JAWAHARLAL INSTITUTE OF POST GRADUATE MEDICAL EDUCATION & RESEARCH (JIPMER)

GOVERNMENT OF INDIA

(An Institution of National Importance under Ministry of Health & Family Welfare) Dhanvantri Nagar, Puducherry–605006

OPEN TENDER
ENQUIRY DOCUMENT
FOR
Rate Contract
For
Supply of Consumables/Non-Consumables
for
Department of
Cardiology

(This document consisting of 45 pages)

JIPMER,
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I N D E X

SECTION-I

JAWAHARLAL INSTITUTE OF POSTGRADUATE MEDICAL EDUCATION AND RESEARCH PUDUCHERRY – 6
(Institution of National Importance under the Ministry of Health & Family Welfare, Govt. of India)
Purchase Section

No. JIP/PUR6(1)/OeT/Cardio/2021-22 Dated: 24.07.2021

NOTICE INVITING e-TENDER (e-NIT)

Sub: Supply of Consumables / Non-Consumables

E- tenders are invited from eligible and qualified domestic manufacturer's or their distributors for the RATE CONTRACT for supply of CONSUMABLES/NON-CONSUMABLES for the Department of **Cardiology** for 2021– 22.

1. Scope of work: Supply of consumables/non-consumables for the Department of Surgery
2. Value of tender: Approximate Cost **Rs. 1,00,00,000**
3. Tender timelines:
   - **i.** Opening date & time for download of Tender document: **06.00 PM** on **24.07.2021**
   - **ii.** Last date for receipt of pre-bid queries: **06.00 PM** on **29.07.2021**
   - **iii.** Pre-bid queries can be made through e-mail to**:** jipmercardiologyoffice@gmail.com
   - **iv.** Opening date & time for submission of online bids: **09.00 AM on 03.08.2021**
   - **v.** Closing date & time for submission of online bids **06.00 PM on 24.08.2021**
   - **vi.** Date and Time of opening of online bids for Technical evaluation: **11.00 AM** on **26.08.2021**
   - **vii.** Date & time of opening of Price Bid: To be intimated later.

4. Earnest Money Deposit/Bid Security: Bid security declaration must be submitted by all bidders.

5. Tender Processing Fee (Non-refundable): **Rs. 590 (including 18% GST) shall be paid through SBI collect only.**

6. Interested bidders are advised to download the complete Tender Enquiry document from the websites **www.jipmer.edu.in** or **https://eprocure.gov.in/eprocure/app** for complete details.
7. The prospective bidders must register with the E-procurement system of **https://eprocure.gov.in/eprocure/app**. Special Instructions to the bidders for the e-submission of the bids online through this eProcurement Portal on completion of the registration process is given in **https://eprocure.gov.in/eprocure/app ,** the bidders will be provided user ID and password upon enrollment. In order to submit the bids electronically, bidders are required to have a valid Class 3 Digital Signature Certificate (signing and encryption/ decryption certificates).

8. Bidders are requested to read the bidders help document on e-tender web site link before proceeding for bidding.
9. Post receipt of User ID & Password, Bidders can log on for downloading & uploading tender document.
10. The bidders shall submit the required bid security declaration (as per G.I.T clause 2) before the due date and time mentioned above.
11. The online submission of tender(s) can only be done through **https://eprocure.gov.in/eprocure/app**
12. Bidders shall ensure that their tender(s), complete in all respects, are submitted online through **https://eprocure.gov.in/eprocure/app** e-portal (as described above) only.
13. Prospective bidders are advised to browse the above websites regularly before submission of their bids as any further amendments will be published in these websites only.

Asst. Officer in Charge
Purchase Section,
For
Director,
JIPMER, Puducherry

Section-II

SCHEDULE OF REQUIREMENTS AND SPECIFICATIONS

LIST OF ITEMS REQUIRED

1. Scope of work: Supply of Consumables/ Non-Consumables for the department of **Cardiology** as detailed below

Sl. No.	Item Description	Qty.	Units
1.001	(S) FEMORAL PCI SHEATH SYSTEMS SIZE 4F with accessories for insertion. Should be FDA'/CE'/DCGI approved	100	Nos
1.002	(S) FEMORAL PCI SHEATH SYSTEMS SIZES 5F-9F with accessories for insertion. Should be FDA'/CE'/DCGI approved	1500	Nos
1.003	(S) FEMORAL PCI SHEATH SYSTEMS SIZES 10, 12, 14 F with accessories for insertion. Should be FDA'/CE'/DCGI approved	100	Nos
1.004	(S)VASCULAR CLOSURE DEVICES Femoral all sizes. Should be FDA'/CE'/DCGI approved	50	Nos
1.005	(S) TRANSRADIAL SHEATH SYSTEMS with hydrophilic coating on sheath, tapering dilator hydrophilic staright polymer 025 or 021 wires and 20G plastic cannula for puncture 5F, 6F,7F. Should be FDA'/CE'/DCGI approved	1500	Nos
1.006	(S) RADIAL ANGIO SET WITH PUNCTURE NEEDLE, SHEATH 6F, STRAIGHT GUIDEWIRE 025/021 AND TIG and TIG II CATHETER 5F, AND HYDROPHILIC POLYMER J tipped 035 145 cm WIRES. Should be FDA'/CE'/DCGI approved	1500	Nos
1.007	(S) RADIAL ANGIOGRAPHY KITS Comprising the following Items a.Radial Sheath Insertion System 6F b. Radial Angiography 5F TIG catheter c. 0.035 J Hydrophilic polymer wire, d.Disposable Sterile Drape for Radial/Femoral angiography e.Disposable Sterile Gowns -2, f.Disposable Gloves-2, g.Finger control syringe 20cc h.3/3 Rotating manifold i.pressure line j.100 mL Non Ionic contrast media. Should be FDA'/CE'/DCGI approved	500	Nos
1.008	(S)Flexible delivery sheaths with steerable and defelctable introducer systems (60 cm and above long). Should be CE' AND DCGI approved	50	Nos
1.009	(S)EXTRALONG (more than 150 cm) pigtail catheters for illiac angiography from radial . Should be FDA'/CE'/DCGI approved	20	Nos
1.01	(S)Prolene sutured femoral access closure devices. Should be FDA' or CE' andDCGI approved	50	Nos
1.011	(S)Radial Compression Devices. Should be FDA'/CE'/DCGI approved	10	Nos
1.012	(S) COURNAND CATHETER 5F, 6F woven dacron. Should be FDA'/CE'/DCGI approved	50	Nos
1.013	(S) PIG TAIL CATHETERS 5F, 6F. Should be FDA'/CE'/DCGI approved	100	Nos
1.014	(S) MARKER PIGTAIL CATHETER 5F, 6F. Should be FDA'/ CE'/ DCGI approved	50	Nos
1.015	(S)JUDKINS LEFT CATHETER 4F,5F,6F all Curves. Should be FDA'/CE'/DCGI approved	200	Nos
1.016	(S)JUDKINS RIGHT CATHETER 4F,5F,6F all Curves. Should be FDA'/CE'/DCGI approved	200	Nos
1.017	(S)AMPLATZ LEFT CATHETER 4F, 5F,6F all Curves. Should be FDA'/CE'/DCGI approved	25	Nos
1.018	(S)AMPLATZ RIGHT CATHETER 4F,5F,6F all Curves. Should be FDA'/CE'/DCGI approved	25	Nos
1.019	(S)LEFT CORONARY BY PASS CATHETER 4F,5F,6F all Curves. Should be FDA'/CE'/DCGI approved	10	Nos

1.02	(S)RIGHT CORONARY BY PASS CATHETER 4F,5F,6F all Curves. Should be FDA'/CE'/DCGI approved	10	Nos
1.021	(S)INTERNAL MAMMARY ARTERY CATHETER 4F, 5F,6F all Curves. Should be FDA'/CE'/DCGI approved	25	Nos
1.022	(S)MULTIPURPOSE CATHETER A1,A2,B1,B2. Should be FDA'/ CE'/ DCGI approved	50	Nos
1.023	(S)PIG-TAIL CATHETER PEDIATRIC 4 F. Should be FDA'/CE'/DCGI approved	10	Nos
1.024	(S)RADIAL CAG TIG CATHETERS 5F ALL Curves. Should be FDA'/CE'/DCGI approved	1200	Nos
1.025	(S) 2/3 ROTATING MANIFOLD FOR CAG. Should be FDA'/ CE'/ DCGI approved	1000	Nos
1.026	(S) 3/3 ROTATING MANIFOLD FOR CAG/PTCA. Should be FDA'/ CE'/ DCGI approved	1000	Nos
1.027	(S) TOUHEY BORST Y CONNECTOR- PUSH PULL TYPE and INTRODUCER NEEDLE for PTCA. Should be FDA'/CE'/DCGI approved	1000	Nos
1.028	(S) Y CONNECTOR 10F Diameter having larger inner lumen with INTRODUCER NEEDLE for PTCA. Should be FDA'/CE'/DCGI approved	250	Nos
1.029	(S) 3 RING CONTROL SYRINGES 20cc. Should be FDA'/CE'/DCGI approved	1000	Nos
1.03	(S)TEFLON COATED GUIDE WIRE145 CM 0.021" ST. & J. Should be FDA'/CE'/DCGI approved	50	Nos
1.031	(S)TEFLON COATED GUIDE WIRE 145 CM 0.025" ST'& J. Should be FDA'/CE'/DCGI approved	50	Nos
1.032	(S)TEFLON COATED GUIDE WIRES 0.032" 145 CM ST& J. Should be FDA'/CE'/DCGI approved	1000	Nos
1.033	(S)TEFLON COATED GUIDE WIRES 145 CMS 0.035"ST & J. Should be FDA'/CE'/DCGI approved	1000	Nos
1.034	(S)TEFLON COATED GUIDE WIRES 145 CMS 0.038" ST. Should be FDA'/CE'/DCGI approved	500	Nos
1.035	(S)Exchange length (260cm and above) 0.018" peripheral guide wire . Should be FDA'/CE'/DCGI approved	50	Nos
1.036	(S)Lunderquist type high support exchange length 0.035" (260cm and above) short and long tip guide wire . Should be FDA'/CE'/DCGI approved	50	Nos
1.037	(S)TEFLON COATED EXCHANGEGUIDE WIRE 260 CM 0.032"ST. & J. Should be FDA'/CE'/DCGI approved	100	Nos
1.038	(S)TEFLON COATED EXCHANGEGUIDE WIRE 260 CM 0.035"ST. & J. Should be FDA'/CE'/DCGI approved	100	Nos
1.039	(S)TEFLON COATED EXCHANGEGUIDE WIRE 260 CM 0.038"ST& J. Should be FDA'/CE'/DCGI approved	100	Nos
1.04	(S)TERUMO GUIDE WIRES 145 CM 018, 021.032,035 ST AND J ALL SIZES. Should be FDA'/CE'/DCGI approved	500	Nos
1.041	(S)TERUMO EXCHANGE GUIDE WIRES 018, 021.032,035 ST AND J ALL SIZES. Should be FDA'/CE'/DCGI approved	200	Nos
1.042	(S)TERUMO STIFF EXCHANGE GUIDE WIRES ALL SIZES. Should be FDA'/CE'/DCGI approved	500	Nos
1.043	(S)AMPLATZ TYPE SUPERSTIFF EXCHANGE GUIDE WIRES 0.035' AND .038". 260 cm and above- Short floppy tip. Should be FDA'/CE'/DCGI approved	50	Nos
1.044	(S)AMPLATZ ULTRA STIFF GUIDE WIRES 0.035' AND 0.038" J. 260 cm and above- LONG floppy tip. Should be FDA'/CE'/DCGI approved	50	Nos
1.045	(EX)FEMORAL ANGIOPLASTY GUIDING CATHETER SYSTEMS 5F,6F,7F,8F ALL SHAPES AND SIZES. Should be FDA'/CE'/DCGI approved	1200	Nos
1.046	(EX)RADIAL ANGIOPLASTY -GUIDING CATHETER SYSTEMS 4F ALL SHAPES AND SIZES. Should be FDA'/CE'/DCGI approved	25	Nos
1.047	(EX)RADIAL ANGIOPLASTY -GUIDING CATHETER SYSTEMS 5F,6F,7F	500	Nos

	ALL SHAPES AND SIZES. Should be FDA'/CE'/DCGI approved		
1.048	(S)PTCA WIRES Floppy standard length . Should be FDA'/CE'/DCGI approved	300	Nos
1.049	(S)PTCA WIRES-Floppy exchange length. Should be FDA'/CE'/DCGI approved	300	Nos
1.05	(S) PTCA WIRES Intermediate Standard Length. Should be FDA'/CE'/DCGI approved	300	Nos
1.051	(S) PTCA WIRES Extra support Standard Length. Should be FDA'/CE'/DCGI approved	300	Nos
1.052	(S) PTCA DOC LINK EXTENSION WIRES. Should be FDA'/CE'/DCGI approved	300	Nos
1.053	(S)EXCHANGE WIRES FOR PTCA . Should be FDA'/CE'/DCGI approved	250	Nos
1.054	(S)Double core spring tip TYPE WIRES FOR PTCA. Should be FDA'/CE'/DCGI approved	300	Nos
1.055	(S)PTCA WIRES Tapering Tip Hydrophilic. Should be FDA'/CE'/DCGI approved	250	Nos
1.056	(S)PTCA WIRE -TOTAL OCCLUSION WIRES-PLATINUM. Should be FDA'/CE'/DCGI approved	250	Nos
1.057	(S)PTCA WIRES for CTO-hydrophlic . Should be FDA'/CE'/DCGI approved	250	Nos
1.058	(S)PTCA WIRES for CTO-hydrophlic Extra support. Should be FDA'/CE'/DCGI approved	250	Nos
1.059	(S)PTCA WIRES Polymer Coated Hydrophilic. Should be FDA'/CE'/DCGI approved	250	Nos
1.06	(S)PTCA WIRES Polymer Coated Hydrophilic Extra support. Should be FDA'/CE'/DCGI approved	250	Nos
1.061	(S)PTCA WIRES- Polymer coated tapering tip. Should be FDA'/CE'/DCGI approved	300	Nos
1.062	(S)PTCA WIRES - Specially Designed for Tortuous vessels. Should be FDA'/CE'/DCGI approved	250	Nos
1.063	(S)PTCA WIRES Extrasupport Polymer coated. Should be FDA'/CE'/DCGI approved	250	Nos
1.064	(S)LINK EXTENSION type 300 cm wires. Should be FDA'/CE'/DCGI approved	250	Nos
1.065	(S)PTCA 009 exchange length wires for Retrograde approach exchange length. Should be FDA'/CE'/DCGI approved	100	Nos
1.066	(EX)Rapid Exchange Balloon Systems with Semi-Compliant/Compliant Design with Excellent Trackability and High Inflation Pressures. Should be FDA' OR CE' AND DCGI approved	600	Nos
1.067	(EX)Rapid Exchange BalloonSystems with Semi-Compliant/Compliant Design with Excellent Trackability and High Inflation Pressures. Should be CE' AND DCGI approved	350	Nos
1.068	(EX)Non-Compliant Balloons for PTCA. Should be FDA' OR CE' AND DCGI approved	600	Nos
1.069	(EX)Non-Compliant Balloons for PTCA. Should be CE' AND DCGI approved	350	Nos
1.07	(S)Laser catheter coronary calcification intervention. Should be CE' AND DCGI approved	2	Nos
1.071	(S)Ultra low profile Non-compliant balloons for PCI. Should be CE' AND DCGI approved	25	Nos
1.072	(S)Ultra low profile semi complaint baloons for PCI. Should be CE' AND DCGI approved	25	Nos
1.073	(S)Lithrotripsy balloons for heavy calcified proximal vessel PCI. Should be CE' AND DCGI approved	4	Nos
1.074	(S)High trackable 0.018" embolisation coils for post op- visceral (GI and Renal artery pseudoaneurysm) . Should be CE' AND DCGI approved	25	Nos
1.075	(S)low profile CTO baloons for PCI starts at 0.75mm in diameter. Should be CE' AND DCGI approved	25	Nos

1.076	(EX)Over the wire Balloons for PTCA. Should be FDA' OR CE' AND DCGI approved	25	Nos
1.077	(EX)Over the wire Balloons for PTSMA. Should be FDA' OR CE' AND DCGI approved	10	Nos
1.078	(EX)Cutting Balloons Coronary. Should be FDA' OR CE' AND DCGI approved	50	Nos
1.079	(EX)Cutting Balloons Peripheral. Should be FDA' OR CE' AND DCGI approved	20	Nos
1.08	(EX)Drug Eluting Balloons Coronary. Should be FDA' OR CE' AND DCGI approved	20	Nos
1.081	(EX)Ultra high pressure Mustang type balloons Peripheral- sizes from 4mm to 10mm all lengths.. Should be FDA' OR CE' AND DCGI approved	50	Nos
1.082	(EX)Drug eluting Balloons for in stent restenosis Coronary. Should be FDA' OR CE' AND DCGI approved	20	Nos
1.083	(EX) Scoring Balloons - for coronary and Peripheral- For severely calcified lesions. Should be FDA' OR CE' AND DCGI approved	10	Nos
1.084	(EX)BALLOON MOUNTED DRUG COATED stain less steel CORONARY STENTS. Should be CE' AND DCGI approved	50	Nos
1.085	(EX)Balloon expandable drug coated Cobalt chromium alloy CORONARY STENTS . Should be FDA' OR CE' AND DCGI approved	100	Nos
1.086	(EX)Balloon expandable drug coated Platinum chromium alloy CORONARY STENTS . Should be FDA' OR CE' AND DCGI approved	100	Nos
1.087	(EX)DRUG ELUTING STENTS FDA and DCGI APPROVED First Generation. Should be FDA approved	250	Nos
1.088	(EX)DRUG ELUTING STENTS FDA and DCGI APPROVED Second Generation. Should be FDA approved	250	Nos
1.089	(EX)DRUG ELUTING STENTS FDA and DCGI APPROVED Third Generation. Should be FDA approved	250	Nos
1.09	(EX)DRUG ELUTING STENTS CE and DCGI APPROVED. Should be CE' and DCGI approved	250	Nos
1.091	(EX)DRUG ELUTING STENTS DCGI APPROVED. Should be DCGI approved	250	Nos
1.092	(EX) DRUG ELUTING CORONARY STENTS TAPERING including longer legths. Should be FDA' OR CE' AND DCGI approved	250	Nos
1.093	(EX) DRUG ELUTING CORONARY STENTS EXTRA LENGTH (beyond 38mm). Should be FDA' OR CE' AND DCGI approved	250	Nos
1.094	(EX) DRUG ELUTING STENTS- POLYMER FREE. Should be FDA' OR CE' AND DCGI approved	100	Nos
1.095	(EX) DRUG ELUTING STENTS- sizes above 48mm for diffuse lesions. Should be FDA' OR CE' AND DCGI approved	100	Nos
1.096	(S) Coronary micro snares (2mm to 5mm). Should be FDA' OR CE' AND DCGI approved	10	Nos
1.097	(EX)COVERED STENTS-CORONARY Low profile ad 0.014" compatable. Should be FDA'/CE'/DCGI approved	20	Nos

1.098	(S) FEMORAL ANGIOPLASTY KIT (1stent): 1. Disposable Cath-Lab Gowns Light Blue Colour full size and elastic wrist bands with water'/ blood impenetrable front and hand shielding - 3 Nos 2. DISPOSABLE STERILE DRAPES FOR CORONARY ANGIOGRAPHY" / ANGIOPLASTY COMPATIBLE WITH Bilateral FEMORAL and RADIAL APPROACHES - 1No 3. DISPOSABLE GAUZE PIECES in (ETO Sterilizable) PACKS OF 10 for use with Cath Lab procedures - 10Nos 4. TEFLON COATED GUIDE WIRES 145 CMS 0.035"ST & J - 1No 5. Arterial Pressure monitoring Transducers (Disposable) - 1No 6. INITIAL PUNCTURE NEEDLES18G - 1No 7. FEMORAL PCI SHEATH SYSTEMS SIZE 5F to 9F with accessories for insertion - 2Nos 8. PTCA WIRE - 1 No 9. 3 RING CONTROL SYRINGES FOR CAG'/PTCA 20cc - 1No 10. 3'/3 ROTATING MANIFOLD FOR CAG'/PTCA-1No 11. TOUHEY BORST Y CONNECTOR- PUSH PULL TYPE and INTRODUCER NEEDLE for PTCA - 1No 12. PTCA GUIDING CATHETER - 1No 13. 3 Ring Finger Control syringe 12 cc - 1No 14. INFUSION ACCELERATOR (PRESSURE BAG) -1No 15. Rapid Exchange Balloon Systems with Semi-Compliant'/Compliant Design with Excellent Trackability and High Inflation Pressures - 1No 16. Non-Compliant Balloons for PTCA - 1No 17. DRUG ELUTING STENTS FDA/ DCGI /CE APPROVED -1 No Should be FDA'/CE'/DCGI approved	150	kit
1.099	(S)FEMORAL ANGIOPLASTY KIT (2 Stent) 1. Disposable Cath-Lab Gowns Light Blue Colour full size and elastic wrist bands with water'/ blood impenetrable front and hand shielding - 3 Nos 2.DISPOSABLE STERILE DRAPES FOR CORONARY ANGIOGRAPHY" / ANGIOPLASTY COMPATIBLE WITH Bilateral FEMORAL and RADIAL APPROACHES - 1No 3.DISPOSABLE GAUZE PIECES in (ETO Sterilizable)PACKS OF 10 for use with Cath Lab procedures - 10Nos 4.TEFLON COATED GUIDE WIRES 145 CMS 0.035"ST & J - 1No 5.Arterial Pressure monitoring Transducers (Disposable) - 1No 6.INITIAL PUNCTURE NEEDLES18G - 1No 7.FEMORAL PCI SHEATH SYSTEMS SIZE 5F to 9F with accessories for insertion - 2Nos 8. PTCA WIRE- 1 No 9.3 RING CONTROL SYRINGES FOR CAG'/PTCA 20cc - 1No 10. 3'/3 ROTATING MANIFOLD FOR CAG'/PTCA-1No 11. TOUHEY BORST Y CONNECTOR- PUSH PULL TYPE and INTRODUCER NEEDLE for PTCA - 1No 12.PTCA GUIDING CATHETER - 1No 13.3 Ring Finger Control syringe 12 cc - 1No 14.INFUSION ACCELERATOR (PRESSURE BAG) -1No 15.Rapid Exchange BalloonSystems with Semi-Compliant'/Compliant Design with Excellent Trackability and High Inflation Pressures - 1No 16.Non-Compliant Balloons for PTCA - 1No 17.DRUG ELUTING STENTS FDA/ DCGI /CE -2 Nos . Should be FDA'/CE'/DCGI approved	150	kit

1.1	(S)RADIAL ANGIOPLASTY KIT (1 Stent) 1. Disposable Cath-Lab Gowns Light Blue Colour full size and elastic wrist bands with water'/ blood impenetrable front and hand shielding - 3 Nos 2.DISPOSABLE STERILE DRAPES FOR CORONARY ANGIOGRAPHY" / ANGIOPLASTY COMPATIBLE WITH Bilateral FEMORAL and RADIAL APPROACHES - 1No 3.DISPOSABLE GAUZE PIECES in (ETO Sterilizable)PACKS OF 10 for use with Cath Lab procedures - 10Nos 4.TEFLON COATED GUIDE WIRES 145 CMS 0.035"ST & J - 1No 5.Arterial Pressure monitoring Transducers (Disposable) - 1No 6.INITIAL PUNCTURE NEEDLES18G - 1No 7.TRANSRADIAL SHEATH SYSTEMS SIZE 6F with accessories for insertion - 1Nos 8. PTCA WIRE - 1 No 9.3 RING CONTROL SYRINGES FOR CAG'/PTCA 20cc - 1No 10. 3'/3 ROTATING MANIFOLD FOR CAG'/PTCA-1No 11. TOUHEY BORST Y CONNECTOR- PUSH PULL TYPE and INTRODUCER NEEDLE for PTCA - 1No 12.PTCA GUIDING CATHETER - 1No 13.3 Ring Finger Control syringe 12 cc - 1No 14.INFUSION ACCELERATOR (PRESSURE BAG) -1No 15.Rapid Exchange BalloonSystems with Semi-Compliant'/Compliant Design with Excellent Trackability and High Inflation Pressures - 1No 16.Non-Compliant Balloons for PTCA - 1No 17.DRUG ELUTING STENTS FDA / DCGI/CE APPROVED -1 No . Should be FDA'/CE'/DCGI approved	150	kit
1.101	(S)RADIAL ANGIOPLASTY KIT (2 Stent) 1. Disposable Cath-Lab Gowns Light Blue Colour full size and elastic wrist bands with water'/ blood impenetrable front and hand shielding - 3 Nos 2.DISPOSABLE STERILE DRAPES FOR CORONARY ANGIOGRAPHY" / ANGIOPLASTY COMPATIBLE WITH Bilateral FEMORAL and RADIAL APPROACHES - 1No 3.DISPOSABLE GAUZE PIECES in (ETO Sterilizable)PACKS OF 10 for use with Cath Lab procedures - 10Nos 4.TEFLON COATED GUIDE WIRES 145 CMS 0.035"ST & J - 1No 5.Arterial Pressure monitoring Transducers (Disposable) - 1No 6.INITIAL PUNCTURE NEEDLES18G - 1No 7.TRANSRADIAL SHEATH SYSTEMS SIZE 6F with accessories for insertion - 1Nos 8. PTCA WIRE - 1 No 9.3 RING CONTROL SYRINGES FOR CAG'/PTCA 20cc - 1No 10. 3'/3 ROTATING MANIFOLD FOR CAG'/PTCA-1No 11. TOUHEY BORST Y CONNECTOR- PUSH PULL TYPE and INTRODUCER NEEDLE for PTCA - 1No 12.PTCA GUIDING CATHETER - 1No 13.3 Ring Finger Control syringe 12 cc - 1No 14.INFUSION ACCELERATOR (PRESSURE BAG) -1No 15.Rapid Exchange BalloonSystems with Semi-Compliant'/Compliant Design with Excellent Trackability and High Inflation Pressures - 1No 16.Non-Compliant Balloons for PTCA - 1No 17.DRUG ELUTING STENT FDA/ DCGI/CE APPROVED -2 No . Should be FDA'/CE'/DCGI approved	100	kit
1.102	(S) 20CC INFLATION DEVICE AS ONE SET capable of inflation pressures upto 30 bar. Should be FDA'/CE'/DCGI approved	1000	nos
1.103	(S)INFLATION DEVICE TO INFLATE LOW AND HIGH PRESSURE UPTO 40 ATM. Should be FDA'/CE'/DCGI approved	1000	nos

1.104	(S) PTCA MANIFOLD WITH FLUSH DEVICE. Should be FDA'/CE'/DCGI approved	1000	nos
1.105	(S) INFUSION ACCELERATOR (PRESSURE BAG). Should be FDA'/CE'/DCGI approved	1000	nos
1.106	(S) 3 RING CONTROL SYRINGES FOR CAG/PTCA 12cc. Should be FDA'/CE'/DCGI approved	1000	nos
1.107	(S) 3 RING CONTROL SYRINGES FOR CAG/PTCA 20cc. Should be FDA'/CE'/DCGI approved	1000	nos
1.108	(D) Thrombus Extraction Catheters for Primary PTCA. Should be FDA'/CE'/DCGI approved	100	nos
1.109	(D) Coronary Microcatheters(0.014" compatable). Should be FDA' or CE' andDCGI approved	100	nos
1.11	(D)Shock wave assisted balloons for calcified lesions. Should be CE' AND DCGI approved	5	nos
1.111	(D)Shock wave assembly and delivery system for ballooons. Should be CE' AND DCGI approved	5	nos
1.112	(D)ULTRA HIGH PRESSURE (30 and above) non compliant CORONARY balloons for sevrely calcified stenosis (All sizes). Should be CE' AND DCGI approved	20	nos
1.113	(D) Coronary Microcatheters for CTO entry- braided with cutting tip. Should be FDA' or CE' andDCGI approved	50	nos
1.114	(D)Steerable Microguide catheters for sidebranch entry -Dual lumen type(over the wire and monorail). Should be FDA' or CE' and DCGI approved	20	nos
1.115	(D)Micro catheters for visceral angio embolisation-Should be hydrophilic and 0.021" compatable. Should be FDA' OR CE' AND DCGI approved	50	nos
1.116	(D)Retrograde coronary collateral Channel dilator . Should be FDA' or CE' and DCGI approved	20	nos
1.117	(S)Superselective cathetor for Mother and Child technique. Should be FDA' or CE' and DCGI approved	20	nos
1.118	(S) Soft Guideliner for mother and child technique. Should be FDA' or CE' and DCGI approved	20	nos
1.119	(D) Intravascular Ultrasound Catheters compatible with Volcano IVUS system . Should be FDA'/CE'/DCGI approved	20	nos
1.12	(D) Intravascular Ultrasound Catheters compatible with ILAB IVUS system with Pull back Slut. Should be FDA'/CE'/DCGI approved	25	nos
1.121	(D)OCT Catheters . Should be FDA'/CE'/DCGI approved	10	nos
1.122	(D)RADI 014 FFR WIRES. Should be FDA'/CE'/DCGI approved	50	nos
1.123	(D)RADI 014 FFR WIRES wireless. Should be FDA'/CE'/DCGI approved	50	nos
1.124	(D)VOLCANO 014 FFR WIRES. Should be FDA'/CE'/DCGI approved	10	nos
1.125	(D) FFR WIRE Compatible with ILAB. Should be FDA'/CE'/DCGI approved	50	nos
1.126	(D)Rotablator Wire 009. Should be FDA'/CE'/DCGI approved	25	nos
1.127	(D)Rotablator Advancer. Should be FDA'/CE'/DCGI approved	25	nos
1.128	(D)OPN type ultra high pressure highly non compliant balloon . Should be FDA' OR CE' AND DCGI approved	25	nos
1.129	(D)Rotablator Burr. Should be FDA'/CE'/DCGI approved	25	nos
1.13	(S)Swan Ganz capable of accommodating 0.038" Guide Wire Large Lumen Size 7F . Should be FDA'/CE'/DCGI approved	25	nos
1.131	(S)Swan Ganz capable of accommodating 0.032/035" Guide Wire Large Lumen Size 6F . Should be FDA'/CE'/DCGI approved	25	nos
1.132	(S)BERMAN BALLOON ANGIO CATHS WITH SIDE HOLES 5F,6F. Should be FDA'/CE'/DCGI approved	25	nos
1.133	(S)BALLOON TEMPORARY PACING LEADS 5F,6F. Should be FDA'/CE'/DCGI approved	25	nos

1.134	(S)Woven Dacron Catheter with platinum Tip Elecrode for Trasvenous Temporary pacing 5F/6F. Should be FDA'/CE'/DCGI approved	1000	nos
1.135	(S)Temporary Pacing Leads sizes with 5F/6F. Should be FDA'/CE'/DCGI approved	1000	nos
1.136	(S)Balloon Flow Assisted Temporary Pacing lead(5F and 6F . Should be FDA'/CE'/DCGI approved	100	nos
1.137	(S)Accessories for Accura based Mitral valvuloplasty systems. Should be CE' AND DCGI approved	10	nos
1.138	(EX)ALL SIZES OF PTMC Balloons- Reusable. Should be FDA'/CE'/DCGI approved	50	nos
1.139	(EX)ALL SIZES OF MITRAL VALVE BALLOON DILATATION SYSTEMS WITH REUSABLE LATEX RUBBER BALLOON DESIGN AND ACCESSORIES TO BE QUOTED. Should be FDA'/CE'/DCGI approved	25	nos
1.14	(S)PRESSURE MANOMETER LINES MALE/FEMALE 150 CM. Should be FDA'/CE'/DCGI approved	1000	nos
1.141	(S)PRESSURE MANOMETER LINES MALE/MALE 150 CM. Should be FDA'/CE'/DCGI approved	1000	nos
1.142	(S)THREE WAY STOPCOCKS. Should be FDA'/CE'/DCGI approved	1500	nos
1.143	(S)HIGH PRESSURE CONTRAST INJECTION TUBE. Should be FDA'/CE'/DCGI approved	250	nos
1.144	(S)HIGH FLOW ANGIO FLUSH CATHS. Should be FDA'/CE'/DCGI approved	50	nos
1.145	(S)LEFT INTERNAL MAMMARY GUIDE CATHETER 5 F,6F and & 7F . Should be FDA'/CE'/DCGI approved	50	nos
1.146	(S)BRAIDED MULTIPURPOSE GUIDE CATHS(5F AND 6F). Should be FDA'/CE'/DCGI approved	60	nos
1.147	(S) SOFT TIPPED braided Guide catheters for (4F,5F) for coronary/visceral use. Should be FDA' or CE' and DCGI approved	25	nos
1.148	(S)BRAIDED SIMMONS SHAPED CEREBRAL CATHS. Should be FDA'/CE'/DCGI approved	50	nos
1.149	(S)RENAL DOUBLE CURVE GUIDE CATHETER 4F, 5F 80CM , 120 CM, 140 CM. Should be FDA'/CE'/DCGI approved	50	nos
1.15	(S) TERUMO GLIDE CATHETER 4F,5F AND 6 F. Should be FDA'/CE'/DCGI approved	50	nos
1.151	(S)YASHIRO GLIDE CATHETER 4F, 5F 80 CM,120 CM,140CM. Should be FDA'/CE'/DCGI approved	50	nos
1.152	(EX)PERIPHERAL BALLOONS(0.035" compatable) for femoro popliteal and iliac lesions. Should be FDA'/CE'/DCGI approved	50	nos
1.153	(EX) RENAL BALLOON(0.014" compatable). Should be FDA'/CE'/DCGI approved	50	nos
1.154	(S) 4 F AND 5 F catheter COMPATABLE PERI MEMBRANEUOUS VSD DEVICE with double screw design for antegrade and retrograde closure. Should be CE' AND DCGI approved	5	nos
1.155	(EX)CAROTID and BTK BALLOONS(0.014" compatable) .Balloon Shaft length should be 160 CM AND ABOVE. Should be FDA'/CE'/DCGI approved	50	nos
1.156	(EX)SMALL VESSEL BALLOON(Diameters should be from 1mm to 4mm) All lengths. Should be FDA'/CE'/DCGI approved	50	nos

1.157	(EX)PERIPH /RENAL/CAROTID GUIDING-100 cm and longer. Should be FDA'/CE'/DCGI approved	50	nos
1.158	(EX)RENAL STENTS -0.014' compatable Balloon Mounted Shaft Length 80 cm- All sizes. Should be FDA'/CE'/DCGI approved	50	nos
1.159	(EX) Self expanding Femoro popliteal stents - all diameters. Should be FDA'/CE'/DCGI approved	50	nos
1.16	(EX)Dilators of Coons type for serial vascular dilation for large access . Should be CE' and DCGI approved	10	nos
1.161	(EX) Balloon expandable STENTS PERIPHERAL- All sizes. Should be FDA'/CE'/DCGI approved	25	nos
1.162	(EX) Self expanding Iliac stents - all diameters. Should be FDA' or CE' and DCGI approved	25	nos
1.163	(EX) CAROTID DISTAL PROTECTION DEVICES. Should be FDA'/CE'/DCGI approved	10	nos
1.164	(EX) CAROTID STENTS TAPERING Self Expanding. Should be FDA'/CE'/DCGI approved	25	nos
1.165	(EX)CAROTID STENTS Straight Self Expanding. Should be FDA'/CE'/DCGI approved	25	nos
1.166	(EX)Nitinol Vascular plugs for embolisation. Should be FDA'/CE'/DCGI approved	10	nos
1.167	(EX)STENT GRAFTS FOR VISCERAL VASCULAR SYSTEM(Should be highy trackable,0.035" compatable). Should be FDA'/CE'/DCGI approved	25	nos
1.168	(S) Sizing BALLOONS for ASD/PDA. Should be FDA'/CE'/DCGI approved	20	nos
1.169	(D)VENA CAVA FILTER- Retrivable, Jugular Deployement. Should be FDA'/CE'/DCGI approved	10	nos
1.17	(D)VENA CAVA FILTER- Retrivable, Femoral Deployement. Should be FDA'/CE'/DCGI approved	10	nos
1.171	(D)VENA CAVA FILTER- RETRIVAL SYSTEM . Should be FDA'/CE'/DCGI approved	10	nos
1.172	(S)PERIPHERAL GUIDE WIRES& SPECIALITY WIRES Exchange Length (0.018"). Should be FDA'/CE'/DCGI approved	25	nos
1.173	(S)PERIPHERAL GUIDE WIRES& SPECIALITY WIRES Exchange length(0.035"). Should be FDA'/CE'/DCGI approved	25	nos
1.174	(S)ROAD RUNNER POLYMER COATED WIRE 035 staright and J. Should be FDA'/CE'/DCGI approved	10	nos
1.175	(S)PERIPHERAL CHRONIC TOTAL OCCLUSION WIRES 0.014, 0.0.35 diameter. Should be FDA'/CE'/DCGI approved	25	nos
1.176	(EX) EMBOLIZATION COILS 035" all sizes. Should be FDA'/CE'/DCGI approved	100	nos
1.177	(EX) EMBOLIZATION COILS 0.038 to 052" all sizes. Should be FDA'/CE'/DCGI approved	100	nos

1.178	(EX) EMBOLIZATION MICRO COILS 018 all sizes - Non Detachable and freely deliverable. Should be FDA'/CE'/DCGI approved	100	nos
1.179	(EX)EMBOLIZATION MICRO COILS 018 all sizes -DETACHABLE 2D. Should be FDA'/CE'/DCGI approved	100	nos
1.18	(EX) Vascular embolisation nitinol based occluders all sizez with delivery system. Should be FDA' or CE' and DCGI approved	10	nos
1.181	(EX) EMBOLIZATION MICRO COILS 018 all sizes -DETACHABLE 3D. Should be FDA'/CE'/DCGI approved	10	nos
1.182	(EX)ENDOVASCULAR SNARES with Metal Tip. Should be FDA'/CE'/DCGI approved	5	nos
1.183	(EX)ENDOVASCULAR SNARES with Polymer Tip. Should be FDA'/CE'/DCGI approved	5	nos
1.184	(EX)ENDOVASCULAR SNARES Gooseneck. Should be FDA'/CE'/DCGI approved	5	nos
1.185	(EX)EMBOLISATION PARTICLES Poly Vinyl Alcohol all sizes. Should be FDA'/CE'/DCGI approved	50	nos
1.186	(S)Super Absorbant Polymer Microspheres for Drug(Doxorubucin) Adsorption for TACE. Should be FDA'/CE'/DCGI approved	10	nos
1.187	(S)Gelfoam (Powder or granule form) for Embolization. Should be FDA'/CE'/DCGI approved	10	nos
1.188	(S)N-butyl cyanoacrylate GLUE for Embolization . Should be FDA'/CE'/DCGI approved	10	nos
1.189	(EX)MICROCATHETERS with Inbuilt hydrophilic wire for Peripheral Angiography and Particle /Coil delivery. Should be FDA'/CE'/DCGI approved	50	nos
1.19	(EX)MICROCATHETERS - STEERABLE/ SHAPABLE with Inbuilt hydrophilic wire for Peripheral Angiography and Particle /Coil delivery. Should be FDA'/CE'/DCGI approved	50	nos
1.191	(EX)LARGE DIAMETER COVERED AORTA-ARCH AND ZONE 2 STENT GRAFTS TAA all sizes. Should be FDA'/CE'/DCGI approved	10	nos
1.192	(EX) Aorto Uni iliac stent covered grafts All sizes. Should be CE' AND DCGI approved	2	nos
1.193	(EX)AORTIC STENT covered GRAFTS for DTA aneurysm stenting . Should be FDA'/CE'/DCGI approved	5	nos
1.194	(EX)Bare AORTIC STENTS FOR Coarctation (Sizes should be above 20mm up to 40mm) -Only Baloon expandable design. Should be FDA'/CE'/DCGI approved	2	nos
1.195	(EX) Covered AORTIC stents (balloon expandable only) for aortic coarctation . Diameters should be 20mm and above. Should be CE' AND DCGI approved	2	nos

1.196	(EX) Low profile Dacron / PTFE covered stent grafts (20-45mm) for sealing extensive aortic dissection . Should be CE' AND DCGI approved	5	nos
1.197	(EX) Low profile hydrophilic sheaths for aortic procedures (12 F and above) . Should be FDA'/CE'/DCGI approved	10	nos
1.198	(EX) Covered stent grafts for aortic arch aneurysm with onsite fenestraion capability. Should be FDA'/CE'/DCGI approved	5	nos
1.199	(EX) Aortic covered Stent grafts (straight) capable for fenestration for justa renal and juxta subclavain aneurysms and traumatic aortic injury. Should be FDA'/CE'/DCGI approved	5	nos
1.2	(EX) ABDOMINAL AORTIC ANEURYSM STENT GRAFTS WITH ILIAC LIMBS for abd aorta aneurysm. Should be FDA'/CE'/DCGI approved	5	nos
1.201	(EX)ASD DEVICES: All cutomised sizes from 8mm to 50 mm. Should be CE' AND DCGI approved	25	nos
1.202	(EX) Ultra compliant linear Sizing balloons with markers (34mm to 44mm) for balloon asssited device deployment compatable with contralateral 8F sheath. Should be CE/DCGI approved	25	nos
1.203	(EX) Ultra compliant linear Sizing balloons with markers (24mm to 34mm) for balloon asssited device deployment compatable with contralateral 7F sheath. Should be CE' and DCGI approved	25	nos
1.204	(EX) COOK SHEATHS (all sizes 6F-15F) with check flo valves) WITH ANGLED TIPS FOR ASD DEVICE DELIVERY. Should be FDA' or CE' and DCGI approved	100	nos
1.205	(EX) Fenestrated ASD device for mutiple septal defects. Should be FDA'/CE'/DCGI approved	10	nos
1.206	(EX)ASD AMPLATZER TYPE SELF CENTRE DEVICES WITH ALL sizes from 8mm and above . Should be FDA approved	25	nos
1.207	(EX)PDA OCCUSION DEVICE --ALL CUSTOMISABLE SIZES WITH FROM 3MM TO 20MM WAIST SIZE . Should be FDA'/CE'/DCGI approved	25	nos
1.208	(EX) Delivery System and cable set for PDA occlusion devie (4F- 12 F). Should be FDA'/CE'/DCGI approved	25	nos
1.209	(EX)ADO-1 device for all morphology. Should be FDA' or CE' andDCGI approved	25	nos
1.21	(EX) DUCT OCCLUSION DEVICE DELIVERY SYSTEM NONTAPERING SHEATHS with sidearm+INTRODUCER CABLE SET . Should be FDA'/CE'/DCGI approved	25	nos
1.211	(EX)PDA OCCUSION DEVICE with Dual Discs of equal size . Should be FDA' or CE' andDCGI approved	5	nos
1.212	(EX)PDA DELIVERY SYSTEMS AND INTRODUCER CABLE SYSTEMS . Should be FDA'/CE'/DCGI approved	10	nos
1.213	(EX)Ventricular septal Closure Devices All sizes (4mm to18 mm). Should be FDA'/CE'/DCGI approved	10	nos

1.214	(EX) Delivery cable and sheath set for Ventricular septal defect closure. Should be FDA' or CE' and DCGI approved	10	nos
1.215	(EX)Muscular VSD Closure Devices-All sizes and types . Should be FDA'/CE'/DCGI approved	10	nos
1.216	(EX) Low profile Delivery sheath and cable for VSD closure devices(5F to 14F). Should be CE' and DCGI approved	10	nos
1.217	(EX) Low profile Duct occluder device for neonates and infants. Should be CE' AND DCGI approved	5	nos
1.218	(EX)Ventricular septal Closure Devices All sizes with delivery system. Should be FDA' or CE' and DCGI approved	5	nos
1.219	(EX) Atrial septal occluder(self centering) for complex morphology with balloon assisted delivery . Should be FDA' or CE' andDCGI approved	10	nos
1.22	(EX) Post Infarct ventricular septal rupture Closure Devices- All sizes and types.FDA approved	10	nos
1.221	(EX) PURPLE Recovery sheaths for embolised ASD devices-16 F and above size. Should be FDA approved	10	nos
1.222	(EX) Large sized atrial septal occluders above 40mm size. Should be CE' AND DCGI approved	5	nos
1.223	(EX) Vascular plug closure devices for coronary and Pulmonary AV fistula. Should be FDA'/CE'/DCGI approved	5	nos
1.224	(EX) Ventricular septal defect (device alone) for antegrade closure . Should be FDA' or CE' and DCGI approved	5	nos
1.225	(EX) Dual disc nitinol biocompatable layer coat secundum type atrial occluders (ALL SIZSES) for Pre assembled delivery system. Should be FDA' or CE' andDCGI approved	10	nos
1.226	(EX) Extra soft Nitinol multi functional Closure Devices compatable with guide catheter delivery. Should be CE/DCGI approved	10	nos
1.227	(EX)Left atrial appendage closure device. Should be FDA'/CE'/DCGI approved	10	nos
1.228	(EX)left atrial appendage closure delivery system. Should be FDA'/CE'/DCGI approved	10	nos
1.229	(EX) Single disc occluders with device alone with sizes from 3mm onwards up to 20 mm .. Should be FDA' or CE' andDCGI approved	10	nos
1.23	(EX)Trans catheter aortic valve (self expanding). Should be FDA'/CE'/DCGI approved	10	nos
1.231	(EX)Trans catheter aortic valve (balloon expanding). Should be FDA'/CE'/DCGI approved	10	nos
1.232	(EX)Trans catheter aortic valve delivery sheath(self expanding). Should be FDA'/CE'/DCGI approved	10	nos
1.233	(EX)Trans catheter aortic valve delivery sheath(balloon expanding). Should be FDA'/CE'/DCGI approved	10	nos

1.234	(EX)Large diametre (14-20mm) balloon expanding aortic/ vena caval stents. Should be FDA'/CE'/DCGI approved	25	nos
1.235	(EX)Large bore low profile manual aspiration system for high thrombus load. Should be FDA'/CE'/DCGI approved	25	nos
1.236	(EX)Ultra low profile semi compliant coronary balloons for Chronic total occlusions. Should be FDA'/CE'/DCGI approved	25	nos
1.237	(EX)Ultra High pressure above 25 atm non compliant balloon system for calcified lesions. Should be FDA'/CE'/DCGI approved	25	nos
1.238	(EX)Balloon expandable covered stents(sizes 5-20mm) for fenestrated TEVAR/EVAR. Should be FDA'/CE'/DCGI approved	25	nos
1.239	(EX) Covered self expanding covered stent grafts for anurysm and leaks (4mm to 12mm). Should be FDA' or CE' andDCGI approved	10	nos
1.24	(EX)Coda balloon for optimal Aortic Stent expansion . Should be FDA' or CE' and DCGI approved	10	nos
1.241	(EX)Anzel sheaths for fenestarted EVAR . Should be FDA'/CE'/DCGI approved	20	nos
1.242	(EX)Braided kink resistant long(100-120cm) delivery sheaths for VSD closure. Should be FDA'/CE'/DCGI approved	25	nos
1.243	(EX)Braided kink resistant flexible sheaths (60-100cm) for fenestarted TEVAR and carotid interventions-straight and curved. Should be FDA'/CE'/DCGI approved	20	nos
1.244	(S) Angioseal type-sponge based vascular closure system. Should be CE' AND DCGI approved	10	nos
1.245	(EX)Radiation protection drape pads. Should be FDA'/CE'/DCGI approved	50	nos
1.246	(S)GEL FOAM POWDER FOR ANGIO EMBOLISATION. Should be FDA' or CE' andDCGI approved	25	nos
1.247	(EX)Cyanoacrylate glue for sealing coronary perforation. Should be FDA'/CE'/DCGI approved	25	nos
1.248	(EX)Dissection re entry balloon for coronary CTO. Should be FDA'/CE'/DCGI approved	25	nos
1.249	(EX)Sub intimal tracking device for CTO. Should be FDA'/CE'/DCGI approved	25	nos
1.25	(EX)Large bore long femoral sheaths 12-18 F for aortic/TAVI interventions. Should be FDA'/CE'/DCGI approved	50	nos
1.251	(EX)Suture based haemostasis device for large bore Femoral access. Should be FDA'/CE'/DCGI approved	10	nos
1.252	(EX)Scoring balloons for debulking calcified lesions. Should be FDA'/CE'/DCGI approved	25	nos
1.253	(EX)Vascular Plugs with delivery systems and sheaths. Should be FDA'/CE'/DCGI approved	10	nos
1.254	(EX)Expandable femoral sheaths. Should be FDA'/CE'/DCGI approved	5	nos

1.255	(S)STR shaped woven dacron intra coronary guide catheter(above 100 cm length). Should be CE' AND DCGI approved	10	nos
1.256	(S)Slender sheaths for radial access -Should be expandable. Should be CE' AND DCGI approved	25	nos
1.257	(S)Coronary micro snares. Should be CE' and DCGI approved	2	nos
1.258	(EX)BIOPTOME for Coil Delivery. Should be FDA'/CE'/DCGI approved	10	nos
1.259	(EX)Pulmonary'/Aortic balloon systems comprising of Pulmonary Valvuloplasty Balloons all sizs, curves and lengths . Should be FDA'/CE'/DCGI approved	10	nos
1.26	(EX)Single diameter Aortic Valvuloplasty balloons of Cribier-Letac all sizes and lengths. Should be FDA'/CE'/DCGI approved	10	nos
1.261	(EX)High pressure non compliant peripheral balloons- (sizes from 4mm-10mm) 0.035" compatable (All lengths). Should be CE' and DCGI approved	10	nos
1.262	(EX)Stepped diameter balloons Cribier- Letac all sizes and lengths. Should be FDA'/CE'/DCGI approved	10	nos
1.263	(EX)Pediatric Valvuloplasty balloons with accelerator, Bifoil balloons, Trifoil Balloons all sizes. Should be FDA'/CE'/DCGI approved	10	nos
1.264	(S)Amplatz apex J extra stiff guide wire exchange lengths. Should be FDA'/CE'/DCGI approved	25	nos
1.265	(S)14 F balloon introducer systems. Should be FDA'/CE'/DCGI approved	10	nos
1.266	(EX)COOK LONG SHEATHS CAPABLE OF ASD/VSD/PDA device deliverySIZES 6Fto 16F. Should be FDA'/CE'/DCGI approved	50	nos
1.267	(S) INITIAL PUNCTURE NEEDLES18G. Should be FDA'/CE'/DCGI approved	1000	nos
1.268	(S) MICROPUNCTURE NEONATAL INTRODUCER SETS AND SHEATH SYSTEMS 4F,5F. Should be FDA'/CE'/DCGI approved	100	nos
1.269	(S)GUIDE WIRE 38,A COOK HIWIRE 150CM. Should be FDA'/CE'/DCGI approved	25	nos
1.27	(S)CARDIAC MULTIPACKS COMPRISING OF VENTRICULAR PIG TAIL CATHETER, LEFT CORONARY CATHETER, RIGHT CORONARY CATHETER, CATHETER INTRODUCER SHEATH ALL SIZES. Should be FDA'/CE'/DCGI approved	100	kit
1.271	(S)Guide Extension Catheter,. Should be FDA'/CE'/DCGI approved	100	nos
1.272	(S Balloon atrial septostomy catheter. Should be FDA'/CE'/DCGI approved	5	nos
1.273	(S)TRANSSEPTAL CATHETER . Should be FDA'/CE'/DCGI approved	10	nos
1.274	(S)BROCKENBOROUGH NEEDLE ADULT . Should be FDA'/CE'/DCGI approved	50	nos
1.275	(S)BROCKENBOROUGH NEEDLE PEDIATRICS . Should be FDA'/CE'/DCGI approved	50	nos
1.276	(S)MULLINS SHEATH with Matching Dilator with sidearm 6F TO 14 F all sizes. Should be FDA'/CE'/DCGI approved	50	nos

1.277	(S)MULLINS SHEATH with Matching Dilator with out sidearm 6F, 7F, 8F. Should be FDA'/CE'/DCGI approved	50	nos
1.278	(S)BALKAN CONTRALATERAL SHEATH 6F,7F,8F. Should be FDA'/CE'/DCGI approved	20	nos
1.279	(S)MULLINS SHEATH CHILD. Should be FDA'/CE'/DCGI approved	10	nos
1.28	(S)Bipolar temporary pacing electrodes, Sizes: 4F,5F,6F. Should be FDA'/CE'/DCGI approved	50	nos
1.281	(S)Contrast Media for Angiography and Angioplasty (Non-Ionic 350'/370 100 ml Bottles). Should be FDA'/CE'/DCGI approved	100	nos
1.282	(S)Contrast Media for Angiography and Angioplasty (Non-Ionic 350'/370 50 ml Bottles). Should be FDA'/CE'/DCGI approved	100	nos
1.283	(S)Iso Osmolar Contrast Media for Angiography and Angioplasty for Patients with Renal Dysfunction (Non-Ionic 350'/370 100 ml Bottles). Should be FDA'/CE'/DCGI approved	100	nos
1.284	(S)Iso Osmolar Contrast Media for Angiography and Angioplasty for Patients with Renal Dysfunction (Non-Ionic 350'/370 50 ml Bottles). Should be FDA'/CE'/DCGI approved	100	nos
1.285	(S)Myocardial Echocardiographic Contrast comprising of Sulphur Hexachloride or equivalent Micro Bubbles. Should be FDA'/CE'/DCGI approved	10	nos
1.286	(S)Bioptomes for Endomyocardial biopsy : Bioptomes ; 5.2F. Should be FDA'/CE'/DCGI approved	10	nos
1.287	(S)Radial Sheaths with Introducer needle, Guide wire and dilator: Complete Set:. Should be FDA'/CE'/DCGI approved	500	nos
1.288	(S)Oximetry Cuvets for AVOXIMETER. Should be FDA'/CE'/DCGI approved	250	nos
1.289	Inj TIROFIBAN 100 mL bottles. Should be FDA'/CE'/DCGI approved	50	nos
1.29	Inj BIVALURIDIN.FDA approved	25	nos
1.291	(S)TROPONIN I QUANTITATIVE BED SIDE TEST KIT . Should be FDA'/CE'/DCGI approved Measuring Device to be supplied	250	nos
1.292	(S)TROPONIN T QUANTITATIVE BED SIDE TEST KIT. Should be FDA'/CE'/DCGI approved Measuring Device to be supplied	200	nos
1.293	(S)HIGH SENSITIVE TROPONIN T QUANTITIVE QUANTITATIVE BED SIDE TEST KIT. Should be FDA'/CE'/DCGI approved Measuring Device to be supplied	200	nos
1.294	(S)HIGH SENSITIVE TROPONIN I QUANTITATIVE- FDA approved-Analyser Should be Supplied. Should be FDA' approved Measuring Device to be supplied	200	nos
1.295	(S)N-TERMINAL PRO BNP QUANTITATIVE BED SIDE TEST KIT. Should be FDA'/CE'/DCGI approved Measuring Device to be supplied	50	nos
1.296	(S)N-TERMINAL BNP QUANTITATIVE BED SIDE TEST KIT. Should be FDA'/CE'/DCGI approved Measuring Device to be supplied	50	nos
1.297	(S)D-TIMER QUANTITATIVE BED SIDE TEST KIT. Should be FDA'/CE'/DCGI approved Measuring Device to be supplied	50	nos
1.298	(S)NGAL QUANTITATIVE BED SIDE TEST KIT. Should be FDA'/CE'/DCGI approved Measuring Device to be supplied	300	nos
1.299	(S)Bedside Lipid Profile Measurement Kits.Reading Machine should be provided free	500	nos
1.3	(S)Bedside INR Measurement Kits.Reading Machine should be provided free	250	nos
1.301	(S)Vinyl layered LEAD FREE aprons superior quality imported types for radiation protection during interventional procedures. COVERING BOTH FRONT AND BACK SINGLE PIECE MEDIUM SIZE. Should be	20	nos

	FDA'/CE'/DCGI approved & Item to be Demonstrated to departmental selection committee		
1.302	(S)Vinyl layered LEAD FREE aprons superior quality imported types for radiation protection during interventional procedures. COVERING BOTH FRONT AND BACK TWO PIECE SKIRT AND BLOUSE TYPE MEDIUM SIZE. . Should be FDA'/CE'/DCGI approved & Item to be Demonstrated to departmental selection committee	20	nos
1.303	(S)Vinyl layered LEAD FREE aprons superior quality imported types for radiation protection during interventional procedures. COVERING BOTH FRONT AND BACK TWO PIECE SKIRT AND BLOUSE TYPE LARGE SIZE. Should be FDA'/CE'/DCGI approved & Item to be Demonstrated to departmental selection committee	20	nos
1.304	(S)Thyroid Collars Radiation Protection. Should be FDA'/CE'/DCGI approved & Item to be Demonstrated to departmental selection committee	20	nos
1.305	(S)Gonadal Shields Radiation Protection. Should be FDA'/CE'/DCGI approved & Item to be Demonstrated to departmental selection committee	10	nos
1.306	(S)Radaiation Protective Eye Goggles. Should be FDA'/CE'/DCGI approved & Item to be Demonstrated to departmental selection committee	10	nos
1.307	(S)Radaiation Protective Head Gear. Should be FDA'/CE'/DCGI approved & Item to be Demonstrated to departmental selection committee	10	nos
1.308	(S) Cath-Lab Cotton Dress Shirt with three pockets and Pyjama with pockets Large and medium Sized Light Blue Colour with label of " Cath-Lab JIPMER" on both the shirt and the pyjama and Size printed.Sample to be supplied	100	nos
1.309	(S) Disposable Cath-Lab Gowns Light Blue Colour full size and elastic wrist bands with water/ blood impenetrable front and hand shielding. Should be FDA'/CE'/DCGI approved	500	nos
1.31	(S) Glucometer Test Strips with four glucometers. Should be FDA'/CE'/DCGI approved	500	nos
1.311	(D)Single Chamber Demand Cum Overdrive Temporary Pulse Generatorsin various Dual Chamber and Single Chamber modes. Should be FDA'/CE'/DCGI approved	5	nos
1.312	(D)Dual Chamber Demand Cum Overdrive Temporary Pulse Generatorsin various Dual Chamber and Single Chamber modes. Should be FDA'/CE'/DCGI approved	5	nos
1.313	5 French fixed curve quadripolar catheter with 2-5-2 interelectrode spacing. One matching connector for every 5 catheters. Should be FDA'/CE'/DCGI approved	20	nos
1.314	6 French deflectable quadripolar catheter with 2-5-2 interelectrode spacing and matching connector. Should be FDA'/CE'/DCGI approved	6	nos
1.315	6 French deflectable decapolar catheter with 2-5-2 interelectrode spacing and matching connector. Should be FDA'/CE'/DCGI approved	20	nos
1.316	6 or 7 French deflectable duodecapolar catheters (Halo) with matching connectors. Should be FDA'/CE'/DCGI approved	10	nos
1.317	Long deflectable sheath with hemostatic valve, side port, atraumatic tip, braided shaft, uni- or bidirectional steerability, availability in small, medium or large curves, minimum inner diameter or 7F, minimum length of 65 cms. Should be FDA'/CE'/DCGI approved	6	nos
1.318	7 French quadripolar Radiofrequency ablation catheter with a 4 mm tip, unidirectional deflection, availability in small, medium and large curves. Should be compatible with Stockert and IBI RF generators and Suitable connecting cable should be provided for every 5 catheters.. Should be FDA'/CE'/DCGI approved	15	nos
1.319	7 French quadripolar Radiofrequency ablation catheter with a 4 mm tip, bidirectional deflection, availability in small, medium and large curves. Should be compatible with Stockert and IBI RF generators and suitable connecting cables should be provided . Should be FDA'/CE'/DCGI approved	10	nos

1.32	7 French quadripolar Radiofrequency ablation catheter with a 8 mm tip, unidirectional deflection, availability in medium and large curves. Should be compatible with Stockert and IBI RF generators and Suitable connecting cable should be provided for every 5 catheters.. Should be FDA'/CE'/DCGI approved	4	nos
1.321	7 French quadripolar Radiofrequency ablation catheter with irrigated tip, uni- or bi-directional deflection and availability in medium and large curves. Should be compatible with Stockert and IBI RF generators and Suitable connecting cable should be provided for every 5 catheters.. Should be FDA'/CE'/DCGI approved	4	nos
1.322	Quadripolar, irrigated tip, Radiofrequency ablation catheter with a 3.5 mm tip and unidirectional deflection for use with the CARTO mapping system. Should include irrigation tubing set for interface between the irrigation pump and the ablation catheter and matching external reference patch for use with the CARTO 3 system. Should be FDA'/CE'/DCGI approved	20	nos
1.323	Reusable indifferent electrode pad compatible with the Stockert ablator. Should be FDA'/CE'/DCGI approved	4	nos
1.324	Surface electrode kit for use with the Ensite NavX system including 3 pairs of surface patches, 10 ECG electrodes and 1 system reference electrode . Should be FDA'/CE'/DCGI approved	4	nos
1.325	SSIR - Single chamber rate responsive implantable multiprogrammable pacemaker with maximum programmable sensitivity of 0.5 mv or more with matching leads and subclavian introducer set to be quoted as one set. Active fixation leads should be supplied as atrial (50%) / ventricular (50%). Should be FDA'/CE'/DCGI approved	50	nos
1.326	VDD implantable multiprogrammable pacemaker with shortest programmable AV delay of 50 ms, with matching single pass steroid eluting VDD lead and matching sublavian introducer. Should be FDA'/CE'/DCGI approved	50	nos
1.327	DDD - Dual chamber implantable multiprogrammable pacemaker with shortest programmable Av delay of 50 ms, with matching set of steroid eluting active fixation atrial and ventricular leads and two matching subclavian introducers.. Should be FDA'/CE'/DCGI approved	50	nos
1.328	DDDR - Dual chamber rate-responsive implantable multiprogrammable pacemaker with shortest programmable Av delay of 50 ms, with matching set of steroid eluting active fixation atrial and ventricular leads and two matching subclavian introducers.. Should be FDA'/CE'/DCGI approved	25	nos
1.329	Bipolar lead specially designed for placement in coronary sinus for LV pacing. Should be supplied with 9F short sheath with valve, long preshaped slittable sheaths in various shapes for CS cannulation and inner subselective sheath. Lead, long sheath, subselective sheath and short sheath should be quoted as one set.. Should be FDA'/CE'/DCGI approved	20	nos
1.33	CRT - Triple chamber pacing system comprising pacemaker, active atrial and ventricular leads and bipolar, over the wire coronary sinus lead. Should include long preshaped slittable sheaths for coronary sinus cannulation in multiple shapes, subselective inner sheaths, guidewire and occlusion balloon catheter for coronary venography. Coronary sinus lead should be available in multiple curves '/ sizes.. Should be FDA'/CE'/DCGI approved	20	nos
1.331	AICD - Single chamber, implantable cardioverter defibrillator for treatment of life threatening ventricular arrhythmias with facility for bradycardia pacing and tiered therapy for tachycardias, maximum shock energy of at least 35 J, with matching single and dual coil, integrated bipolar active fixation ventricular leads.. Should be FDA'/CE'/DCGI approved	20	nos
1.332	AICD - Dual chamber, implantable cardioverter defibrillator for treatment of life threatening ventricular arrhythmias with facility for bradycardia pacing and tiered therapy for tachycardias, maximum shock energy of atleast 35 J, with matching single and dual coil, integrated bipolar active fixation atrial and ventricular leads.. Should be FDA'/CE'/DCGI approved	20	nos

1.333	CRT-D - Triple Chamber Pacemaker System with cardioverter defibrillator for treatment of life threatening ventricular arrhythmias with facility for CRT pacing and tiered therapy for tachycardias, maximum shock energy of atleast 35 J, with matching single and dual coil, integrated bipolar, active fixation atrial and ventricular leads. . Should be FDA'/CE'/DCGI approved	10	nos
1.334	Steroid eluting bipolar active fixation atrial leads with introducer sheath. Should be FDA'/CE'/DCGI approved	20	nos
1.335	Steroid eluting bipolar active fixation ventricular leads with introducer sheath. Should be FDA'/CE'/DCGI approved	30	nos
1.336	Single pass VDD leads with introducer sheath. Should be FDA'/CE'/DCGI approved	10	nos
1.337	Single and Dual coil (50-50%), integrated bipolar active fixation ICD leads. Should be FDA'/CE'/DCGI approved	10	nos
1.338	Active fixation epicardial leads . Should be FDA'/CE'/DCGI approved	10	nos
1.339	Quadripolar lead specially designed for placement in coronary sinus for LV pacing. Should be supplied with 9F short sheath with valve, long preshaped slittable sheaths in various shapes for CS cannulation and inner subselective sheath. Lead, long sheath, subselective sheath and short sheath should be quoted as one set.. Should be FDA'/CE'/DCGI approved	10	nos
1.34	Implantable loop recorder for subcutaneous implantation, capable of recording at least one channel subcutaneous ECG. Matching hand-held activator should be provided. . Should be FDA'/CE'/DCGI approved	5	nos
1.341	Steerable, long sheath introducer of at least 8F inner diameter, with atraumatic tip and bidirectional steering, designed for use in epicardial mapping. Should be FDA'/CE'/DCGI approved	2	nos
1.342	Single chamber MRI conditional pacemaker with matching atrial or ventricular active or passive fixation lead and introducer sheath set. . Should be FDA'/CE'/DCGI approved	5	nos
1.343	Dual chamber MRI conditional pacemaker with matching atrial and ventricular active or passive fixation leads and 2 introducer sheath sets.. Should be FDA'/CE'/DCGI approved	5	nos
1.344	Disposable Heartstart adult plus mutifunction defibrillation pads for use with the Philips defibrillator. Should be FDA'/CE'/DCGI approved	10	nos
1.345	Disposable pads for defibrillation / transcutaneous pacing for use with the Zoll defibrillator. Should be FDA'/CE'/DCGI approved	10	nos
1.346	(S)One set of high grade stainless steel surgical instruments for minor surgical procedure. Must include the following instruments – Sponge forceps 2 Nos, Towel clamp – 4, Hemostatic Crile forceps straight – 4, hemostatic Crile forceps curved – 6, Allis tissue forceps – 2, Mayo Hegar needle holder 6 inches – 1, Mayo Hegar needle holder 7 inches – 1, Mayo scissors straight – 1, Metzenbaum scissors curved – 1, Small curved scissors – 1, small straight scissors – 1, thumb forceps – 2, adsons toothed forceps – 2, Knife handle #7 – 1, knife handle #4 – 1, Weitlaner self retaining retractor sharp 5 inches – 1, double hook retractor – 2, right angle forceps – 1. Should be FDA'/CE'/DCGI approved & Item to be Demonstrated to departmental selection committee	2	kit
1.347	(S)LEAD locking device for transvenous extraction of chronically implanted pacemaker and defibrillator leads with mechanical rotating dilator 9F/11F/13F diameter with a flexible shaft and dilator sheath set . Should be FDA'/CE'/DCGI approved	25	nos
1.348	(S)SPO2 Probe Compaitable with Monitor Model : SPACE LABS. Should be FDA'/CE'/DCGI approved	10	nos

1.349	(S)ECG Probe Compaitable with Monitor Model : SPACE LABS. Should be FDA'/CE'/DCGI approved	10	nos
1.35	(S)BP Cuff Compaitable with Monitor Model : SPACE LABS. Should be FDA'/CE'/DCGI approved	10	nos
1.351	(S)SPO2 Probe Compaitable with Monitor Model : Mindray MEC 2000. Should be FDA'/CE'/DCGI approved	10	nos
1.352	(S)ECG leads Compaitable with Monitor Model : Mindray MEC 2000. Should be FDA'/CE'/DCGI approved	10	nos
1.353	(S)NIBP cuff (Adult)Compaitable with Monitor Model : Mindray MEC 2000. Should be FDA'/CE'/DCGI approved	10	nos
1.354	(S)NIBP cuff (paediatrics)Compaitable with Monitor Model : Mindray MEC 2000. Should be FDA'/CE'/DCGI approved	10	nos
1.355	(S)BP Cuff (Adult) Compaitable with Monitor Model : Mindray MEC 2000. Should be FDA'/CE'/DCGI approved	10	nos
1.356	(S)BP Cuff (paediatrics) Compaitable with Monitor Model : Mindray MEC 2000. Should be FDA'/CE'/DCGI approved	10	nos
1.357	(S)SPO2 Probe Compaitable with Monitor Model : STAR 55. Should be FDA'/CE'/DCGI approved	10	nos
1.358	(S)ECG leads Compaitable with Monitor Model : STAR 55. Should be FDA'/CE'/DCGI approved	10	nos
1.359	(S)NIBP cuff (Adult)Compaitable with Monitor Model : STAR 55. Should be FDA'/CE'/DCGI approved	10	nos
1.36	(S)NIBP cuff (paediatrics)Compaitable with Monitor Model : STAR 55. Should be FDA'/CE'/DCGI approved	10	nos
1.361	(S)BP Cuff (Adult) Compaitable with Monitor Model : STAR 55. Should be FDA'/CE'/DCGI approved	10	nos
1.362	(S)BP Cuff (paediatrics) Compaitable with Monitor Model : STAR 55. Should be FDA'/CE'/DCGI approved	10	nos
1.363	(S)SPO2 Probe Compaitable with Monitor Model : Philips MP 30. Should be FDA'/CE'/DCGI approved	10	nos
1.364	(S)ECG leads Compaitable with Monitor Model : Philips MP 30. Should be FDA'/CE'/DCGI approved	10	nos
1.365	(S)NIBP cuff (Adult)Compaitable with Monitor Model : Philips MP 30. Should be FDA'/CE'/DCGI approved	10	nos
1.366	(S)NIBP cuff (paediatrics)Compaitable with Monitor Model : Philips MP 30. Should be FDA'/CE'/DCGI approved	10	nos
1.367	(S)BP Cuff (Adult) Compaitable with Monitor Model : Philips MP 30. Should be FDA'/CE'/DCGI approved	10	nos
1.368	(S)BP Cuff (paediatrics) Compaitable with Monitor Model : Philips MP 30. Should be FDA'/CE'/DCGI approved	10	nos
1.369	ACT Catridges compatible with medtronic ACT machine. Should be FDA'/CE'/DCGI approved	300	nos
1.37	(S)Arterial Pressure monitoring Transducers (Disposable). Should be FDA'/CE'/DCGI approved	100	nos
1.371	(S)Syringe For pressure Injector (Disposable) - Liebel Flarsheim injector 100ml. Should be FDA'/CE'/DCGI approved	10	nos

1.372	(S)SPO2 Probe disposable compaitable with Multi Purpose Monitor PHILIPS A3. Should be FDA'/CE'/DCGI approved	10	nos
1.373	(S)ECG Cable compaitable with Multi Purpose Monitor PHILIPS A3. Should be FDA'/CE'/DCGI approved	10	nos
1.374	(S)NIBP Cuff (Adult) compaitable with Multi Purpose Monitor PHILIPS A3. Should be FDA'/CE'/DCGI approved	10	nos
1.375	(S)NIBP Cuff (Child) compaitable with Multi Purpose Monitor PHILIPS A3. Should be FDA'/CE'/DCGI approved	10	nos
1.376	(S)SPO2 Probe disposable compaitable with Multi Purpose Monitor AGILENT. Should be FDA'/CE'/DCGI approved	10	nos
1.377	(S)ECG Cable compaitable with Multi Purpose Monitor AGILENT. Should be FDA'/CE'/DCGI approved	10	nos
1.378	(S)NIBP Cuff (Adult) compaitable with Multi Purpose Monitor AGILENT. Should be FDA'/CE'/DCGI approved	10	nos
1.379	(S)NIBP Cuff (Child) compaitable with Multi Purpose Monitor AGILENT. Should be FDA'/CE'/DCGI approved	10	nos
1.38	(S)SPO2 Probe disposable compaitable with Multi Purpose Monitor MEC 1200. Should be FDA'/CE'/DCGI approved	10	nos
1.381	(S)ECG Cable compaitable with Multi Purpose Monitor MEC 1200. Should be FDA'/CE'/DCGI approved	10	nos
1.382	(S)NIBP Cuff (Adult) compaitable with Multi Purpose Monitor MEC 1200. Should be FDA'/CE'/DCGI approved	10	nos
1.383	(S)NIBP Cuff (Child) compaitable with Multi Purpose Monitor MEC 1200. Should be FDA'/CE'/DCGI approved	10	nos
1.384	(S)Superior Quality ECG Jelly in bottles of 250 g long lasting . Should be FDA'/CE'/DCGI approved	2000	nos
1.385	(S)Superior Quality ECG ECHO Jelly in bottles of 250 g long lasting . Should be FDA'/CE'/DCGI approved	2000	nos
1.386	(S)Quick Clot - Hydrophilic non woven gauze roll impregnated with a naturally occuring inorganic mineral -Kaolin, Along with a unique direct pressure adhesive bandage packed in a foil pouch for aseptic removal.. Should be FDA'/CE'/DCGI approved	1000	nos
1.387	(S)MORTARA ELI-250 MACHINE cable. Should be FDA'/CE'/DCGI approved	5	nos
1.388	(S)MORTARA ELI 350 MACHINE cable. Should be FDA'/CE'/DCGI approved	5	nos
1.389	(S)CHEST BULBS ADULT. Should be FDA'/CE'/DCGI approved	5	nos
1.39	(S)CHEST BULBS PEDIATRICS. Should be FDA'/CE'/DCGI approved	5	nos
1.391	(S)LIMB LEADS ADULT. Should be FDA'/CE'/DCGI approved	5	nos
1.392	(S)LIMB LEADS PEDIATRICS. Should be FDA'/CE'/DCGI approved	5	nos
1.393	(S) BioCare ECG recorder paper. Should be FDA'/CE'/DCGI approved	2000	nos
1.394	(S) Mortara ELI-250 ECG Recorder Head. Should be FDA'/CE'/DCGI approved	5	nos
1.395	(S) Mortara ELI-350 ECG Recorder Head. Should be FDA'/CE'/DCGI approved	5	nos
1.396	(S) Mortara TMT Paper. Should be FDA'/CE'/DCGI approved	1000	nos
1.397	(S) Mortara ELI-350 ECG Paper.. Should be FDA'/CE'/DCGI approved	1000	nos
1.398	(S) Mortara ELI-250 ECG Paper.. Should be FDA'/CE'/DCGI approved	1000	nos
1.399	(S) Edan Multichannel Machine Paper. Should be FDA'/CE'/DCGI approved	1000	nos
1.4	(S) NIHON KODEN Multichannel Machine Paper Roll Type. Should be FDA'/CE'/DCGI approved	1000	nos
1.401	(S) AG/AGCL IMPORTED PATIENT PREJELLED ELECTRODES FOR LONG-TERM MONITORING SUPERIOR QUALITY . Should be FDA'/ CE'/DCGI approved	500000	nos
1.402	(S)PATIENT TUBING SYSTEM COMPATIBLE WITH MAQUET S VENTILATOR. Should be FDA'/CE'/DCGI approved	10	nos
1.403	(S)SPOT TEST KITS for HIV . Should be FDA'/CE'/DCGI approved	300	nos
1.404	(S)SPOT TEST KITS for HBsAg . Should be FDA'/CE'/DCGI approved	300	nos
1.405	(S)SPOT TEST KITS for HCV . Should be FDA'/CE'/DCGI approved	300	nos

1.406	(S)StickyMats'/Tacky Mats 1 to 30 on 30 Peel Tags, anti ststic, with Holding Stick Time : > 1000 mins, Price per box to be quoted with number of units per box. Should be FDA'/CE'/DCGI approved	300	nos
1.407	(S) DISPOSABLE STERILE DRAPES FOR CORONARY ANGIOGRAPHY'/ANGIOPLASTY COMPATIBLE WITH Bilateral FEMORAL and RADIAL APPROACHES with absorbable material.Sample to be supplied for testing	1000	nos
1.408	(S) DISPOSABLE STERILE DRAPES FOR PERMANENT PACEMAKER IMPLANTATION L and R Pectoral Approaches.Sample to be supplied for testing	250	nos
1.409	(S) Tissue paperbased disposable bed covers in continious rolls of Width "90Cm" for use on trolleys & Cardiac Cath Lab Tables..Sample to be supplied for testing	5000	nos
1.41	(S) DISPOSABLE STERILE ANGIOPLASTY DRAPES KITS WITH A. One Unit -DRAPE ANGIOPLASTY COMPATIBLE WITH Bilateral FEMORAL and RADIAL APPROACHES with absorbable material B. Three Units DISPOSABLE STERILE GOWNS C. One Unit Disosable Centre Hole towel D. One Unit Cath Trolley Drape.Sample to be supplied for testing	5000	kit
1.411	(S) DISPOSABLE GOWN & PYJAMA SET FOR CATH LAB PROCEDURE (SMALL/MEDIUM /LARGE).Sample to be supplied for testing	2000	nos
1.412	(S) DISPOSABLE GAUZE PIECES in (ETO Sterilizable)PACKS OF 10 for use with Cath Lab procedures.Sample to be supplied for testing	10000	nos
1.413	(D) BATTERY OPERATED AUTOMATED DIGITAL BP APPARATUS with ADULT,LARGE ADULT and PEDIATRIC CUFFS with 3 yr warranty.Item to be Demonstrated to departmental selection committee.	10	nos
1.414	(D) DIGITAL BP APPARATUS HEAVY DUTY, with ADULT,LARGE ADULT and PEDIATRIC CUFFS RUNNING IN AC POWER FOR USE IN OPD AND WARD with 3 yr warranty.Item to be Demonstrated to departmental selection committee.	2	nos
1.415	(D)DIGITAL WEIGHING SCALE FOR PATIENT WEIGHT with 3 yr warranty.Item to be Demonstrated to departmental selection committee.	2	nos
1.416	(D) SYRINGE INFUSION PUMPS with 3 yr warranty.Item to be Demonstrated to departmental selection committee.	4	nos
1.417	(D)MULTIPARAMETER MONITORS ECG, SPO2,NIBP with 3yr warrranty and 2 sets of cables.Item to be Demonstrated to departmental selection committee.	10	nos
1.418	(D) OT LIGHT- FLOOR MOUNTED, LED SOURCE , Brightness and Focus ADJUSTABLE with 3 yr warranty.Item to be Demonstrated to departmental selection committee.	3	nos
1.419	(D) SMALL SIZE Packing material compatible with automated sealing machine for ETO sterilization of consumables..Item to be Demonstrated to departmental selection committee.	10	nos
1.42	(D) INTERMEDIATE SIZE Packing material compatible with automated sealing machine for ETO sterilization of consumables..Item to be Demonstrated to departmental selection committee.	10	nos
1.421	(D) LARGE SIZE Packing material compatible with automated sealing machine for ETO sterilization of consumables..Item to be Demonstrated to departmental selection committee.	10	nos
1.422	(D) Battery Operated Pocket Sized SpO2 Probes.Item to be Demonstrated to departmental selection committee.	5	nos
1.423	(D)Compact Disk- Recordable (CD-R) with hard shell box.Item to be Demonstrated to departmental selection committee.	2000	nos
1.424	(D)DVD- Recordable (DVD-R) with hard shell box.Item to be Demonstrated to departmental selection committee.	200	nos

1.425	(S)CD Sticker Labels with Department of Cardiology ,JIPMER Stickers, With Space for printing Angio No and CD No.Item to be Demonstrated and discussed to departmental selection committee.	3000	nos
1.426	(S)Permanent Marker Pens- Thin Script- Black.Item to be Demonstrated and discussed to departmental selection committee.	50	nos
1.427	(S)White Board Markers Black and Blue.Item to be Demonstrated and discussed to departmental selection committee.	50	nos
1.428	(S)Batteries Compatible with Zoll M Series defibrillators. Should be FDA'/CE'/DCGI approved	15	nos
1.429	(D)Catridge for HP 1020 series. Should be FDA'/CE'/DCGI approved	5	nos
1.43	(S)Disposable Paper based Sterile Front Buttoned Black/ Brown/ Dark blue half sleeve shirt and pyjama for cath lab patient dress. Should be FDA'/CE'/DCGI approved	1000	nos
1.431	(S)Rechargable Ni- MH Batteries 1.2 to 1.5 V AA 2500 mAh with 80 batteries and one fast charger which can charge 4 batteries at a time should be supplied with every 8 batteries.. Should be FDA'/CE'/DCGI approved	80	nos
1.432	(D)Long term external loop recorder with capability to record at least one ECG lead for a period of at least 7 days. Recording patch / electrodes should be water resistant allowing patient to bathe. Episodes should be identified automatically and it should also be possible to record patient triggered episodes. Transmission should be wireless and monitored in a 24x7 staffed monitoring center. Quoted cost should include disposable electrode set, recorder, hand-held activator, data transmission and reporting for the period of recording.. Should be FDA'/CE'/DCGI approved	10	nos
1.433	(S)Disposable, sterile cover for Fluoroscopy system Image Intensifier. Should fit Image Intensifier of different sizes and should have elastic band for easy retention.. Should be FDA'/CE'/DCGI approved & Item to be Demonstrated to departmental selection committee	100	nos
1.434	(S)Washable and Autoclavable Cut Shoes (Clogs) Specially designed for wearing in Operation rooms /ICU's wear slippers at least 12 inch foot length.(roomy fit). Should have non collapsible sole to support the arch of foot. The brochure mentioning the details of the product should be submitted .Mention various sizes available.(SPR).Item to be Demonstrated to departmental selection committee	150	nos

Additional Tender Conditions:

1. ITEMS WITH (Prefix "S") sample MUST be submitted to the department with item number superscripted within 15 days of online tender opening
2. All implantable devices should be Licensed for sale in India by DCGI, and should be USFDA approved failing which Human Clinical Trial Data Published in Indexed Journals should be submitted
3. All implantable devices should be supplied only by companies that has supplied to major hospitals (500 beds) in India or abroad (proof to be enclosed)
4. ITEMS WHICH COME IN PACKS- mention price per unit and pack size.
5. Patient Size Variable Implantable Devices and Consumables (Prefix EX) (Stents/ ASD/VSD/PDA/Balloons/Coils/ etc- the supplier should exchange the desired size if one size is exhausted so as to avoid device patient mismatch till the last item is used.
6. ITEMS WHICH HAS A PREFIX (D) has to be demonstrated by the company to the departmental selection committee.
7. Items supplied should have an Expiry Date of more Than 12 months from date of supply.
8. All the items should be Should be FDA'/CE'/DCGI approved

Section-III

GENERAL INSTRUCTIONS TO TENDERERS

1. **Period of contract**:

The contract shall initially be for a period of **one year** that may be extended to **6 months**. The rates approved shall remain unchanged during the period of contract.

2. Bid Security (Earnest Money Deposit)

i. In compliance with the OM No. F.9/4/2020-PPD dated 12.11.2020 on the subject ―Bid Security/ Earnest Money Deposit‖, issued by the Procurement Policy Division, Department of Expenditure, Ministry of Finance, Government of India, there is no need for bidders to pay Bid Security/ Earnest Money Deposit for participating in this tender.

ii. However, in lieu of Bid Security, the bidder must print this ―Bid Security Declaration‖ on his/her firm‘s letterhead duly sign the undertaking and upload the document in the fee cover.

iii. The bidder hereby declares that they accept the condition that if they withdraw or modify their bids during period of validity etc., they will be suspended for a period of two years from participating in any tender invited by JIPMER, Puducherry with effect from the date of their withdrawal or modification of their bid.

3. <u>Eligibility conditions of bidders:</u>

i. The Tenderer must be a domestic manufacturer that is a ‘Class-I local supplier' or a ‘Class-II local supplier’ as defined under Public Procurement (Preference to Make in India), order 2017 of MoC and I (DIPP), Govt. Of India, as further amended by orders of even number dated 28.05.2018, 29.05.2019, 04.06.2020 and 16.09.2020. In case the manufacturer does not quote directly, they may authorize an agent as per proforma of Manufacturer authorization form as given in the Tender enquiry document to quote and enter into a contractual obligation.

ii. In compliance with order (Public Procurement No.1) No. 6/18/2019-PPD dated 23rd July 2020 issued by the Public Procurement Division, Dept. of Expenditure, Min of Finance under Rule 144(xi) of GFR 2017 any bidder 'from a country which shares a land border with India will be eligible to bid in this tender only if the bidder is registered with the Competent Authority.

a. "Bidder" (including the term 'tenderer', 'consultant' or 'service provider’ in certain contexts) means any person or firm or company, including any member of a consortium or joint venture (that is an association of several persons, or firms or companies), every artificial juridical person not falling in any of the descriptions of bidders stated hereinbefore, including any agency branch or office controlled by such person, participating in a procurement process.

b. "Bidder from a country which shares a land border with India" for the purpose of this Order means: -

I. An entity incorporated, established or registered in such a country; or
II. A subsidiary of an entity incorporated, established or registered in such a country; or
III. An entity substantially controlled through entities incorporated, established or registered in such a country; or
IV. An entity whose beneficial owner is situated in such a country; or
V. An Indian (or other) agent of such an entity; or
VI. A natural person who is a citizen of such a country; or
VII. A consortium or joint venture where any member of the consortium or joint venture falls under any of the above

c. The beneficial owner for the purpose of (5.2.2) above will be as under:

I. In case of a company or Limited Liability Partnership, the beneficial owner is the natural person(s), who, whether acting alone or together, or through one or more juridical person, has a controlling ownership interest or who exercises control through other means.

Explanation—

- "Controlling ownership interest" means ownership of or entitlement to more than twenty-five per cent, of shares or capital or profits of the company;
- "Control" shall include the right to appoint majority of the directors or to control the management or policy decisions including by virtue of their shareholding or management rights or shareholders agreements or voting agreements;

II. In case of a partnership firm, the beneficial owner is the natural person(s) who, whether acting alone or together, or through one or more juridical person, has ownership of entitlement to more than fifteen percent of capital or profits of the partnership.
III. In case of an unincorporated association or body of individuals, the beneficial owner is the natural person(s), who, whether acting alone or together, or through one or more juridical person, has ownership of or entitlement to more than fifteen percent of the property or capital or profits of such association or body of individuals.
IV. Where no natural person is identified under (a) or (b) or (c) above, the beneficial owner is the relevant natural person who holds the position of senior managing official;
V. In case of a trust, the identification of the beneficial owner(s) shall include the identification of the author of the trust, the trustee, the beneficiaries with 15% or more interest in the trust and any other natural person exercising ultimate effective control over the trust through a chain of control or ownership.

d. An Agent is a person employed to do any act for another, or to represent another in dealings with third person.

e. In case of turnkey contracts, the successful bidder shall not be allowed to sub-contract works to any contractor from a country which shares a land border with India unless such contractor is registered with the Competent Authority.

f. Bidders must certify compliance with the above mentioned provisions in the tender form as per Section VII.
g. The bidder should have successfully completed the delivery of the item of at least 25% of the quantity required in this tender to a government hospital in the last 3 years.
h. The bidder or the manufacturer must have an average annual turnover of at least Rs.1,00,00,000/-(Rupees one crore only) each during the last three financial years i.e. 2017-18, 2018-19, 2019-20.
i. Bidder should have ISO Certification.

4. Purchase Preference

i. The Procurement of goods and services under this e-tender will be regulated as per the applicable provisions of Public Procurement (Preference to Make in India), order 2017 of MoC and I (DIPP), Govt. Of India, as further amended by orders of even number dated 28.05.2018, 29.05.2019, 04.06.2020 and 16.09.2020. Salient portions of the order are reproduced in Appendix A of this tender document by way of information. Bidders are advised to see the original orders and satisfy themselves that they qualify to participate in the tender. Bidders who are claiming eligibility to participate in this tender must submit a certificate in format given in Appendix A along with documentary evidence in support of their claim wherever necessary failing which their bid will be summarily rejected. The purchaser reserves the right to give preference to the 'Class-I local supplier'.

ii. The Purchaser reserves the right to give the purchase preference to small-scale sectors, Micro and small scale enterprises etc. as per the instruction in vogue while evaluating, comparing and ranking the responsive Tenders as given in the MSMED Act 2006 reproduced below:

a. In exercise of powers conferred in Section 11 of the Micro, Small and Medium Enterprises Development (MSMED) Act 2006, the Government has notified a new Public Procurement Policy for Micro & Small Enterprises effective from 1st April 2012. The policy mandates that 25% of procurement of annual requirement of goods and services by all Central Ministries/Public Sector Undertakings will be from the micro and small enterprises. The Government has also earmarked a sub-target of 4% procurement of goods & services from MSEs owned by SC/ST entrepreneurs out of above said 25% quantity.

b. In accordance with the above said notification, the participating Micro and Small Enterprises (MSEs) in a tender, quoting price within the band of L1+15% would also be allowed to supply a portion of the requirement by bringing down their price to the L1 price, in a situation where L1 price is from someone other than an MSE. Such MSEs would be allowed to supply up to 25% of the total tendered value. In case there are more than one such eligible MSE, the 25% supply will be shared equally. Out of 25% of the quantity earmarked for supply from MSEs, 5% quantity is earmarked for procurement from MSEs owned by SC/ST entrepreneurs. However, in the event of failure of such MSEs to participate in the Tender process or meet the

tender requirements and the L1 price, the 5% quantity earmarked for MSEs owned by SC/ST entrepreneurs will be met from other participating MSEs.

c. The MSEs fulfilling the prescribed eligibility criteria and participating in the Tender shall enclose with their tender a copy of their valid registration certificate with District Industries Centres or Khadi and Village Industries Commission or Khadi and Village Industries Board or Coir Board or National Small Industries Corporation or any other body specified by Ministry of Micro and Small enterprises in support of their being an MSE, failing which their tender will be liable to be ignored.

d. Special provision for Micro and Small Enterprise owned by women: – Out of the total annual procurement from Micro and Small Enterprises, 3 per cent from within the 25 per cent target shall be earmarked for procurement from Micro and Small Enterprises owned by women.

e. Note: "If the bidder is a MSME, it shall declare in the bid document the Udyog Aadhar Memorandum Number issued to it under the MSMED Act, 2006. If a MSME bidder do not furnish the UAM Number along with bid documents, such MSME unit will not be eligible for the benefits available under Public Procurement Policy for MSEs Order 2012.

5. Demonstration of sample

The tenderer must give a sample within **ten** days of the closing of online submission of bids failing which the bids will be rejected.

6. Instructions for the filling the tender form

i. E-Tender form shall be completed in all respect, signed in full and stamped at appropriate places and initialed and stamped on all remaining pages. Incomplete or e-tenders without tender processing fee, bid security declaration, Make-in-India self-certification as in Appendix-A only by original manufacturer, GFR 144 (xi) compliance certificate as in Section-X shall be treated as invalid.

ii. Bidders have to ensure that all the documents are properly filled.

iii. Conditional tenders are liable to be rejected.

iv. Bids received and found valid will be evaluated by JIPMER to ascertain the complete work/services under the specification and documents. The bidder should take care to submit all the information sought by JIPMER in prescribed formats.

v. Incomplete bids, bids in paper format, conditional bids, telephonic bids or tenders submitted after the due date and time will not be considered and summarily rejected. Vendors are, therefore, advised to submit their bids well on time.

vi. The bidder can quote for one or more items mentioned in the list. Bidder has to give all details (HSN, MSME, Make-in-India, make/brand, model, pack size and remark) mentioned in BOQ for all quoted items, failure of that the bid will be rejected summarily.

7. Submission of tenders

The bidders must ensure that they submit the **on-line bids** within the scheduled closing date & time.

8. Late Tender:

There is NO PROVISION of uploading late tender beyond stipulated date & time in the e- tendering system.

9. Alteration and Withdrawal of Tender

i. The bidder, after submitting its bid, is permitted to alter/modify its bid, within the deadline for submission of bids. Alterations/modifications to bids received after the prescribed deadline will not be possible on the e-tender portal.

ii. No tender should be withdrawn or modified after the deadline for submission of tender and before expiry of the tender validity period. If a bidder withdraws or modifies the tender during this period, it will result in the tenderer being suspended from participating in all tenders invited by JIPMER for a period of two years with effect from the date of withdrawal, as per the terms and conditions of bid security declaration furnished by the bidder in its tender.

10. Preparation of e-tenders

This is a Two-Bid Tender system, consisting of the **Techno-Commercial Bid** and **Price Bid** that are to be uploaded in the prescribed formats in the e-tendering portal. The tender(s) shall only be submitted online as mentioned below:

I. Techno-commercial Bid shall comprise

a. Fee Cover

i. **E-tender Processing fee** of Rs.590/- (Rupees five hundred and ninety only) inclusive of 18% GST payment **receipt** duly self-attested and rubber stamped should be uploaded.

ii. Scanned copy (100 or 200 DPI) in pdf format of **Bid Security Declaration** as per format attached in Section-IX

b. In the cover named "Prequal/Technical" the scanned copy in pdf format of the following documents are to be uploaded:

i. The **Bidder's Profile** as in Section-VI, **Tender Form** as in Section-VII must be downloaded duly filled signed and stamped, and an **Authorization letter** for signing tender documents if a person other than the Owner, Partner, Managing Director is signing the documents, must be uploaded as a single PDF file.

ii. **Copies of Supply orders/Completion certificate** in support of Eligibility condition 1 and 2 and **Copies of abridged Annual report of last 03 years** (Income tax return acknowledgement, Assets and Liabilities, Balance sheet and Profit & Loss Account) must be uploaded as a single PDF file

iii. **Manufacturer's Authorization letter** in company letterhead in format in given Section-VI and, in case the bidder is empaneled by the Competent Authority under **GFR 144 (xi) (mandatory)** a copy of the same or **GFR 144(xi) compliance certificate** as in Section-X must be uploaded.

iv. Copy of Self Certified **GST registration certificate** and Copy of **PAN Card** must be uploaded as a single PDF file.

v. **Bank Details** (Beneficiary name, Bank name, Account number, IFSC code, Branch address on letterhead) and **Check list** as in Section-VIII in the prescribed format duly filled and signed must be uploaded as a single PDF file.

vi. Self-certification for supporting the claim to be a local supplier under the "Public Procurement preference to Make in India" order in format as in **Appendix-A (mandatory)** and any other document that the bidder wishes to submit as a single PDF file.

vii. A **file mentioning the list of items without price** for which bidder is quoting must be uploaded as a single PDF file.

II. <u>Price Bid:</u>

Prices are to be quoted in the prescribed Price Bid format provided in the e-tender portal using the BOQ template only. The price should be quoted for the **<u>accounting unit</u>** indicated in the e-tender document.

<u>Note:</u>

i) The bidder has to be diligent while filling up the Techno-Commercial Bid and Price Bid provided in prescribed formats and must not tamper with the contents of the sheets.

ii) Bidders must ensure that the documents uploaded in pdf format are legible.

iii) It is the responsibility of bidder to go through the Tender document to ensure furnishing all required documents in addition to above, if any.

iv) The Make in India self-certification as in Appendix-A must be given and uploaded by original manufacture. The distributor can upload the Appendix-A given by original manufacture.

v) ITE- Item-wise Eligibility Sheet should be downloaded, the items that the bidder wishes to quote must be selected as "Eligible", and this "ITE file" must also be uploaded for the price bid to be considered by the system. The selected items will be displayed once uploaded and the bidder can verify that all items he wishes to quote for, are present in the list

vi) A person signing (manually or digitally) the tender form or any documents forming

part of the contract on behalf of another shall be deemed to warrant that he has authority to bind such other persons and if, on enquiry, it appears that the persons so signing had no authority to do so, the purchaser may, without prejudice to other civil and criminal remedies, cancel the contract and hold the signatory liable for all cost and damages.

vii) A bid, which does not fulfill any of the above requirements and/ or give evasive information/reply against any such requirement, shall be liable to be ignored.

viii) Tender sent by fax/telex/cable shall be ignored.

11. Digital Signing of Tender

The tenderers shall submit their tenders as per the instructions contained as above. Tenders shall be uploaded with all relevant tender documents in the prescribed format. The relevant tender documents should be uploaded by an authorized person having Class 3 digital signature certificate.

12. Tender currencies.

The tender shall be quoted only in INR.

13. Additional information and instruction on GST:

If the Tenderer desires to ask for GST or any other taxes to be paid extra, the same must be specifically stated. In the absence of any such stipulation, the price will be taken inclusive of such taxes and no claim for the same will be entertained later. The rate of GST quoted in the tender shall be taken for price comparison. However, the rate of GST quoted in the tender or the actual rate of GST applicable, whichever is lower shall be payable by the purchaser. The supplier can charge a higher GST than quoted in the tender only if the rate of GST was revised by the government after the tender closing date.

14. Bid Security (Earnest Money Deposit)

i. In compliance with the OM No. F.9/4/2020-PPD dated 12.11.2020 on the subject ―Bid Security/ Earnest Money Deposit‖, issued by the Procurement Policy Division, Department of Expenditure, Ministry of Finance, Government of India, there is no need for bidders to pay Bid Security/ Earnest Money Deposit for participating in this tender.

ii. However, in lieu of Bid Security, the bidder must print this ―Bid Security Declaration‖ on his/her firm‘s letterhead duly sign the undertaking and upload the document in the fee cover.

iii. The bidder hereby declares that they accept the condition that if they withdraw or modify their bids during period of validity etc., they will be suspended for a period of two years from participating in any tender invited by JIPMER, Puducherry with effect from the date of their withdrawal or modification of their bid.

15. <u>Tender opening</u>

i. The Tender Inviting Authority will open the e-tenders at the specified date and time and at the specified place as indicated in the NIT. In case the specified date of tender opening falls on / is subsequently declared a holiday or closed day for the Tender Inviting Authority, the tenders will be opened at the appointed time and

place on the next working day.

ii. Authorized representatives of the tenderers, who have submitted tenders on time, may attend the tender opening provided they bring with them letters of authority from the corresponding tenderers. The tender opening official(s) will prepare a list of the representatives attending the tender opening. The list will contain the representatives' names & signatures and corresponding tenderers' names and addresses.

iii. This being a Two-Bid Tender system, the **Techno-Commercial Tenders** are to be opened in the first instance, at the prescribed time and date as indicated in NIT. These Tenders shall be scrutinized and evaluated by the competent committee/ authority with reference to parameters prescribed in the tender document. During the Techno- Commercial Tender opening, the tender opening official(s) will read the salient features of the tenders like brief description of the goods offered, delivery period, bid security declaration and any other special features of the tenders, as deemed fit by the tender opening official(s). Thereafter, in the second stage, the Price bids of only the Techno - Commercially acceptable offers (as decided in the first stage) shall be opened for further scrutiny and evaluation on a date notified after the evaluation of the Techno-Commercial tender.

16. Scrutiny and evaluation of tenders

A. Basic Principle

Tenders will be evaluated on the basis of the terms & conditions already incorporated in the tender enquiry document, based on which tenders have been received and the terms, conditions etc. mentioned by the tenderers in their tenders. No new condition will be brought in while scrutinizing and evaluating the tenders.

B. Scrutiny of Tenders

i. The Purchaser will examine the Tenders to determine whether they are complete, whether any computational errors have been made, whether required sureties have been furnished and, whether the documents uploaded are in legible form.

ii. The Purchaser's determination of a Tender's responsiveness is to be based on the contents of the tender itself without recourse to extrinsic evidence.

iii. The tenders will be scrutinized to determine whether they are complete and meet the essential and important requirements, conditions etc. as prescribed in the tender document. The tenders, which do not meet the basic requirements, are liable to be treated as non-responsive and will be summarily ignored.

iv. The following are some of the important aspects, for which a tender shall be declared non-responsive during the evaluation and will be ignored;

   i. Tender validity is shorter than the required period.
   ii. Non-submission of Bid Security Declaration.
   iii. Non-submission of receipt of tender processing fee.
   iv. Non-submission of self-certification in format as given in Appendix-A only by original manufacturer, for determining eligibility to participate in the tender under the "Public Procurement preference to Make in India" order.
   v. Non-submission of GFR-144 (xi) compliance certificate.

vi. Tenderer has not agreed to give the required performance security of required amount in an acceptable form.
vii. Non-submission of samples within ten days of the closing of online submission of bids
viii. Poor/ unsatisfactory past performance.
ix. Tenderers who stand de-registered/banned/blacklisted by any Central Govt. Ministries/Departments/Hospitals/Institutes.
x. Tenderer is not eligible as per tender conditions.
xi. Tenderer has not quoted for the entire quantity as specified in the List of Requirements/ BOQ for the item quoted.
xii. Non-submission of all details of quoted items (HSN, MSME, Make-in-India, make/brand, model, pack size and remark).
xiii. Tenderer has not agreed to other essential condition(s) specially incorporated in the tender enquiry, like delivery terms, delivery schedule, terms of payment, liquidated damages clause, warranty clause, dispute resolution mechanism applicable law.

17. Minor Informality/Irregularity/Non-Conformity

If during the preliminary examination, the purchaser find any minor informality and/or irregularity and/or non-conformity in a tender, the purchaser may waive the same provided it does not constitute any material deviation and financial impact and, also, does not prejudice or affect the ranking order of the tenders. Wherever necessary, the purchaser will convey its observation on such 'minor' issues to the tenderer by registered/speed post etc. asking the tenderer to respond by a specified date. If the tenderer does not reply by the specified date or gives evasive reply without clarifying the point at issue in clear terms, that tender will be liable to be ignored.

18. Award of work

i. The selection of the agency will be at the sole discretion of the JIPMER who reserves its right to accept or reject any or all the proposals without assigning any reason thereof.
ii. The lowest bid will be evaluated on the basis of annual estimated amount as quoted by bidders in the quotation including taxes.
iii. Upon evaluation of offers the decision on the award of contract will be intimated to the successful bidder.
iv. The annual estimate is given only as an indication. The actual quantity procured may increase or decrease. No assurance is given that the quantity stated will actually be procured.

Section-IV

GENERAL CONDITIONS OF CONTRACT

1. Price of goods

The rate quoted in the e-tender will be fixed for the whole contract period.

2. Technical Specifications and Standards

The Goods & Services to be provided by the supplier under this contract shall conform to the technical specifications mentioned in "Technical Specification" under Section II.

3. Terms of Delivery

i. Goods shall be delivered by the supplier within 30 days of issue of supply order. Please note that the time shall be the essence of the contract.

ii. Any unexcused delay by the supplier in maintaining its contractual obligations towards delivery of goods and performance of services shall render the supplier liable to any or all of the following sanctions:

a) Imposition of liquidated damages,
b) Forfeiture of its performance security and
c) Termination of the contract for default.

4. Liquidated Damages

If the supplier fails to deliver any or all of the goods or fails to perform the services within the time frame(s) incorporated in the contract, the Purchaser/Consignee shall, without prejudice to other rights and remedies available to the Purchaser/Consignee under the contract, deduct from the contract price, as liquidated damages, a sum equivalent to 0.5% per week of delay or part thereof on delayed supply of goods, installation, commissioning and/or services until actual delivery or performance subject to a maximum of 10% of the contract price. Once the maximum is reached Purchaser/Consignee may consider termination of the contract as per GCC 11. Since the Liquidated damages are in virtue of non-performance of services, it will attract GST or any other applicable taxes which in turn shall be deducted from the bidder.

5. Performance Security

The successful bidder shall have to deposit an amount 3% of the value of contract as Performance Security Deposit (PSD) within two weeks after award of contract, through SBI Collect available on JIPMER website. In the event of any failure /default of the supplier with or without any quantifiable loss to the purchaser, the amount of the performance security is liable to be forfeited.

Subject to the condition mentioned above the Performance Security will be released without any interest to the supplier on completion of the supplier's all contractual obligations

including the warranty obligations and extension of time (with or without Liquidated Damages).

6. <u>Payment Procedure</u>

The contractor shall submit bill in triplicate upon satisfactory supply of goods. Bill must be raised based on the rate quoted in e-tender. Every effort shall be made to ensure that the payment will be made within 45 days of submission of bill. No advance payment will be considered. TDS will be deducted as per provision of Income Tax Act, GST Acts and other statutes as relevant.

7. Risk Clause

The contractor shall at all times have standby arrangements for carrying out the work under the contract, in case of any failure of the existing arrangements. JIPMER reserves the right for termination of the contract at any time by giving 30 days written notice, if the items delivered are found to be unsatisfactory and also has the right to award the contract to the next higher bidder willing to supply the item at the cost, risk and responsibilities of contractor and excess expenditure incurred on account of this will be recovered by JIPMER from the contractor's Performance Security Deposit or pending bills or by raising a separate claim.

8. Termination clause:

During the period of agreement if it is found that the agency is not providing proper services, the JIPMER reserves rights to make the vendor forfeit the security deposit deposited with JIPMER or part thereof in favour of JIPMER and agreement will be terminated after giving 30 days' notice. Furthermore, in such situations, tender can be allotted to second lowest bidder and the difference in cost shall be recovered from the earlier vendor who is breach of the contract. In addition in case it is found that the supplier is charging by fraudulent means or indulging in criminal activities the contract will be terminated immediately.

9. Jurisdiction of the courts

Jurisdiction of the courts for settlement of disputes:- Jurisdiction for the settlements of disputes if any is Puducherry only.

In above mentioned conditions Director JIPMER reserves all the rights.

Section-V

BIDDER'S PROFILE

This form duly filled and signed by authorized representative of the bidder and the scanned copy must be uploaded online

1.	Name & Designation of the contact person	
2.	Name and Address of the Tenderer	
3.	Phone No a) Land line number (functional between 9 am and 5pm)	
4.	Mobile No of contact person (available from 9am to 6pm)	
5.	Email ID of the Tenderer	
6.	Email ID of the contact person	
7.	Local supplier/Distributor in Chennai/ Puducherry or any other place (complete address must be written)	
8.	Manufacture Name	
9.	Manufacture Address	
10	Whether Tenderer is registered MSE Manufacture the product quoted. (If registered MSE, submit copy of the Udyog Aadhaar certificate or Equivalent Certificate)	Yes / No

If there is any change in the above details, I will immediately intimate you by speed post or fax or email

I.....hereby declare that the details given above are true to the best of my knowledge and I have thoroughly read and understood the terms and conditions of the tender and shall abide by the rules,

Signature

Dated: (Name and Designation & Seal)

NB: This form must be duly filled in by an authorized person

Section-VI

MANUFACTURER'S AUTHORISATION FORM

(Letterhead)

The Director,
JIPMER, Puducherry

Dear Sir,

Ref: Your TE document No:_____dated: _____

We,_____who are proven and reputable manufacturers of_____(name and description of the goods offered in the Tender) having factories at_, hereby authorize Messrs. _____(name and address of the agent) to submit a tender, process the same further and enter into a contract with you against your requirement as contained in the above referred TE documents for the above goods manufactured by us.

We also state that we are not participating directly in this tender for the following reason(s): _____(Please provide reason here).

We further confirm that no supplier or firm or individual other than Messrs_____(name and address of the above agent) is authorised to submit a tender, process the same further and enter into a contract with you against your requirement as contained in the above referred TE documents for the above goods manufactured by us. We also hereby extend our full warranty, CMC/AMC as applicable as per the General Conditions of Contract, read with modification, if any, in the Special Conditions of Contract for the goods and services offered for supply by the above firm against this TE document. We also hereby confirm that we would be responsible for the satisfactory execution of contract placed on the authorised agent.

We also confirm that the price quoted by our agent shall not exceed the price which we would have quoted directly"

Yours faithfully,
[Signature with date, name, designation and Email]
for and on behalf of Messrs _____
[Name & address of the manufacturers]

Note:

(1) This letter of authorization should be on the letter head of the manufacturing firm and should be signed by a person competent and having the power of attorney to legally bind the manufacturer.
(2) Original letter may be sent.
(3) The purchaser reserves the right to verify this document with its signatory.

SECTION–VII
TENDER FORM
(On Firms' Letter Head)

To Date_____
The Director
JIPMER, Puducherry 605006

Ref. Your TE document No._____dated _____

We, the undersigned have examined the above mentioned TE document, including amendment/corrigendum No._____, dated_____(if any), the receipt of which is hereby confirmed. We now offer to supply and deliver_____(Description of goods and services) in conformity with your above referred document **for the sum as shown in the price schedules attached herewith and made part of this tender**. If our tender is accepted, we undertake to supply the goods and perform the services as mentioned above, in accordance with the delivery schedule specified in the List of Requirements.

We further confirm that, if our tender is accepted, we shall provide you with a performance security of required amount in an acceptable form in terms of GCC, Special Conditions of Contract", for due performance of the contract.

We have read the clause regarding restrictions on procurement from a bidder of a country which shares a land border with India and we certify that this bidder is not from such a country/ from such a country and has been registered with the Competent Authority and a copy of the valid registration by the Competent Authority is attached as evidence of the same (Strike out what is not applicable). In case there are Turnkey works to be carried out this bidder will not sub-contract any work to a contractor from such countries unless such contractor is registered with the Competent Authority. We hereby certify that this bidder fulfills all requirements in this regard and is eligible to be considered.

We agree to keep our tender valid for acceptance as required in the GIT, Special Instructions to Tenderers" or for subsequently extended period, if any, agreed to by us. We also accordingly confirm to abide by this tender up to the aforesaid period and this tender may be accepted any time before the expiry of the aforesaid period. We further confirm that, until a formal contract is executed, this tender read with your written acceptance thereof within the aforesaid period shall constitute a binding contract between us.

We further understand that you are not bound to accept the lowest or any tender you may receive against your above-referred tender enquiry.

We confirm that we do not stand deregistered/banned/blacklisted by any Govt. Authorities in the last 7 years.

We confirm that we fully agree to the terms and conditions specified in above mentioned TE document, including amendment/ corrigendum if any.

(Signature with date)
(Name and designation)

Duly authorized to sign tender for and on behalf of

Section-VIII

CHECK LIST FOR SUBMISSION OF TENDER

(To be filled by the tenderer and submitted along with the bid)

Sl. No.	Particular	Yes / No
1	i. Tender Processing Fee (Mandatory for all bidders) ii.Bid security declaration	
2	Copies of Supply orders/Completion certificate in support of Eligibility condition 2(ii).	
3	Copy of PAN Card	
4	Copy of ISO Certification of bidder, ISI/CE Certification of each product wherever asked	
5	Copies of last three years Income Tax Return statement with Balance sheets & Profit & Lose A/c i.e. 2017-18, 2018-19, 2019-20.	
6	Copy of GST Registration Certificate	
7	Authorization letter for signing tender documents if a person other than the Owner, Partner, Managing Director is signing/uploading the tender	
8	Tender form as in Section – VII duly signed and stamped	
9	Manufacturer's authorization form in format as in Section VI and	
10	Self-certification for supporting the claim to be a local supplier under the "Public Procurement preference to Make in India" order in format as in Appendix A	
11	A copy bidder's empanelment by the Competent Authority under GFR 144 (xi) or GFR 144 (xi) compliance certificate	
12	A PDF file containing list of all items quoted by the bidder without price bid in technical cover	
13	All details of the items (HSN, MSME, Make-in-India, make/brand, model, pack size and remark) quoted by the bidder.	
14	Any other document(s) enclosed (To be specified)	

I/We certify that the information furnished above is true and correct. The terms and conditions are acceptable to us and have the authority to bid a tender.

Signature of the owner/
Managing Partner/Director

Name:
Seal:

Date:
Place:

SECTION – IX

Bid Security Declaration
(To printed on the Firm's letterhead)

(In compliance with the OM No. F.9/4/2020-PPD dated 12.11.2020 on the subject ―Bid Security/ Earnest Money Deposit‖, issued by the Procurement Policy Division, Department of Expenditure, Ministry of Finance, Government of India, there is no need for bidders to pay Bid Security/ Earnest Money Deposit for participating in this tender. However, in lieu of Bid Security, the bidder must print this ―Bid Security Declaration‖ on his/her firm's letterhead duly sign the undertaking and upload the document in the fee cover.)

I/We hereby declare that we accept the condition that if I/We withdraw or modify our bids during period of validity etc., we will be suspended for a period of two years from participating in all tenders invited by JIPMER, Puducherry with effect from the date of our withdrawal or modification of our bid.

(Signature with date)
(Name and designation)
Duly authorized to sign this tender and declaration
for and on behalf of _____ (Name of the Bidder's Firm) Seal

Appendix-A

Self-certification format for claiming purchase preference under the "Public Procurement preference to Make in India" order

As per the order issued by

(i) Department of Industrial Policy and Promotion (DIPP) vide No. P-45021/2/2017-BE-II dated 15.06.2017 as further amended by Order No.P-45021/2/2017-B.E.-ll dated 28.05.2018, Order No.P-45021/2/2017-B.E.-II dated 29.05.2019, Order No. P-45021/2/2017-PP (BE-II) dated 04.06.2020 and Order No. P-45021/2/2017-PP (BE-II) dated 16.09.2020; and

(ii) Department of Pharmaceuticals vide No. F- 31026/36/2016-MD dated 18.05.2018 and the subsequent orders thereof; The purchaser reserves the right to give preference to the local supplier.

A local supplier (definition of "local supplier" is given in clause 2 of the aforesaid order of DIPP as amended from time to time) has to submit the following along with their e-tender(s) failing which their bid will be evaluated without considering such preference mentioned in the DIPP order dated 15.06.2017 further amended on 28.05.2018, 25.09.2019 and 04.06.2020:

a. The local supplier at the time of e-tender, bidding or solicitation shall be required to provide self-certification that the item offered meets the minimum local content and shall give details of the location(s) at which the local value addition is made in the format in Annexure A.

"Certified that the following items quoted has more than 50% or 20% to 50% of value addition in India at the location(s) mentioned against each and is eligible for purchase preference as per the Govt. of India "Public Procurement preference to Make in India" order Dt.15.06.2017 as further amended by Order No.P-45021/2/2017-B.E.-ll dated 28.05.2018, Order No.P-45021/2/2017-B.E.-II dated 29.05.2019, and Order No. P-45021/2/2017-PP (BE-II) dated 04.06.2020.

Sl. No	Name of the item	Details of the location(s) at which the local value addition was made.

Authorized Signature:
Name:
Designation:

b. In cases of procurement for a value in excess of Rs. 10 Crore. the local supplier shall be required to provide a certificate from the statutory auditor or cost auditor of the company (in the case of companies) or from a practicing cost accountant or practicing chartered accountant (in respect of suppliers other than companies) giving the percentage of local content.

c. Minimum Local Content: 'Class-I local supplier' means a supplier or service provider, whose goods, services or works offered for procurement, has local content equal to or more than 50%, as defined

under the Order. “Class-II local supplier” means a supplier or service provider, whose goods, services or works offered for procurement, has local content more than 20% but less than 50%, as defined under this Order.

d. Margin of Purchase Preference: The margin of purchase preference shall be 20%.

e. Manufacture under license/technology collaboration agreements with phased indigenization are exempted from meeting the stipulated local content if the product is being manufactured in India under a license from a foreign manufacturer who holds intellectual property rights and where there is a technology collaboration agreement/transfer of technology agreement for indigenous manufacture of a product developed abroad with clear phasing of increase in local content

f. Decisions on complaints relating to implementation of this Order shall be taken by the competent authority which is empowered to look into procurement-related complaints relating the procuring entity.

g. A constituted committee with internal and external experts will examine for independent verification of self-declarations and auditor’s/accountant’s certificates on random basis and in the case of complaints.

h. A fees of Rs.10000/- in the form of demand draft favoring The Director, JIPMER, payable at Puducherry, is required to be deposited with complaints for verification of local content.

i. False declarations will be breach of the Code of Integrity under Rule 175(1)(i)(h) of the General Financial Rules for which a bidder or its successors can be debarred for up to two years as per Rule 151(iii) of the General Financial Rules along with such other actions as may be permissible under law.

j. A supplier who has been debarred by any procuring entity for violation of this Order shall not be eligible for preference under this Order for procurement by any other procuring entity for the duration of the debarment. The debarment for such other procuring entities shall take effect prospectively from the date on which it comes to the notice of other procurement entities.

Officer in Charge
Purchase Section
JIPMER Puducherry
For Director, JIPMER

SECTION – X

GFR-144 (xi) compliance certificate
(To printed on the Firm's letterhead)

Tender No:

GFR-144(xi) compliance certificate (as per order F.No. 6/18/2019-PPD, Ministry of Finance, GOI)

I have read the clauses regarding restrictions under GFR144(xi) on procurement from a bidder of a country which shares a land border with India. I certify that ……., the vendor

- ☐ is not such a country
- ☐ is from a country and has been registered with a competent authority (attached evidence of valid registration).

(Select one of the above and strike off the other)

I hereby certify that we fulfill all requirement in this regard and is eligible to be considered for the procurement on CPP portal.

Thanking you.

Authorized Signatory