Manufacturer with references of the supply orders, for any of the one financial years in last five financial years except FY 2024-25 & 2025-26 with at least one satisfactory performance of submitted Government Purchase Order for the quoted item (Annexure V) - Authorization from principal manufacturer / Importer (On the letter head of Principal manufacturer / Sole Importer) In case authorization to the bidder is furnished by the Sole Importer/Indian Subsidiary, document confirming authorization from foreign Principal Manufacturer in favour of Indian Subsidiary / Sole Importer is to be submitted (strictly as per annexure VI) - Copies of Audited Balance sheet & profit loss account for last three financial years certified by Chartered Accountant of the Importer/ Indian Subsidiary for (2022-23, 2023-24 and 2024-25) with UDIN. In case of foreign manufacturer, the balance sheets of Indian Subsidiary/Sole Importer only shall be considered (Submitted balance sheets are without UDIN) - Self-attested photocopy of IEC certificate and Permission/ Authorization for sale for sale from the foreign principal manufacturer (in case of imported product) Annexure A - Quality Certifications on the products viz. ISI/CE/USFDA etc. (As per the list of quoted items, model is Avvigo+ Latest Generation AI enabled IVUS whereas as per USFDA certificate, the model is Avvigo+ multi modality guidance system. Clarify & submit all the relevant documents) - Copy of CDSCO Registration Certification (Submitted document is Market Standing Certificate only) (Submit Valid MD-15 certificate for the quoted item) - Valid MD-42 or equivalent document of the bidder
2/2	M/s S R Technomed, Jammu (Bidder) M/s Phillips India Ltd. (Importer) M/s Phillips Medical Systems, Nederland (Manufacturer) Model: Intra Sight 5	To Submit: - Authorization from principal manufacturer / Importer (On the letter head of Principal manufacturer / Sole Importer) In case authorization to the bidder is furnished by the Sole Importer/Indian Subsidiary, document confirming authorization from foreign Principal Manufacturer in favour of Indian Subsidiary / Sole Importer is to be submitted (strictly as per annexure VI) - Self-attested photocopy of IEC certificate and Permission/ Authorization for sale for sale from the foreign principal manufacturer (in case of imported product) Annexure A - Copies of Audited Balance sheet & profit loss account for last three financial years certified by Chartered Accountant of the Importer/ Indian Subsidiary for (2022-23, 2023-24 and 2024-25) with UDIN. In case of foreign manufacturer, the balance sheets of Indian Subsidiary/Sole Importer only shall be considered (Submitted document is without UDIN) - Quality Certifications on the products viz. ISI/CE/USFDA etc. whichever applicable. (Submitted USFDA certificate in favour of M/s Volcano Corporation for Intrasight system)

		whereas as per list of quoted items M/s Phillips Medical Systems Nederland is the Principal Manufacturer for the model Intrasight 5. Clarify the relation between two & submit all relevant documents accordingly) - Copy of CDSCO Registration Certification, wherever applicable. (As per submitted MD 15, legal manufacturer is M/s Volcano Corporation, California US for the quoted item, whereas as per list of quoted items M/s Phillips Medical Systems Nederland is the Principal Manufacturer for the model Intrasight 5. Clarify the relation between two & submit all relevant documents accordingly) - Valid MD-42 or equivalent document of the bidder Service escalation matrix (As asked at Serial 07 under Section VI B: - Special Conditions of Contract (SCC) on Page 60 of NIT 688 of JKMSCL) - Declaration of Availability of spare parts and consumables for the quoted equipment for at least 10 years/life of the item, from the date of installation.(As asked at 2.6 Section VI A: - General Conditions of Contract (GCC) on Page 42 of NIT 688 of JKMSCL) - Declaration of 5 years comprehensive warranty on labour and all supplied items (As asked at J.1 on Page 80 of NIT 688 of JKMSCL) Bidder has to submit an affidavit that the quoted model is in compliance with all the specifications required as per NIT 688 of JKMSCL along with subsequent corrigendum with no deviation in any aspect Note: - Bidder has submitted letter under Cover-A (Technical Bid) with warranty & payment terms along with other conditions which are not in accordance with Terms & Conditions of the NIT 688 of JKMSCL. Clarify (T&C beyond the scope of NIT 688 of JKMSCL are not acceptable)
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ECHO Machine

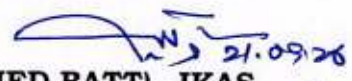
1/2	M/s Erbis Engineering Co. Ltd., New Delhi (Bidder/ Importer) M/s Canon Medical Systems Corp./Canon Inc., Japan (Foreign Manufacturer) Model: APLIO ME (CUS-AME00)	- Compliance Sheet for each equipment (self-attested) (Bidder has submitted document with deviation in compliance w.r.t Serial. No. 08, 10, 11, 19, 20 & 21 of the Technical specifications required for the quoted item as per NIT 688 of JKMSCL, which cannot be considered) (Note: Bidder has to submit an affidavit that the quoted model is in compliance with all the specifications required as per NIT 688 of JKMSCL along with subsequent corrigendum with no deviation in any aspect) - Declaration for Latest Non-Conviction, non-blacklisting on non-judicial Stamp paper of Rs 100 furnished by the Principal Manufacturer / Sole Importer/ Indian Subsidiary as per proforma duly notarised. (Annexure IV B) Client Base on Letter Head of the Bidder / Manufacture/Indian Subsidiary of Principal Manufacturer with references of the supply orders, for any of the two financial years in last five financial years except FY 2025-26 with at least One Government Purchase order along with along its satisfactory performance (Copies of reference supply orders
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		and satisfactory performance certificate need to be attached) . Annexure V 4. Self-attested photocopy of IEC certificate and Permission/ Authorization for sale for sale from the foreign principal manufacturer (in case of imported product) Annexure A 5. Specify point of supply with full Address. Annexure G Note: As per list of quoted items & Authorization letter, Manufacturer is M/s Canon Inc. Japan whereas as per the CDSCO & quality certificates Manufacturer is M/s Canon Medical Systems Corporation Japan. Clarify and submit all relevant documents accordingly.
2/2	M/s S R Technomed, Jammu (Bidder) M/s Phillips India Ltd., Pune (Manufacturer) Model: Affiniti 50	To Submit: 1. Bidder has to submit an affidavit that the quoted model is in compliance with all the specifications required as per NIT 688 of JKMSCL along with subsequent corrigendum with no deviation in any aspect 2. 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. Annexure IV B (As M/s Phillips India is the Manufacturer for the quoted item, resubmit Annexure IV B as OEM) 3. Satisfactory Performance Certificate for any one of the submitted Purchase order from Government Institution for the quoted item 4. Authorization from principal manufacturer (On the letter head of Principal manufacturer) (Bidder has to submit Authorization from M/s Phillips India as Principal Manufacturer to M/s S R Technomed Jammu for the quoted item strictly as per annexure VI) 5. Copies of Audited Balance sheet & profit loss account for last three financial years certified by Chartered Accountant of the Importer/ Indian Subsidiary for (2022-23, 2023-24 and 2024-25) with UDIN. In case of foreign manufacturer, the balance sheets of Indian Subsidiary/Sole Importer only shall be considered (Submitted document is without UDIN) 6. Quality Certifications on the products viz. ISI/CE/USFDA etc. whichever applicable. (Submitted USFDA certificate in favour of M/s Philips Ultrasound Inc. for the quoted item whereas as per list of quoted items M/s Phillips India is the Principal Manufacturer. Clarify & submit all relevant documents) 7. Valid MD-42 or equivalent document of the bidder 8. Service escalation matrix (As asked at Serial 07 under Section VI B: - Special Conditions of Contract (SCC) on Page 60 of NIT 688 of JKMSCL) 9. Declaration of Availability of spare parts and consumables for the quoted equipment for at least 10 years/life of the item, from the date of installation.(As asked at 2.6 Section VI A: - General Conditions of Contract (GCC) on Page 42 of NIT 688 of JKMSCL) 10. Declaration of 5 years comprehensive warranty on labour and all supplied items (As asked at J.1 on Page 80 of NIT 688 of JKMSCL) 

		Note: 1. Bidder has submitted letter under Cover-A (Technical Bid) with warranty & payment terms along with other conditions which are not in accordance with Terms & Conditions of the NIT 688 of JKMSCL. Clarify (T&C beyond the scope of NIT 688 of JKMSCL are not acceptable)
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The bidder(s) are requested to upload their infirmities within 07 days from the issuance of this notice on the email of tender inviting authority **gmjkmscl.mil@gmail.com** which shall be forwarded to technical evaluation sub-committee for scrutiny and recommendations in the matter.

In case the documents are not submitted in time, your bid shall be evaluated only on the basis of documents submitted earlier.


(WAKEEL AHMED BATT), JKAS
General Manager (Adm.), JKMSCL

No. NIT/JKMSCL/GM (ADM)/2026/4832-34

Dated: 21/09/2026.

Copy to the:

1. Dy. General Manager (Tendering) JKMSCL.
2. Assistant Programmer to upload the minor infirmities on the portal.
3. PA to the Managing Director, JKMSCL
4. Office Copy