Rate Contract for Consumables & Implants for Structural Heart Intervention on Consignment Basis

at

All India Institute of Medical Sciences, Jodhpur

NIT No.	**:**	**PROC-2/RC/05/2026-AIIMS.JDH**
NIT Issue Date	**:**	**14th May, 2026**
Last Date of Submission	**:**	**16th June, 2026**
Pre-Bid Meeting	**:**	**01st June, 2026 (3:00 PM)**

Tender documents may be downloaded from institute's web site www.aiimsjodhpur.edu.in (for reference only) and GeM-CPP Portal site www.eprocure.gov.in (for bidding)

All India Institute of Medical Sciences, Jodhpur
Basni Phase - II, Jodhpur - 342 005, Rajasthan
Phone: 0291-2740741,
Email: ***Proc2RateContract@aiimsjodhpur.edu.in***
Website: www.aiimsjodhpur.edu.in

All India Institute of Medical Sciences, Jodhpur, Rajasthan, is an apex healthcare institute established by an Act of Parliament of India under aegis of Ministry of Health & Family Welfare, Government of India, invites Online Bids in Two Bid System for **Rate Contract for Consumables & Implants for Structural Heart Intervention on Consignment Basis for Department of Cardiology at AIIMS Jodhpur.– PROC-2/RC/05/2026** at AIIMS Jodhpur. Bidders are requested to quote their best offers along with the complete details as required in the succeeding paragraphs.

General Instructions to Bidders

1. Bids shall be submitted online only through the GeM-CPP Portal at www.eprocure.gov.in

2. **Only Original Manufacturer (OM) or The Authorized Indian Agent (Importer) of Foreign Firm can participate in the bid.**

3. The complete bidding process is ONLINE. The Bidder should be in possession of valid Digital Signature Certificate (DSC) for online submission of bids. Prior to the bidding, the DSC needs to be registered on the above-mentioned website. For any assistance regarding e-Bidding Process, the bidder may contact to the Help Desk at 0291-2740741.

4. Bidders are advised to follow the instructions provided in the **'General Instructions to the Bidders'** for the e-submission of the bids online through the GeM - *Central Public Procurement Portal*.

5. **Bid Security (Earnest Money):** The bidder shall be required to submit the Bid Security (Earnest Money) for an amount of **₹ 2,00,000 /-** by way of ***Demand Draft only*** (no other mode is acceptable). The Bid Security (Earnest Money) shall be drawn in favour of '*All India Institute of Medical Sciences, Jodhpur'*.
   a) The Bid Security (Earnest Money) has to be deposited by all the bidders except Micro and Small Enterprises (MSEs) as defined in MSE Procurement Policy issued by Department of Micro, Small and Medium Enterprises (MSME) or are registered with the Central Purchase Organization or the concerned Ministry or Department [or Start-ups as recognized by Department for Promotion of Industry and Internal Trade (DPIIT)]. *The exemption is applicable only for the goods that are produced and quoted in this contract by the participating MSE bidder.* However, traders/ distributors/ sole agent/ Works Contract are excluded from the purview of Public Procurement Policy for MSEs Order, 2012. Bidders seeking exemption from Bid Security must submit their registration certificates/documents in the technical bid. No other relaxation shall be allowed.
   b) The Bid Security (Earnest Money) of Successful Bidders shall be returned after the receipt of Performance Security from all the Awardee of Contract.
   c) The Bid Security (Earnest Money) of the Unsuccessful Bidders shall be returned after the Award of Contract.
   d) No request for transfer of any pervious Bid Security or Performance Security or Payment of any Pending Bill held by the institute, in respect of any previous work, will be entertained.
   e) The bidder is not permitted to withdraw his offer or modify the terms and conditions thereof. In case the bidder fails to observe and comply with the conditions made herein or backs out after quoting the rates, the Bid Security will be forfeited.
   f) No Claim shall lie against AIIMS- Jodhpur in respect of interest on the amount of Bid Security (Earnest Money).

   The Original Hard Copy of only of the Bid Security (*Demand Draft)* must be delivered to the office of Deputy Director (Administration), AIIMS, Jodhpur before the last date / time of Bid Submission. The bid without Bid Security (Earnest Money) will be summarily rejected. The copy of Bid Security must be attached with the Technical Bid Document.

6. The bidder must read and examine all the terms, conditions, instructions, etc., contained in the NIT. Failure to provide and/or comply with the required information and instructions incorporated in the Tender Documents will result in the rejection of their bid.

7. **SUBMISSION OF TENDER**
   a) The bid shall be submitted online in Two Parts viz. ***(A) TECHNICAL BID*** and ***(B) FINANCIAL BID*** (BoQ in an Excel Sheet).
   The **(A) TECHNICAL BID** consists of two parts: (i) Technical Bid Document (PDF) AND (ii) List of Quoted Items (Excel file).
   b) All the pages of bid being submitted must be **signed - stamped and sequentially numbered** by the bidder, irrespective of nature of content of the documents before uploading.
   c) **The Technical Bid documents must be submitted and arranged in the same order as specified in the Form-A [Check List].**
8. **Consignment Basis** means when AIIMS Jodhpur requires a particular implant construct which is already in Rate Contract Agreement, the appointed dealer/supplier of that company supplies the required implant construct immediately on chalan basis and the payment is made to the dealer/supplier for the consignment supplied after the issuing of consumption certificate.
9. AIIMS Jodhpur will sign a Rate Contract with the successful bidders/companies of the list of items of **Rate Contract for Consumables & Implants for Structural Heart Intervention on Consignment Basis for Department of Cardiology at AIIMS Jodhpur.– PROC-2/RC/05/2026** on Consignment Basis for Department of Cardiology at AIIMS Jodhpur to be supplied. The items will be kept on **consignment basis.**
10. The bidder has to **maintain the inventory of the items** in consultation with the user department so that in any case the patient care is not compromised.

GENERAL TERMS AND CONDITIONS

Sub.: Notice Inviting bids for **Rate Contract for Consumables & Implants for Structural Heart Intervention on Consignment Basis– PROC-2/RC/05/2026 for Department of Cardiology at All India Institute of Medical Sciences- Jodhpur**.

1. **PARTIES**
   The parties to the contract are the contractor (the bidder to whom the Award of Contract have been awarded) and the AIIMS- Jodhpur through Deputy Director, All India Institute of Medical Sciences, Jodhpur for and on behalf of the Executive Director, AIIMS, Jodhpur.

2. **PRE-BID MEETING**

a) The Pre-Bid Meeting will be held at the Conference Hall, Medical Superintendent Office, OPD Building, AIIMS, Jodhpur. Bidders are advised to submit their queries or representations only via email to **Proc2RateContract@aiimsjodhpur.edu.in**, no later than **Two Days after the Pre-Bid Meeting, by 05:00 PM.**

b) The bidder MUST keep the subject of the eMail as "**Rate Contract for Consumables & Implants for Structural Heart Intervention on Consignment Basis for Department of Cardiology at AIIMS Jodhpur.– PROC-2/RC/05/2026** ".

c) Bidders are Bidders are instructed to send all their representations in a single email. If it becomes necessary to send a new representation, the bidder may send a new email, but it must include all previously submitted representations in consolidated form. Only the latest email will be considered.

d) Representations received thereafter, or sent to any other email ID of the Institute, will not be entertained.

3. The proposal for rate contract may be submitted in the prescribed format and all columns may be filled up. Incomplete proposals and tenders received after due date shall not be entertained. Any bids received by the Institute, which does not fulfill the terms and conditions shall be rejected out rightly and no communication in this regard shall be sent.
4. Bids comprising of such vague and indefinite expression such as '*Subject to prior confirmation*', '*Subject to immediate acceptance*' etc. will be treated as vague offers and rejected accordingly. Any conditional tender shall be rejected summarily.

5. The bidder shall not be permitted to withdraw his offer or modify the terms and conditions thereof. In case the tenderer fails to observe and comply with stipulation made herein or backs out after quoting the rates, **the amount of Bid Security (Earnest Money) will be forfeited**.
6. **Only Original Manufacturer (OM) or The Authorized Indian Agent (Importer) of Foreign Firm can participate in the bid (the AUTHORIZATION CERTIFICATE as per the Form-E, showing the Validity Period, must be submitted / attached in the Technical Bid). The Original Hard copies of such authorization certificates must be submitted on or before the last date of bid submission, failing which the bid shall be rejected.**

7. At any time prior to date of submission of tender, AIIMS-Jodhpur may, for any reason or decision, modify the terms & conditions of the tender document by a corrigendum displayed on the website of AIIMS-Jodhpur and the CPP portal. In order to provide reasonable time to take the amendment into account in preparing the bid, AIIMS-Jodhpur may, at its discretion, change the date and time for submission of bids.

8. **DOCUMENTS COMPRISING THE BID:** The Two Bid System i.e. ***(A) TECHNICAL BID*** and ***(B) FINANCIAL BID***

(A) TECHNICAL BID

To qualify in the Technical Bid the firm must fulfill the minimum eligibility criteria as under and the firm in this regard must submit / attach all the following documents in support of their eligibility criteria:

8.01. **Form-A** : Check List [on the Letter Head of the bidder]

8.02. **Form-B** : Details of the **(a) Bidder** and **(b) Key Person** for any communication regarding Rate Contract [on the Letter Head of the bidder].

8.03. **Form-C** : Details of the Original Manufacturers [on the Letter Head of the bidder]

8.04. **Bid Security (Earnest Money)**

8.05. **Form-D** : Tender Acceptance Form [on the Letter Head of the bidder]

8.06. **Form-E** : Authorization Certificates issued to the participating bidder by the Principal Manufacturer, Indian Subsidiary, Indian Agent, or Importer [on the Letter Head of the issuer]

- ***The Original Hard copies of such authorization certificates must be submitted on or before the last date of bid submission, failing which the bid shall be rejected.***
- ***If, the bidders are participating through Indian Importers, the bid must include the Original Manufacturer's Authorization Certificate issued to those importers*** *(no Hard Copy is required to be submitted to the institute).*

8.07. **Form-F** : No Debarment / Non-Blacklisting Certificate [on ₹ 500/- Non-Judicial Stamp Paper]

8.08. **Form-G** : No Case Pending Declaration [on ₹ 500/- Non-Judicial Stamp Paper]

8.09. **Form-H** : No Deviation Certificate [on ₹ 500/- Non-Judicial Stamp Paper]

8.10. **Form-I** : Price Justification Certificate [on ₹ 500/- Non-Judicial Stamp Paper]

8.11. **Form-J** : Land Border Declaration [on ₹ 500/- Non-Judicial Stamp Paper]

(In compliance of the terms and conditions mentioned in Department of expenditure OM No. 6/18/2019-PPD dated: 23rd July, 2020 and subsequent guidelines issued thereafter).

8.12. **Form-K** : Self Certification regarding Local Content [on ₹ 500/- Non-Judicial Stamp Paper] – if applicable.

8.13. **Form-L** : Bank Details / Mandate Form [on the Letter Head of the bidder]

8.14. **Annexure-1** 'List of Quoted Items' (TO BE SUBMITTED IN THE **GIVEN EXCEL SHEET**)

8.15. **Annexure-2** 'Calculation of Local Content' (To be submitted along with BoQ only) – if applicable.

Note:
1. This Annexure-2 [Calculation of Local Content] is to be submitted with BoQ [Financial Bid] only.
2. Your bid shall be REJECTED, if the Annexure-2 [Calculation of Local Content] is submitted in the Technical Bid Document.

8.16. Valid Registration Certificate / Certificate of Incorporation of the bidder's firm

8.17. GST Registration Certificate of the bidder's firm

8.18. PAN Card of the bidder's firm

8.19. Bidder's **Average Annual Turnover** from similar jobs should be at least **₹ 100 Crores** during the Last 03 Financial Years : 2022-23 , 2023-24 & 2024-25, from Indian business. A certificate or statement to this effect, duly verified and audited by a Chartered Accountant, must be submitted. Bidder's GST and/or PAN Number must be mentioned on this document.

8.20. Bidder's **Income Tax Return Acknowledgement** of the 03 Assessment Years: 2022-23, 2023-24 & 2024-25.

8.21. Bidder's Profit & Loss A/c and Balance Sheet of the 03 Financial Years : 2022-23 , 2023-24 & 2024-25, duly verified and audited by a Chartered Accountant.

8.22. If an Authorized Distributor is participating in this tender, they shall also be required to submit the following documents issued by the Principal Manufacturer, Indian Subsidiary, Indian Agent, or Importer, as applicable.

- **Form-F** : No Debarment / Non-Blacklisting Certificate [on ₹ 500/- Non-Judicial Stamp Paper]
- **Form-J** : Land Border Declaration [on ₹ 500/- Non-Judicial Stamp Paper]

8.23. Bidder's valid ***Drug Licenses*** issued by the Central / State Drug Controller (If applicable).

8.24. The bidder must upload required Quality Assurance Certifications (If applicable for any item) /documents in techno-commercial bid for each items along with item no. i.e., self- attested copies of CE/USFDA/DCGI/ISO/BSP/USP/WHO/GMP/BIS as mentioned in the specification of particular tender items, failing which the offer for such items will be rejected.

8.25. ***Import Licenses*** issued to the Authorized Agent / Importer by the Government of India or ***Manufacturing License*** issued by CDSCO as the case may be.

8.26. This ***NIT Document and its Corrigendum*** (issued by the institute), all pages duly signed and stamped.

8.27. Any other relevant document (left to be included in this list).

Note: The concerned firm / company whose product has been declared as of spurious or adulterated quality and any criminal cases is filled and is pending in any court shall not be eligible to participate in the bidding process. Convicted firms/company shall also not be eligible to participate in the bid. Similarly, blacklisted / banned / debarred firms / company by any central / state govt. or its organization or autonomous bodies or central drug procurement agency is not eligible to participate in the bid.

(B) FINANCIAL BID

Financial Bid as per BoQ format filled up with all the details of the goods offered to be uploaded.

Schedule of price bid in the form of BOQ_XXXX.xls:
Price bid format is provided as BoQ_XXXX.xls along with this Tender Document at www.eprocure.gov.in. Bidders are advised to download this BoQ_XXXX.xls as it is and quote their offer/rates in the permitted column and upload the same in the Financial Bid. Bidder shall not tamper/modify the downloaded price bid template in any manner. In case if the same is found to be tampered / modified in any manner, tender will be completely rejected out rightly.

9. The Item Names and Specifications are mentioned in the **Annexure-3 'Item List'**

10. Full description & specifications, Make (Company) / Brand / Name of the manufacturing firm must be clearly mentioned in the tender failing which the tender will not be considered. The tendered must also mention whether the goods are imported / indigenous. Descriptive literature / catalogues must be attached with the tender in original failing which tender may be rejected.

11. **SIGNING OF TENDER**
Individual signing the tender or other documents connected with contract must specify whether he sign as:
a) A sole proprietor of the concern or constituted attorney of such sole proprietor;
b) A partner of the firm, if it is a partnership firm in which case he must have authority to execute the contracts on behalf of the firm and to refer to arbitration disputes concerning the business of the partnership either by virtue of the partnership agreement or by a power of attorney duly executed by the partners of the firm.
c) Director or a principal officer duly authorized by the Board of Directors of the Company, if it is a company.
d) A person signing the tender form or any document forming part of the tender on behalf of another person should have an authority to bind such other person and if, on enquiry it appears that the person so signing had no authority to do so, AIIMS, Jodhpur may without prejudice, cancel the contract and hold the signatory liable for all costs, consequences and damages under the civil and criminal remedies available.

12. **EXCLUSION**
The near relatives of employees of AIIMS, Jodhpur are prohibited from participation in this tender. The near relative for this purpose is defined as:
a) Members of a Hindu Undivided Family.
b) Their spouses
c) The one related to the other in the manner as father, son(s), Son's wife (daughter-in-law), daughter(s) and daughter's husband (sons-in-law) brother (s) and brother's wife, sister(s) and sister's husband, brother(s)-in-law.

13. **CODE OF INTEGRITY :** No official of the bidder shall act in contravention of the codes which includes prohibition of:
    a) Making offer, solicitation or acceptance of bribe, reward or gift or any material benefit, either directly or indirectly, in exchange for an unfair advantage in the procurement process or to other wise influence the procurement process.
    b) Any omission, or misrepresentation that may mislead or attempt to mislead so that financial or other benefit may be obtained or an obligation avoided.
    c) Any collusion, bid rigging or anti-competitive behavior that may impair the transparency, fairness and the progress of the procurement process.
    d) Improper use of information provided by the procuring entity to the bidder with an intent to gain unfair advantage in the procurement process or for personal gain.
    e) Any financial or business transactions between the bidder and any official of the procuring entity related to tender or execution process of contract; which can affect the decision of the procuring entity directly or indirectly.
    f) Any coercion or any threat to impair or harm, directly or indirectly, any party or its property to influence the procurement process.
    g) Obstruction of any investigation or auditing of a procurement process.
    h) Making false declaration or providing false information for participation in a tender process or to secure a contract.
    i) Disclosure of conflict of interest.
    j) Disclosure by the bidder of any previous transgressions made in respect of the provisions of sub-clause with any entity in any country during the last three years or of being debarred by any other procuring entity.

14. **DOCUMENTS**
    a) All pages of the Tender should be numbered and indexed.
    b) The bidder shall provide in its tender the required as well as the relevant documents like technical data, literature, drawings etc. to establish that the goods and services offered in the tender fully confirm to the goods and services specified by the AIIMS, Jodhpur in the tender documents. For this purpose, the bidder shall also provide a clause-by-clause commentary on the technical specifications and other technical details incorporated by the AIIMS, Jodhpur in the tender documents to establish technical responsiveness of the goods and services offered in its tender duly indicating relevant page numbers in the product literature.

15. **BID CURRENCIES**
    The bidder supplying indigenous goods or imported goods shall quote only in Indian Rupees (INR). Bids, where prices are quoted in any other currencies shall be treated as non-responsive and will be rejected.

16. **BID PRICES**
    a) The Bidder shall indicate in the Price Schedule provided in BoQ all the specified components of prices shown therein including the unit prices on Free Delivery at Site basis, applicable GST, HSN Code, it proposes to supply against the requirement. The Bidders shall indicate MRP in the relevant column against each item of BoQ. The details about make & model, if applicable, may also be indicated. All the columns shown in the Price Schedule should be filled up as required.
    b) In no case the quoted rates should be more than MRP at the time of submission of quotation. If subsequently during the currency of Rate Contract there is decreased in MRP, the bidder shall inform the purchaser promptly along with revised reduced rates. In case, if bidder quotes more than MRP and/or does not inform purchaser about reduction in MRP, it will be viewed seriously and appropriate administrative action will be taken including de-barring the firm.

17. **VALIDITY OF THE BIDS**

The bids shall be valid for a period of **12 Months** after the date of bid opening prescribed in the Tender Document. In case, the Contract is not finalized within stipulated period, its validity shall be updated via corrigendum(s) on the institute's website.

18. **RIGHT OF ACCEPTANCE**

a) The AIIMS, Jodhpur reserve the right to accept the whole or any part or portion of the bid; and the bidder shall provide the same at the rates quoted. The AIIMS Jodhpur reserve the right to reject any or all tenders / quotations or all offers received in response to the tender or cancel or withdraw the tender notice without assigning any reason thereof and also does not bind itself to accept the lowest quotation or any tender and no claim in this regard shall be entertained.

b) The AIIMS- Jodhpur does not bind itself to accept the lowest bid or any bid and reserves the right of accepting the whole or any part of the bid or portion of the job offered; and the bidder shall provide the same at the rates quoted. AIIMS- Jodhpur, reserves the right to reject any or all offers received in response to tender or cancel or withdraw the tender notice without assigning any reason, whatsoever.

c) The AIIMS- Jodhpur reserves the right to place an order for purchase of any items mentioned in the Financial Bid or otherwise, to any other firm(s) in emergency/unavoidable situation.

d) AIIMS- Jodhpur reserves the right to conclude more than one rate contract for the same item.

19. **FIRM PRICE**

Prices quoted by the bidder shall remain firm and fixed during the period of the Rate Contract and not subject to variation on any account. Purchase Orders will be placed by Centers / Hospital / Departments / Store Sections against this Rate Contract till the period of Rate Contract. Statuary variation in GST will be applicable.

20. **ALTERNATIVE MODELS / BRANDS / QUALITY**

Alternative Models / Brands / Quality are not permitted. The Bidder are required to quote Models / Brands / Quality of best quality meeting tender specifications. Wherever, a bidder quotes alternative Models / Brands / Quality, there bid will not be considered for that item.

21. **PURCHASE PREFERENCE TO LOCAL SUPPLIERS**

In pursuance of Government of India Order No. P-45021/2/2017-B.E.-II dated 16th September 2020 (as amended from time to time) and F.No. Z.28018/67/2017-EPW dated 12th June 2018 purchase preference shall be given to local suppliers in all procurements undertaken in the manner specified hereunder and the procurement shall be made as per terms and conditions contained in the said order.

22. **MINIMUM LOCAL CONTENT**

The minimum local content shall be maintained as per Government of India Order No. P-45021/2/2017-B.E.-II dated 16th September 2020 (as amended from time to time) and F.No. Z.28018/67/2017-EPW dated 12/06/2018, as amended from time to time.

23. **MARGIN OF PURCHASE PREFERENCE**

The margin of purchase preference benefit will be as per the Order No. P-45021/2/2017-B.E.-II dated 16th September 2020. Bidder shall submit an undertaking within 7 days of opening of financial bid, that he would be ready to supply the product at L1 price. In case of non-receipt of the same, he would not be given purchase preference.

24. **Affidavit of Self Certification regarding Local Content** [***Form-XI***] on ₹ 500/- Non-Judicial Stamp Paper, except 100% importers.

25. **TECHNICAL EVALUATION**
    a) Detailed technical evaluation shall be carried out by Tender Finalization Committee pursuant to conditions in the tender document to determine the substantial responsiveness of each tender. For this clause, the substantially responsive bid is one that conforms to all the eligibility and terms and condition of the tender without any material deviation. The Institute's determination of bid's responsiveness is to be based on the contents of the bid itself without recourse to extrinsic evidence. The Institute shall evaluate the technical bids also to determine whether they are complete, whether required sureties have been furnished, whether the documents have been properly signed and whether the bids are in order.
    b) Financial bids of only those bidders who qualify the technical criteria will be opened provided all other requirements are fulfilled.
    c) AIIMS Jodhpur shall have right to accept or reject any or all bidders without assigning any reasons thereof.

26. **SAMPLE/DEMONSTRATION**
    a) The bidders may be asked to submit samples of the items (if required) for which they have quoted (without indicating price), with clear marking of Firm's Name in each of item, Tender Reference number and all the expenses for sample/demonstration shall be borne by the bidder.
    b) The samples are required to be submitted at the designated location, in original packing, duly labelled (printed) and sealed, bearing all relevant details such as manufacturing date, expiry date, batch number, etc. The specific submission location at AIIMS Jodhpur will be communicated separately.
    c) The sample received from bidders will be evaluated by the Technical Evaluation Committee for their quality and the decision of the Committee will be final.

27. **FINANCIAL EVALUATION**
    a) The financial bid shall be opened for only those bidders who are found to be technically eligible through the CPP Portal only.
    b) Arithmetical errors shall be rectified on the following basis. If there is a discrepancy between the unit price and total price that is, the unit price shall prevail and the total price shall be corrected by the Institute. If there is a discrepancy between words and figures, the lesser amount shall be considered as valid. If the Supplier does not accept the correction of the errors, that bid shall be rejected.

28. **RIGHT TO CALL UPON INFORMATION**
    a) The AIIMS, Jodhpur will have the right to call upon information regarding any document related to the tender/bid at any point of time.
    b) To assist in the analysis, evaluation and computation of the bids, the Tender Finalization Committee of AIIMS, Jodhpur, may ask bidders individually for clarification of their bids. The request for clarification and the response shall be in writing but no change in the price or substance of the bid offered shall be permitted.

29. The Tender Finalization Committee of AIIMS, Jodhpur shall go into all aspects including cost factors of Consumables and then decide for awarding of the tender, by quoting lower rates in respect of items, a firm does not become entitled to awarding the contract in its favor of those item(s). In order to get selection / consideration in the panel of two or three vendors for awarding of contract (in case the contract is to be awarded to more than one vendor), the criteria of selection for awarding contract will be calculating / comparing the rate of items consumed by the AIIMS-Jodhpur throughout the year and as per the requirement in view of quality, as deemed fit by the Tender Finalization Committee.

30. **PERFORMANCE SECURITY**
    a) The successful bidder shall be required to submit a Performance Security of a minimum amount of ₹ 2,00,000/-. If the bidder is awarded more than one item, the Performance Security shall be calculated at ₹ 10,000/- per awarded item, subject to a maximum of ₹ 5,00,000/-.
    b) The Performance Security of the bidders will be refunded / released after the completion of all contractual obligations.
    c) The Performance Security shall be forfeited by the Institute in the event of any breach or negligence or non-observance of any condition of contract or for unsatisfactory performance or non–observance of any condition of the contract.
    d) No interest on Performance Security shall be paid by the Institute.

31. **AWARD OF CONTRACT**
    a) The Institute shall consider placement of orders to those bidders whose offers have been found Technically and Financially Acceptable. The Institute reserves the right to counter offer price(s) against price(s) quoted by any bidder. The L-1 Bidder will be decided on individual item basis.
    b) AIIMS Jodhpur has the option to renegotiate the price with the Awardee of Contract.
    c) The Award of Contract shall be issued to the L-1 bidders for ***Two Years***, valid from the date of issue of Award of Contract (**extendable on mutual agreement**).
    d) AIIMS-Jodhpur shall reserve the right to terminate the contract at any time without assigning any reason thereof.

32. **AUTHORIZED DISTRIBUTOR**
    a) The bidder after receipt of *Award of Contract*, may furnish the details of their Authorized Distributor, so that the copies of orders can be endorsed to them for expeditious supply. In such cases where Indian Agent has been nominated by the manufacturer, the bills raised by them against our purchase order will be accepted.
    b) Any addition and deletion of Authorized Distributorship (in extreme cases) shall be intimated to AIIMS-Jodhpur, immediately on authorization of a new party.

33. **SUBLETTING OF WORK**
    The Authorized Distributor shall not assign or sublet the work/job or any part of it to any other person or party.

34. The Awardee of Contract shall provide the name and mobile number of a Key Person, who can be contacted at any time, even beyond the office hours on holidays. The person should be capable of taking orders and making arrangement for supply of the desired items even on short notice to AIIMS, Jodhpur.

35. **PLACEMENT OF PURCHASE ORDER**
    a) The Purchase Orders shall be issued for requirement on actual need basis.
    b) All India Institute of Medical Sciences (AIIMS), Jodhpur shall be the sole authority to cancel or amend the Purchase Order, as per requirement, and also to place Purchase Orders beyond Office Hours / Holidays / Place of Supply, for which no additional payment shall be made.

36. **PACKING**
    a) Supplies to be made in a Proper Boxes with proper packaging.
    b) Packing should be able to prevent damage or deterioration during transit.
    c) All containers, i.e., bottles, tins, cartons, tubes etc. are required to be secured with pilfer-proof seals to ensure genuineness of the products packed and the correctness of the contents.
    d) Stamping "**AIIMS JODHPUR SUPPLY ONLY**" is mandatory on each unit.

37. **DELIVERY**
    a) Delivery of Goods shall be made by the supplier within 30 days of placing of Purchase Order on challan basis, at the location mentioned in the Purchase Order issued.
    b) We being a Healthcare Institute, time is the essence of every Purchase Order, hence, the supplier firm should have availability of a responsible person on call 24x7 on all days, to accept and deliver the items at a very short notice i.e. within 24 hours also. The delivery period will be mentioned in the Purchase Order.
    c) The supply must be made in full against the Purchase Orders and shortage will be procured from other supplier on the risk and cost of the Original Supplier.
    d) If any item is found to be defective, or not of the desired quality, the same shall be replaced immediately, for which no extra payment shall be made by AIIMS, Jodhpur.

38. **LIQUIDATED DAMAGES**
    a) Supply of material will have to be completed within the delivery period or period mentioned in the purchased order. The liquidated damages charges @ 0.5% per week shall be imposed if supply made after expiry of delivery period subject to maximum 10% of the total value of relevant goods. Quantum of liquidated damages assessed and levied by the purchaser shall be final and not challengeable by the supplier.
    b) Non supply of items shall attract a penalty of 10% of the value of non-supplied goods/material.

39. **INSPECTION**
    a) AIIMS, Jodhpur shall have the right to inspect and/or to test the goods to confirm their conformity to the NIT Specifications at no extra cost to the AIIMS, Jodhpur.
    b) AIIMS, Jodhpur shall have the right to inspect, test and, where necessary, reject the Goods after the goods arrival at the final destination shall in no way be limited or waived by reason of the Goods having previously been inspected, tested and passed by AIIMS, Jodhpur prior to the goods shipment.
    c) The Executive Director, AIIMS Jodhpur shall be the final authority to reject full or any part of the supply which is not confirming to the specification and other terms and conditions.
    d) No payment shall be made for rejected/ unconsumed Stores. Rejected items must be removed by the Bidders within Two Days of the date of rejection at their own cost and replaced immediately. In case these are not removed, these will be auctioned at the risk and responsibility of the suppliers without any further notice.
    e) In case the quality of goods supplied are not in conformity with the standard given in tender and as per the samples supplied or the supplies are found defective at any stage these goods shall immediately will be taken back by the supplier and will be replaced with the tender quality goods, without any delay. The Purchase Committee reserves all right to reject the goods if the same are not found in accordance with the required description / specifications and liquidates damages shall be charged.
    f) Supply should be made from the latest batch of production with at **least 75% shelf life & original packing**.

40. **REPLACEMENT OF EXPIRED ITEM**
    a) In the event that the supplied item remains unutilized and the balance shelf life is less than 10%, the supplier shall be liable to replace the same with fresh stock of equal quantity, without any additional cost, as and when required. No communication in this regard will be entertained regarding timely information / prior to expiry of items. If the supplier fails to exchange the expired goods / items within the timeframe, it will be termed as breach of contract conditions, AIIMS Jodhpur reserves the right to initiate any suitable action against the supplier in this regard.
    b) The supplier shall arrange to effect free replacement of any quantity which may deteriorate in potency, strength approaching expiry or expired etc. before the date of expiry marked on the labels.

41. **THE PAYMENT CLAUSE**
    a) Payment of the bill will be made to the dealer/supplier for the consignment supplied after the issuing of consumption certificate of the goods utilized by the concerned department.
    b) The GST Invoice in Triplicate must be sent to the Delivery Location along with the supplied goods. The bill should have full particulars of the items(s) viz. Item's complete name, Tender Reference Number, HSN Code, Basic Rate, GST Percentage, GST Amount, Total Amount (in numbers & words).
    c) Freight, insurance charges, if any will be borne by the supplier, similarly shortage, pilferage in transit will be sole responsibility of the supplier and the same will be intimated to the supplier on receipt of goods by the purchaser for resupply. The defective supply will have to be replaced by the supplier within Two Days without additional freight / transport charge etc.
    d) GST and other Government Levies shall be paid extra, as applicable.
    e) No payment shall be made in advance nor shall the loan from any bank or financial institutions be recommended on the basis of the order of award of work.
    f) Delivery of goods will be taken at the risk and cost of the supplier and on F.O.R. basis to the Institute from railway / road transport.
    g) No revision in rate (on higher side) will be accepted during contract period.
    h) No payment will be made for goods rejected.

42. **RISK & COST PURCHASE**
    If supplier fails to supply material within the stipulated delivery date or material supplied other than specification specified in our NIT, AIIMS Jodhpur reserves the right to terminate contract for that item(s), forfeiture of security deposit and to procure same or equivalent material from alternative sources at the vendor's risk, responsibility and cost. Any extra cost incurred in the procurement of the material from alternative source will be recovered from the Security Deposit / Bank Guarantee and Pending Bills of existing firm and if the value of the materials under risk purchase exceeds, the amount of Security Deposit and / or Bank Guarantee and Pending Bills, the same may be recovered if necessary.

43. **FORCE MAJEURE**
    If, at any time during the subsistence of this contract, the performance in whole or in part by either party of any obligation under this contract is prevented or delayed by reasons of any war or hostility, act of public enemy, civil commotion, sabotage, fire, floods, explosion, epidemics, quarantine restriction, strikers lockout or act of God provided notice of happening of any such eventuality is given by party to other within 21 days from the date of occurrence thereof, neither party shall by reason of such event be entitled to terminate this contract nor shall either party have any claim for damages against other in respect of such non-performance or delay in performance, and deliveries have been so resumed or not shall be final and conclusive.

44. **INSOLVENCY / BANKRUPTCY**
    In the event of the firm being adjudged insolvent or having a receiver appointed for it by a court or any other order under the Insolvency Act made against them or in the case of a company the passing any resolution or making of any order for winding up, whether voluntary or otherwise, or in the event of the firm failing to comply with any of the conditions herein specified. AIIMS, Jodhpur shall have the power to terminate the contract without any prior notice and any other action as per the government guidelines will be taken against the bidder.

45. **ARBITRATION**
If any conflict or difference arises concerning this agreement, its interpretation on payment to make there-under, the same shall be settled out by mutual consultation and negotiation. If attempts for conciliation do not yield any result within a period of 30 days, either of the parties may make a request to the other party for submission of the dispute for decision by an Arbitral Tribunal containing Sole Arbitrator to be appointed by the Executive Director, AIIMS Jodhpur. Such requests shall be accompanied with a panel of names of three persons to act as the sole arbitrator. In case of such arbitrator refusing, unwilling or becoming incapable to act or his mandate having been terminated under law, another arbitrator shall be appointed in the same manner from among the panel of three persons to be submitted by the claimant. The provision of Arbitration and Conciliation Act, 1996 and the rule framed there under and in force shall be applicable to such proceedings.

46. **LEGAL JURISDICTION**
The agreement shall be deemed to have been concluded in Jodhpur, Rajasthan and all obligations hereunder shall be deemed to be located at Jodhpur, Rajasthan and Court within Jodhpur, Rajasthan will have Jurisdiction to the exclusion of other courts.

47. The tendering Firm/Agency/Company shall be bound by the details furnished by him/her to the All India Institute of Medical Sciences (AIIMS), Jodhpur while submitting the tender or at subsequent stage. Upon selection of the tendering Firm/Agency/Company, if at any stage, the documents furnished by him/her is found to be false or the quality of the articles found of poor quality/different specifications, it would be deemed to be a breach of terms of contract and the contract shall be cancelled at the discretion of competent authority and performance security shall be stand forfeited.

Medical superintendent

---------- ON THE LETTER HEAD OF THE BIDDER ----------

Form-A

[CHECK LIST]

To

The Executive Director,
All India Institute of Medical Sciences, Jodhpur
MIA Phase-II, Basani, Jodhpur

The bidder is advised to fill prescribed proforma & enclosed relevant document as per requirement & sequence of given proforma. The bidder should mention Page Number with all required details of relevant documents in the prescribed Proforma. If the bidder does not fill the prescribed given Proforma their offer shall be ***summarily rejected & no correspondence will be entertained***.

This **Check List** also needs to be submitted in the given Excel Sheet with the Technical Bid document.

Sr.	Details of the documents to be submitted	Page	Remarks
1	**Form-A** : Check List [on the Letter Head of the bidder]		
2	**Form-B** : Details of the bidder [on the Letter Head of the bidder]		
3	**Form-C** : Details of the Original Manufacturers [on the Letter Head of the bidder]		
3.1	Copy of Bid Security (Demand Draft only)		
4	**Form-D** : Tender Acceptance Form [on the Letter Head of the bidder]		
5.01	**Form-E** : Authorization Certificate [write here the name of the Authorization Certificate issuer-1] (for importer). **Only Original Manufacturer (OM) or The Authorized Indian Agent (Importer) of Foreign Firm can participate in the bid.**		
5.02	**Form-E** : Authorization Certificate [write here the name of the Authorization Certificate issuer-2]		
5.03	**Form-E** : Authorization Certificate [write here the name of the Authorization Certificate issuer-3]		
5.xx	and so on… [bidder can add more rows as per number of Authorization Certificates issued to them]		
6	**Form-F** : No Debarment / Non-Blacklisting Certificate [on ₹ 500/- Non-Judicial Stamp Paper]		
7	**Form-G** : No Case Pending Declaration [on ₹ 500/- Non-Judicial Stamp Paper]		
8	**Form-H** : No Deviation Certificate [on ₹ 500/- Non-Judicial Stamp Paper]		
9	**Form-I** : Rate Reasonableness Certificate [on ₹ 500/- Non-Judicial Stamp Paper]		
10	**Form-J** : Land Border Declaration [on ₹ 500/- Non-Judicial Stamp Paper]		
11	**Form-K** : Self-Certification regarding Local Content [on ₹ 500/- Non-Judicial Stamp Paper]		
12	**Form-L** : Bank Details / Mandate Form [on the Letter Head of the bidder]		
12.01	**Form-M :** Integrity Pact [on ₹ 500/- Non-Judicial Stamp Paper]		
12.02	**Annexure-1** 'List of Quoted Items' (TO BE SUBMITTED IN THE **GIVEN EXCEL SHEET**)		
13	Valid Registration Certificate / Certificate of Incorporation of the bidder's firm		

14	GST Registration Certificate of the bidder's firm		
15	PAN Card of the bidder's firm		
16	Bidder's Average Annual Turnover Certificate / Statement duly verified and audited by Charted Accountant for the F.Y. 2022-23 , 2023-24 & 2024-25. Bidder's GST and/or PAN Number must be mentioned on this document.		
17	Bidder's Income Tax Return Acknowledgement for Last Three Assessment Years: 2022-23, 2023-24 & 2024-25		
18.1	Bidder's Profit & Loss A/c and Balance Sheet (F.Y. 2022-23)		
18.2	Bidder's Profit & Loss A/c and Balance Sheet (F.Y. 2023-24)		
18.3	Bidder's Profit & Loss A/c and Balance Sheet (F.Y. 2024-25)		
20	Bidder's valid ***Drug Licenses*** issued by the Central / State Drug Controller (If applicable).		
21	Valid ***Licenses/***CE/USFDA/DCGI/ISO/BSP/USP/WHO/GMP/BIS (wherever applicable).		
22	***Import Licenses*** issued to the Authorized Agent / Importer by the Government of India or ***Manufacturing License*** issued by CDSCO as the case may be.		
23	This ***NIT Document and its Corrigendum*** (issued by the institute), *all pages duly signed and stamped*.		
24	Any other relevant document (*left to be included in this list*).		
TOTAL NUMBER OF PAGES IN TECHNICAL BID DOCUMENT			

- Page Number ***Must be marked*** on each and every page of Tender Documents, Corrigendum(s) and all copies of the documents attached.
- The bidder ***Must Mention*** Page number in the above Table, wherever the copy(s) of the document(s) are arrange / kept.
- In case of non-fulfilment of any of the above information / document(s), the bid will be summarily rejected, without giving any notice.

Undertaking

1. That I/we have carefully studied all the terms & conditions of NIT and shall abide by it.
2. That I/We undertake that the information given in this tender are true and correct in all respect and I/We hold the responsibility for the same.
3. That I/We undertake that sample of items will be kept ready for inspections by the AIIMS, Jodhpur. I/We shall be responsible for the cancellation of tender if samples are not up to mark.

Name of the Bidder:	_____
Signature	_____
Stamp	_____
Designation:	_____
Contact:	_____
eMail ID:	_____
Office Address:	_____

Place: _____Date: _____

---------- ON THE LETTER HEAD OF THE BIDDER ----------

Form-B

To

The Executive Director,
All India Institute of Medical Sciences, Jodhpur
MIA Phase-II, Basani, Jodhpur

(a) DETAILS OF THE BIDDER	
Name	
Designation in the firm	
Contact	
e-mail	
Bidder Type (Original Manufacturer/ Importer / any other	
Firm / Business Type (Sole Proprietorship, HUF, OPC, Partnership, Company or any other)	
Office Address	
Country of Origin/Registration of the **Bidder**	

(b) DETAILS OF THE KEY PERSON FOR HANDLING QUERRIES / CLARIFICATION REGARDING THE BID DOCUMENT **(preferably from Jodhpur or nearby area)**	
Name	
Designation	
Contact	
email	
Office Address	
Note: The individual designated for handling queries or providing clarifications may be contacted by the institute at any stage until the completion of the Rate Contract process. As this person will serve as the official representative of the bidder, it is advised that bidders appoint a responsible and competent individual for this role.	

---------- ON THE LETTER HEAD OF THE BIDDER ----------

Form-C

To

The Executive Director,
All India Institute of Medical Sciences, Jodhpur
MIA Phase-II, Basani, Jodhpur

Sr.	Name of the Original Manufacturer	eMail / Contact details of the Original Manufacturer	Country of Origin / Registration of the Original Manufacturer	Number of Items Quoted
01				
02				
03				
04				
05				

---------- ON THE LETTER HEAD OF THE BIDDER ----------

Form-D

[TENDER ACCEPTANCE FORM]

To

The Executive Director,
All India Institute of Medical Sciences, Jodhpur
MIA Phase-II, Basani, Jodhpur

NIT No. _____ _____ _____ _____ ____ dated: _____ for Rate Contract _____ _____ _____ _____ _____ at AIIMS Jodhpur.

1. We, the undersigned have examined the above-mentioned Tender Enquiry Document, including amendment / corrigendum (if any), the receipt of which is hereby confirmed. We now offer to supply and deliver in conformity with your above referred document for the sum as shown in the Price Schedules (BoQ) uploaded herewith and made part of this bid. If our bid is accepted, we undertake to supply the items for which Rate Contract has been concluded, in accordance with the delivery schedule as specified in the Schedule of Requirements.

2. We further confirm that, if our bid is accepted, we shall provide you with a Performance Security of required amount in an acceptable form as mentioned in your NIT.

3. We agree to keep our bid valid for acceptance as required in your NIT Document, subsequently for the extended period, if any, agreed to by us. We also accordingly confirm to abide by this bid up to the aforesaid period. We further confirm that, until a formal Rate Contract is executed, this bid read with your written acceptance thereof within the aforesaid period shall constitute a binding contract between us.

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

5. We confirm that we do not stand deregistered/banned/blacklisted by Central / State Govt. / PSU / Autonomous Bodies etc.

6. We confirm that we fully agree to the terms and conditions specified in the Tender Enquiry Document, including amendment / corrigendum if any.

7. We hereby certify that if at any time, information furnished by us is proved to be false or incorrect, we are liable for any action as deemed fit by AIIMS Jodhpur in addition to forfeiture of the Bid Security / Performance Security.

Name: _____

(Signatures of the Bidder with Name, Designation & Company's Seal)

Address: _____

_____

Place: _____

Date: _____

---------- **ON THE LETTER HEAD OF THE ISSUER** ----------

Form-E

[AUTHORIZATION CERTIFICATE]

Dated:

To

The Executive Director,
All India Institute of Medical Sciences, Jodhpur
MIA Phase-II, Basani, Jodhpur

NIT No. _____ _____ _____ _____ ____ dated: _____ for Rate Contract _____ _____ _____ _____ _____ at AIIMS Jodhpur.

Respected Sir,

1. We, _____ _____ _____ _____ , who are proven and reputable Manufacturers / Importer of _____ _____ _____. (name and description of the Items / Category offered in the Quotation) having factories at _____ _____ _____ _____, hereby authorize M/s._____ _____ _____(name and address of the agent) to submit a Quotation, process the same further, against your requirement as contained in the above referred Tender Form for the above items manufactured by us.

2. We further confirm that no supplier or firm or individual other than Messrs. _____ _____ _____ _____ (name and address of the above agent) is authorized to submit a tender, process the same further against your requirement as contained in the above referred Quotation Form for the above items manufactured by us.

3. We also hereby confirm that we would be responsible for the satisfactory execution of supply placed on the authorized agent. We also confirm that the price quoted by our agent shall not exceed than that which we would have quoted directly.

4. The validity of this authorization certificate is valid till……..

[*Signature with date and designation*]

For and on behalf of Messrs._____

[Name, address & contact detail of the manufacturer]

Note:

1. No other format of Authorization Certificate is acceptable.
2. This Certificate must be signed by a person competent and having the power of attorney to legally bind the issuer of the Authorization Certificate.

---------- on ₹ 500/- Non-Judicial Stamp Paper ----------

Form-F

[NO DEBARMENT / NON-BLACKLISTING CERTIFICATE]

To

The Executive Director,
All India Institute of Medical Sciences, Jodhpur
MIA Phase-II, Basani, Jodhpur

NIT No. _____ _____ _____ _____ ____ dated: _____ for Rate Contract _____ _____ _____ _____ _____ at AIIMS Jodhpur.

(a) I/We hereby certify that the ……..[Name of the company / firm]…….. has not been ever debarred /blacklisted by any Central Government / State Government / Public Sector Undertaking / Institute on any account.

(b) I/We also certify that the information given in bid is true and correct in all aspects and in any case at a later date it is found that any details provided are false and incorrect, the contract may be summarily terminated at any stage, and AIIMS Jodhpur may imposed any action as per NIT rules.

OR

(a) I/We hereby certify that the [Name of the company / firm] has been debarred / blacklisted by (Name of Central / State Government / Public Sector Undertaking / Institute) vide order No. ____ ____ _____ ____ _____ dated _____ ____. However, I/We (Name of the company / firm) state that the said debarment / blacklisting has been revoked vide order No. __ ____ __ ___ ____ ____ _ dated ____ _____ ___ ____. I certify that as on date, I/We (Name of the company / firm) is not debarred / blacklisted by any Central Government / State Government / Public Sector Undertaking / Institute on any account.

(b) I/We also certify that the information given in bid is true and correct in all aspects and in any case at a later date it is found that any details provided are false and incorrect, the contract may be summarily terminated at any stage, and AIIMS Jodhpur may imposed any action as per NIT rules.

Note:

1. Strike out whichever is not applicable.
2. Do NOT delete any of the given paragraph from this form.

Name: _____

(Signatures of the Bidder with Name, Designation & Company's Seal)

Address: _____

_____

Place: _____

Date: _____

---------- on ₹ 500/- Non-Judicial Stamp Paper ----------

Form-G

[NO CASE PENDING DECLARATION]

To

The Executive Director,
All India Institute of Medical Sciences, Jodhpur
MIA Phase-II, Basani, Jodhpur

NIT No. _____ _____ _____ _____ ____ dated: _____ for Rate Contract _____ _____ _____ _____ _____ at AIIMS Jodhpur.

I, …… …… ……, proprietor / partner / owner of M/s …… …… …… ……, (*Name of firm & Address*), resident of …… …… …… …… ……. …… …… (*Complete Address*), do hereby solemnly affirm and declare that there is no legal or court case pending or being contemplated against my firm in any form.

Name: _____

(Signatures of the Bidder with Name, Designation & Company's Seal)

Address: _____

_____

Place: _____

Date: _____

---------- on ₹ 500/- Non-Judicial Stamp Paper ----------

Form-H

[NO DEVIATION CERTIFICATE]

To

The Executive Director,
All India Institute of Medical Sciences, Jodhpur
MIA Phase-II, Basani, Jodhpur

NIT No. _____ _____ _____ _____ ____ dated: _____ for Rate Contract _____ _____ _____ _____ at AIIMS Jodhpur.

1. I/We, M/s_____ hereby certify that notwithstanding any contrary indication / conditions elsewhere in our offer documents, I/We have neither set any terms and conditions nor there is any deviation taken from the conditions of AIIMS Jodhpur's tender specification, either Technical or Financial.

2. I/We agree to all the terms and conditions mentioned in AIIMS Jodhpur's tender specification with associated amendments & clarification. If any deviation is found in my / our tender documents, AIIMS Jodhpur may take any suitable decision / action against my / our firm.

Name: _____

(Signatures of the Bidder with Name, Designation & Company's Seal)

Address: _____

_____

Place: _____

Date: _____

---------- on ₹ 500/- Non-Judicial Stamp Paper ----------

Form-I

RATE REASONABLENESS CERTIFICATE

To

The Executive Director,
All India Institute of Medical Sciences,
Jodhpur MIA Phase-II, Basani, Jodhpur

NIT No. _____ _____ _____ _____ _____ dated: _____ _____ for Rate Contract _____ _____ _____ _____ _____ _____ _____ _____ _____ _____ at AIIMS Jodhpur.

I/We, M/s. _____ _____ _____ _____ _____ _____ hereby certify that the rates quoted by us are our best and most competitive, and are reasonable, taking into account the specifications, quality, technology, and prevailing market conditions.

Name: _____

(Signatures of the Bidder with Name, Designation & Company's Seal)

Address: _____ _____ _____

Date: _____

---------- on ₹ 500/- Non-Judicial Stamp Paper ----------

Form-J

[Land Border Declaration]

To

The Executive Director,
All India Institute of Medical Sciences, Jodhpur
MIA Phase-II, Basani, Jodhpur

NIT No. _____ _____ _____ _____ ____ dated: _____ for Rate Contract _____ _____ _____ _____ at AIIMS Jodhpur.

1. *"We have read the clause regarding restrictions on procurement from a bidder of a country which shares a land border with India; and solemnly certify that we are not from such a county or, if from such a country, we are registered with the Competent Authority (copy enclosed). We hereby certify that we fulfil' all requirement in this regard and are eligible to be considered."*

AND

2. *"We have read the clause regarding restrictions on procurement from a bidder of a county which shares a land border with India and on sub-contracting to contractors from such a country; and solemnly certify that we are not from such a county or, if from such a country, we are registered with the Competent Authority (copy enclosed) and we shall not subcontract any work to a contractor from such countries unless such contractor is registered with the Competent Authority. We hereby certify that we fulfil all requirement in this regard and are eligible to be considered."*

3. It is to declare that if, our bid/offer is accepted by the purchaser, as per undertaking given by us and subsequently the certificate is to be found as false, this would be ground for immediate termination of our bid/offer and further legal action in accordance with the law may be initiated on us by the procuring entity i.e. AIIMS, Jodhpur.

Name: _____

(Signatures of the Bidder with Name, Designation & Company's Seal)

Address: _____

_____

Place: _____

Date: _____

---------- on ₹ 500/- Non-Judicial Stamp Paper ----------

Form-K

[Self-Certification regarding Local Content]

Dated:

To

The Executive Director,
All India Institute of Medical Sciences, Jodhpur
MIA Phase-II, Basani, Jodhpur

NIT No. _____ _____ _____ _____ ____ dated: _____ for Rate Contract _____ _____ _____ _____ _____ at AIIMS Jodhpur.

1. I_____ _____ S/o.D/o,W/o_____ _____ _____, Resident of _____ _____ _____ _____ _____ do hereby solemnly affirm and declare as under.

2. That I will agree to abide by the terms and conditions of the policy of Government of India issued vide order no. P-45021/2/2017-B.E.-II dated 15/06/2017.

3. That the information furnished hereinafter is correct to best of my knowledge and belief and I undertake to produce relevant records before the procuring entity or any authority so nominated by the Government of India for the purpose of assessing the local content.

4. That the local content for all inputs which constitute the said drugs has been verified by me and I am responsible for the correctness of the claims made therein.

5. That in the event of the domestic value addition of the product mentioned herein is found to be incorrect and not meeting the prescribed valve-addition norms, based on Government of India for the purpose of assessing the local content, action will be taken against me as per Order No. P-45021/2/2017-B.E.-II dated 15.06.2017.

6. I agree to maintain the following information in the Company's record for a period of 8 years and shall make this available for verification to any statutory authorities:

i) Name and details of the Domestic Manufacturer (Registered Officer, Manufacturing unit location, nature of legal entity)
ii) Date on which this certificate is issued.
iii) Medicine for which the certificate is product.
iv) Procuring entity to whom the certificate is furnished.
v) Percentage of local content claimed.
vi) Name and contact details of the unit of the manufacturer.
vii) Sale Price of the product.
viii) Ex-Factory Price of the product.
ix) Freight, insurance and handling.
x) Total Bill of Material.
xi) List and total cost valve of inputs used for manufacture of the medicine certificates from suppliers, if the input is not in-house to be attached.
xii) List and cost of inputs which and imported, directly or indirectly.

For and on behalf of **(Name of firm/ entity)**

Authorized signatory (To be duly authorized by the Board of Director)

Note:
This is an optional form. If the bidder submits this form, he/she will be required to furnish more details by submitting a separate Annexure-2 [Calculation of Local Content], in an Excel Sheet as given in the BoQ - Financial Bid.

---------- **ON THE LETTER HEAD OF THE BIDDER** ----------

Form-L

[Bank Details / Mandate Form]

To

The Executive Director,
All India Institute of Medical Sciences, Jodhpur
MIA Phase-II, Basani, Jodhpur

Your Firm's Name:	
Bank Name:	
Account Number:	
IFS Code:	
Bank's Branch Address:	

---------- *on ₹ 500/- Non-Judicial Stamp Paper* ----------

Form-M

Integrity Pact

Between

All India Institute of Medical Sciences, Jodhpur hereinafter, referred to as "AIIMS Jodhpur", and
..... hereinafter referred to as "The Bidder(s)/Contractor(s)"

Preamble

AIIMS Jodhpur values full compliance with all relevant laws of the land, rules, regulations, economic use of resources and of fairness/ transparencies in its relations with its Bidder(s) and / or Contractor(s). In order to achieve this goal, AIIMS Jodhpur will appoint Independent External Monitor (IEMs) who will monitor the tender process and the execution of the contract for compliance with the principles mentioned above.

Section - 1 Commitments of AIIMS Jodhpur

1) AIIMS Jodhpur commits itself to take all measures necessary to prevent corruption and to observe the following principles:-

a) No employee of AIIMS Jodhpur, personally or through family members, will in connection with the tender for, or the execution of a contract demand, take a promise for or accept, for him/herself or third person, any material or immaterial benefit which he/she is not legally entitled to.

b) AIIMS Jodhpur will, during the tender process treat to all Bidder(s) with equity and reason. The AIIMS Jodhpur will in particular, before and during the tender process, provide to all Bidder(s) the same information and will not provide to any Bidder(s) confidential/additional information through which the Bidder(s) could obtain an advantage in relation to the tender process or the contract execution.

c) The AIIMS Jodhpur will exclude from the process all known prejudiced persons.

2) If AIIMS Jodhpur obtains information on the conduct of any of its employees which is a criminal offence under the relevant Anti-Corruption Laws of India, or if there be a substantive suspicion in this regard, AIIMS Jodhpur will inform its Chief Vigilance Officer and in addition can initiate disciplinary actions.

Section - 2 Commitments of the Bidder(s)/Contractor(s)

1) The Bidder(s)/Contractor(s) commits himself to take all measures necessary to prevent corruption. The Bidder(s)/Contractor(s) commits himself to observe the following principles during his participation in the tender process and during the contract execution.

a) The Bidder(s)/Contractor(s) will not, directly or through any other person or firm, offer, promise or give to any of AIIMS Jodhpur's employees involved in the tender process or the execution of the contract or any third person any material or other benefit which he/she is not legally entitled to, in order to obtain in exchange any advantage of any kind whatsoever during the tender process or during the execution of the contract. The Bidder(s)/Contractor(s) will not enter with other Bidder(s) into any illegal agreement or understanding, whether formal or informal. This applies in particular to prices, specifications, certifications, subsidiary contracts, submission or non-submission of bids or any other actions to restrict competitiveness or to introduce cartelization in the bidding process.

b) The Bidder(s)/Contractor(s) will not commit any criminal offence under the relevant Anti- Corruption

Laws of India; further the Bidder(s) will not use improperly, for purposes of competition or personal gain, or pass on to others, any information provided by AIIMS Jodhpur as part of the business relationship, regarding plans technical proposals and business details, including information contained or transmitted electronically.

c) The Bidder(s)/Contractor(s) of foreign origin shall disclose the name & address of the Agents/representatives in India, if any. Similarly the Bidder(s)/Contractor(s) of Indian Nationality shall furnish the name and address of foreign principals, if any. Further details as mentioned in the "Guidelines on Indian Agents of Foreign Suppliers" shall be disclosed by the Bidder(s)/Contractor(s). Further, as mentioned in the Guidelines all the payment made to the Indian agent/representative have to

be in Indian Rupees only.

d) The Bidder(s)/Contractor(s) will, when presenting his bid, disclose any and all payments he has made, is committed to or intends to make to agents, brokers or any other intermediaries in connection with the award of the contract.

e) The Bidder(s)/Contractor(s) who have signed the Integrity Pact shall not approach the courts while representing the matter to IEMs and shall wait for their decision in the matter.

2. The Bidder(s)/Contractor(s) will not instigate third persons to commit offences outlined above or be an accessory to such offences.

Section - 3 Disqualification from tender process and exclusion from future contracts

If the Bidder(s)/Contractor(s), before award or during execution has committed a transgression through a violation of Section 2, above or in any other form such as to put their reliability or credibility in question, AIIMS Jodhpur is entitled to disqualify the Bidder(s)/Contractor(s) from the tender processor take action as per rule & regulations.

Section - 4 Compensation for Damages

1. If AIIMS Jodhpur has disqualified the Bidder(s) from the tender process prior to the award according to Section 3 above, The AIIMS Jodhpur is entitled to demand and recover the damage equivalent to Earnest Money Deposit /Bid security.

2. If AIIMS Jodhpur has terminated the contract according to Section 3, or if AIIMS Jodhpur is entitled to terminate the contract according to Section 3, AIIMS Jodhpur shall be entitled to demand and recover from the Bidder(s) liquidated damages of the Contract value or the amount equivalent to performance bank Guarantee.

Section - 5 Previous Transgression

1. The Bidder declares that no previous transgressions occurred in the last 3 years with any other company in any country conforming to the anti- corruption approach or with any Public Sector Enterprise in India that could justify his exclusion from the tender process.

2. If the Bidder makes incorrect statement on this subject, he can be disqualified from the tender process or action can be taken the contract, if already awarded, can be terminated.

Section - 6 Equal treatment of all Bidder (s)/Contractor (s)

In case of Sub-contracting, the AIIMS Jodhpur Contractor shall take the responsibility of the adoption of Integrity Pact by the Sub-contractor.

1. The AIIMS Jodhpur will enter into agreements with identical conditions as this one with all Bidders and Contractors.
2. The AIIMS Jodhpur will disqualify from the tender process all bidders who do not sign this Pact or violate its provisions.

Section - 7 Criminal Charges against violating Bidder (s)/Contractor (s)/ Subcontractors (s)

If the AIIMS Jodhpur obtains knowledge of conduct of a Bidder, Contractor or subcontractor, or of an employee or a representative or an associate of a Bidder, Contractor or Subcontractor which constitutes corruption, or if the AIIMS Jodhpur has substantive suspicion in this regard, the AIIMS Jodhpur will inform the same to the Chief Vigilance Officer.

Section - 8 Independent External Monitor

1. AIIMS Jodhpur has appointed competent and credible Independent External Monitors (hereinafter referred to as monitors) for this Pact as per the recommendations of Central Vigilance Commission. The particulars of Independent External Monitors are as hereunder:-

Sr. No.	Name of IEM	Email ID	Address
1	Dr. M. Malakondaiah, IPS (Retd.)	mannam1958@gmail.com	156, Prashasan Nagar, Jubilee Hills, Road No. 72, Hyderabad-500110

2. The Monitor is not subject to instructions by the representatives of the parties and performs his/her functions neutrally and independently. The Monitor would have access to all contract documents, whenever required. It will be obligatory for him / her to treat the information and documents of the Bidders / Contractors as confidential. He/ she reports to the Executive Director AIIMS Jodhpur.
3. The Bidder (s) Contractor (s) accepts that the Monitor has the right to access, without restriction to all Project documentation of the AIIMS Jodhpur including that provided by the Contractor. The Contractor will also grant the Monitor, upon his/her request and demonstration of a valid interest, unrestricted and unconditional access to their project documentation. The same is applicable to Sub- contractors.
4. The Monitor is under contractual obligation to treat the information and documents of the Bidder (s)/ Contractor(s)/ Sub-contractor(s) with confidentiality. The Monitor has also signed declarations on Non-Disclosure of Confidential Information and of 'Absence of conflict of Interest'. In case of any conflict of interest arising at a later date, the IEM shall inform Executive Director AIIMS Jodhpur and recuse himself/herself from that case.
5. The AIIMS Jodhpur shall provide to the Monitor sufficient information about all meetings among the parties related to the Project provided such meetings could have an impact on the contractual relations between the Principal and the Contractor. The parties offer to the Monitor the option to participate in such meetings.
6. As soon as the Monitor notices, or believes to notice, a violation of this agreement, he/she will so inform the Executive Director AIIMS Jodhpur and request the Management to discontinue or take corrective action, or the take other relevant action. The monitor can in the regard submit non-binding recommendations. Beyond this, the Monitor has no right to demand from the parties that they actin a specific manner, refrain from action or tolerate action.
7. The Monitor shall submit a written report to the Executive Director AIIMS Jodhpur, within 8 to 10

weeks from the date of reference or intimation to him by the AIIMS Jodhpur and, should the occasion arise, submit proposals for correcting problematic situations.

8. If the Monitor has reported to the Executive Director AIIMS Jodhpur, a substantiated suspicion of an offence under relevant IPC/PC Act, and the Executive Director, AIIMS Jodhpur has not, within the reasonable time taken visible action to proceed against such offence or reported it to the Chief Vigilance officer, the Monitor may also transmit this information directly to the Central Vigilance Commissioner.
9. The word Monitor, would include both singular and plural.

Section - 9 Pact Duration

This Pact begins when both parties have legally signed it. It expires for the Contractor 12 months after the last payment under the respective contract, and for all other Bidders 6 months after the contract has been awarded. Any violation of the same would entail disqualification of the bidders and exclusion from future business dealings.

If any claim is made / lodged during this time the same shall be binding and continue to be valid despite the lapse of this pact as specified above, unless it is discharged / determined by Executive Director of AIIMS Jodhpur.

Section - 10 Other Provisions

1. This agreement is subject to Indian Law. Place of performance and jurisdiction is the AIIMS Jodhpur.
2. Changes and supplements as well as termination notices need to be made in writing. Side agreements have not been made.
3. If the Contractor is a partnership or a consortium, this agreement must be signed by all partners or consortium members.
4. Should one or several provisions of this agreement turn out to be invalid, the remainder of this agreement remains valid. In this case, the parties will strive to come to an agreement to their original intentions.
5. Issues like comprehensive Warranty / Guarantee etc. shall be outside the purview of IEMs.
6. In the event of any contradictions between the Integrity Pact and its Annexure, the Clause in the Integrity Pact will prevail.

For and on behalf of the AIIMS Jodhpur	For & on behalf of Bidder/Contractor
Office Seal	Office Seal
Place:	Witness 1:
Date :	Witness 2:

Annexure-1 'List of Quoted Items'

(THIS IS FOR REFERENCE ONLY

SUBMIT THIS ANNEXURE IN THE GIVEN EXCEL SHEET)

To

The Executive Director,
All India Institute of Medical Sciences, Jodhpur
MIA Phase-II, Basani, Jodhpur

NIT No. _____ _____ _____ _____ ____ dated: _____ for Rate Contract _____ _____ _____ _____ at AIIMS Jodhpur.

Sr.	Item Code	*Item Name*	Original Manufacture's name	Name of the Authorization Certificate issuer (this column is to be filled, if the Authorization Certificate has been issued by other than the Original Manufacturer)	Brand Name	(Make/Model No.) of the Item	Pack Size	Accounting Unit A/U	Page No. manufacturer/ import licences attached in Technical Bid Document	Page No. USFDA/CE/DCGI certificate attached in Technical Bid Document	Whether complying with NIT's Specs (Yes/No)	Mention here - if bidder is quoting their item with any different specification or size
	a	b	c	d	e	f			g	h	i	j
1.												
2.												
3.												
4.												
.												
..												
...												

Name of the Key Person: _____
Designation: _____
Contact: _____
eMail ID: _____
Office Address: _____
_____

Place: _____

Date: _____

Note:

1. This Annexure-1 [List of Quoted Items] is a part of Technical Bid Document.
2. Bidder needs to submit this Annexure-1 [List of Quoted Items] in the given Excel Sheet along with the Technical Bid Document.

Annexure-2 'Calculation of Local Content'

(To be submitted along with BoQ only)

To

The Executive Director,
All India Institute of Medical Sciences, Jodhpur
MIA Phase-II, Basani, Jodhpur

NIT No. _____ _____ _____ _____ ____ dated: _____ for Rate Contract _____ _____ _____ _____ at AIIMS Jodhpur.

Name of Manufacture	Calculation by Manufacturer (Cost per unit of product)			
Cost Component	Cost (Domestic Component) A	Cost (Imported Component) B	Total Cost (INR/ US $) C=a+b	Percentage of Local Content D=(a/c)*100
I. …… II. …… III. Total Cost (Excluding tax and duties)				

Note:-

I. Cost (Domestic Component): Sum of the costs of all inputs which go into the product (including duties and taxes levied on procurement of inputs except those for which credit/ set-off can be taken) which have not been imported directly or through a domestic trader or an intermediary.

II. Cost (Imported Component): Sum of the costs of all inputs which go into the product (including duties and taxes levied on procurement of inputs except those for which credit/ set-off can be taken).

Name: _____

(Signatures of the Bidder with Name, Designation & Company's Seal)

Address: _____

_____

Place: _____

Date: _____

Note:

1. This Annexure-2 [Calculation of Local Content] is to be submitted with BoQ [Financial Bid] only.
2. Your bid shall be REJECTED, if the Annexure-2 [Calculation of Local Content] is submitted in the Technical Bid Document.

Annexure-3 'Item List'

(Items proposed to be procured through this Rate Contract)

	Item Code	Name of Item	Specifications (Accounting Unit, Size, shape & other details)
S. No.	**1.00**	**CLOSURE DEVICES – ASD**	
1	1.01	ASD CLOSURE DEVICE WITHOUT DELIVERY SYSTEM – USFDA APPROVED	• Must be available in all approved sizes • Must be approved for pediatric and adult use • Self-centering, detachable device with delivery cable • Device and all accessories must be made up of biologically inert material • Device should be coated to prevent nickel leaching
2	1.02	ASD CLOSURE DEVICE WITHOUT DELIVERY SYSTEM – CE MARKED	• Must be available in all approved sizes • Must be approved for pediatric and adult use • Self-centering, detachable device with delivery cable • Device and all accessories must be made up of biologically inert material • Device should be coated to prevent nickel leaching
3	1.03	ASD CLOSURE DEVICE WITH EXISTING FENESTRATION WITHOUT DELIVERY SYSTEM – USFDA APPROVED	• Must be available in all approved sizes • Must be approved for pediatric and adult use • Self-centering, detachable device with delivery cable • Device and all accessories must be made up of biologically inert material • Device should be coated to prevent nickel leaching
4	1.04	ASD CLOSURE DEVICE WITH EXISTING FENESTRATION WITHOUT DELIVERY SYSTEM – CE MARKED	• Must be available in all approved sizes • Must be approved for pediatric and adult use • Self-centering, detachable device with delivery cable • Device and all accessories must be made up of biologically inert material • Device should be coated to prevent nickel leaching

5	1.05	DELIVERY SYSTEM FOR ASD DEVICE – USFDA APPROVED	• Available in all approved sizes • Approved for use in children and adults • Should comprise of 45° curve delivery sheath with a check valve mechanism • Sheath should be kink resistant with reinforcement with internal braided layer made of stainless steel • Should have a soft radiopaque tip • Dilator with tapering tip should fit in the sheath locking mechanism • Loader, plastic vise and delivery cable should be included
6	1.06	DELIVERY SYSTEM FOR ASD DEVICE – CE MARKED	• Available in all approved sizes • Approved for use in children and adults • Should comprise of 45° curve delivery sheath with a check valve mechanism • Sheath should be kink resistant with reinforcement with internal braided layer made of stainless steel • Should have a soft radiopaque tip • Dilator with tapering tip should fit in the sheath locking mechanism • Loader, plastic vise and delivery cable should be included
7	1.07	SPECIAL ASD CLOSURE DEVICE FOR VERY LARGE ASD	• Must be available in sizes greater than 40 mm • Must be approved for use in adults • Self-centering, detachable device with delivery cable • Device and all accessories must be made up of biologically inert material • Device should be coated to prevent nickel leaching
8	1.08	SPECIAL ASD CLOSURE DEVICE FOR FENESTRATED DEFECTS	• Must be available in all approved sizes • Must be approved for pediatric and adult use • Self-centering, detachable device with delivery cable • Device and all accessories must be made up of biologically inert material • Device should be coated to prevent nickel leaching
9	1.09	ASD Device-DGCI Approved	• Braided and made up of Nitinol mesh that is tightly woven to reduce the residual shunt after positioning. • Available sizes: Dia Waist-6,8,10,12,14,16,18,20,22,24,26,28,30,32,34,36,38,40,42mm. • Compatible with 6F - 14F Sheath

	2.00	**CLOSURE DEVICES – VSD**	
10	2.01	MUSCULAR VSD CLOSURE DEVICES WITHOUT DELIVERY SYSTEM – USFDA APPROVED	• Must be available in all approved sizes • Must be approved for pediatric and adult use • Self-centering, detachable device with delivery cable • Device must be made up of biologically inert material • Device should be coated to prevent nickel leaching
11	2.02	MUSCULAR VSD CLOSURE DEVICES WITHOUT DELIVERY SYSTEM – CE MARKED	• Must be available in all approved sizes • Must be approved for pediatric and adult use • Self-centering, detachable device with delivery cable • Device must be made up of biologically inert material • Device should be coated to prevent nickel leaching
12	2.03	ASYMMETRIC VSD CLOSURE DEVICES WITHOUT DELIVERY SYSTEM – USFDA APPROVED	• Must be available in all approved sizes • Must be approved for pediatric and adult use • Self-centering, detachable device with delivery cable • Device must be made up of biologically inert material • Device should be coated to prevent nickel leaching
13	2.04	ASYMMETRIC VSD CLOSURE DEVICES WITHOUT DELIVERY SYSTEM – CE MARKED	• Must be available in all approved sizes • Must be approved for pediatric and adult use • Self-centering, detachable device with delivery cable • Device must be made up of biologically inert material • Device should be coated to prevent nickel leaching
14	2.05	SPECIAL DOUBLE DISC MULTIPURPOSE OCCLUDER FOR VSD WITHOUT DELIVERY SYSTEM	• Must be available in all approved sizes • Must be approved for pediatric and adult use • Device must be made up of biologically inert material • Central waist and double-disc design for occluding various defects • Screwable from both the discs to allow retrograde and antegrade delivery of device • Soft material high conformability • Very Low profile • The vendor should be able to supply all the components of the delivery system (delivery sheath and cable)

	3.00	**CLOSURE DEVICES- PDA**	
15	3.01	PDA CLOSURE DEVICE WITHOUT DELIVERY SYSTEM – USFDA APPROVED	• Must be available in all approved sizes • Must be approved for pediatric and adult use • Self-centering, detachable device with delivery cable • Device must be made up of biologically inert material • Device should be coated to prevent nickel leaching
16	3.02	PDA CLOSURE DEVICE WITHOUT DELIVERY SYSTEM – CE MARKED	• Must be available in all approved sizes • Must be approved for pediatric and adult use • Self-centering, detachable device with delivery cable • Device must be made up of biologically inert material • Device should be coated to prevent nickel leaching
17	3.03	SYMMETRICAL PDA CLOSURE DEVICE WITH RETENTION DISC ON BOTH SIDES FOR USE IN PRETERM NEONATES	• Must be available in all approved sizes • Must be approved for use in neonates and preterm neonates • Self-centering, detachable device with delivery cable • Device and all accessories must be made up of biologically inert material • Device should be coated to prevent nickel leaching • Device should have a symmetric design enabling both antegrade and retrograde delivery
18	3.04	DELIVERY SYSTEM FOR PDA DEVICE – USFDA APPROVED	• Approved for use in infants, children and adults• Available in all approved sizes (including 4 and 5 Fr)• Should comprise of 180° curve delivery sheath with check valve mechanism• Soft radiopaque tip• Dilator with tapering tip should fit in the sheath locking mechanism• Loader, plastic vise and delivery cable should be included

19	3.05	DELIVERY SYSTEM FOR PDA DEVICE – CE MARKED	• Approved for use in infants, children and adults • Available in all approved sizes (including 4 and 5 Fr) • Should comprise of 180° curve delivery sheath with check valve mechanism • Soft radiopaque tip • Dilator with tapering tip should fit in the sheath locking mechanism • Loader, plastic vise and delivery cable should be included • Available in all approved sizes for adult and pediatric usage
20	3.06	DELIVERY SYSTEM FOR PDA DEVICE WITH SYMMETRICAL DISC ON BOTH SIDES	• Approved for use in infants, children and adults • Compatible with 4 and 5 Fr sheath • Should comprise of 180° curve delivery sheath with check valve mechanism • Soft radiopaque tip • Dilator with tapering tip should fit in the sheath locking mechanism • Loader, plastic vise and delivery cable should be included
21	3.07	PDA CLOSURE DEVICES FOR LARGE SIZE PDAs	• Must be approved for adult use • Self-centering, detachable device with delivery cable • Device and all accessories must be made up of biologically inert material • Amplatzer duct occlude type devices with aortic end of device measuring greater than 24 mm
22	3.08	PDA Occluder- DCGI Approved	•Avialable sizes: 4-6,6-8,8-10,10-12,12-14,14-16,16-18,18-20,20-22,22-24 mm. • Compatible with 5F - 10F
	4.00	**CLOSURE DEVICES - PFO**	
23	4.01	PFO CLOSURE DEVICES WITHOUT DELIVERY SYSTEM – USFDA APPROVED	• Must be available in all approved sizes • Must be approved for pediatric and adult use • Self-centering, detachable device with delivery cable • Device and all accessories must be made up of biologically inert material • Device should be coated to prevent nickel leaching

24	4.02	PFO CLOSURE DEVICES WITHOUT DELIVERY SYSTEM – CE MARKED	• Must be available in all approved sizes • Must be approved for pediatric and adult use • Self-centering, detachable device with delivery cable • Device and all accessories must be made up of biologically inert material • Device should be coated to prevent nickel leaching
	5.00	**VASCULAR PLUGS & CLOSURE DEVICES**	-
25	5.01	VASCULAR PLUG (SINGLE LOBE DESIGN) – USFDA APPROVED	• Must be available in all approved sizes • Must be approved for pediatric and adult use • Self-centering, detachable device with delivery cable • Device and all accessories must be made up of biologically inert material
26	5.02	VASCULAR PLUG (SINGLE LOBE DESIGN) – CE MARKED	• Must be available in all approved sizes • Must be approved for pediatric and adult use • Self-centering, detachable device with delivery cable • Device and all accessories must be made up of biologically inert material
27	5.03	VASCULAR PLUG (MULTISEGMENTED AND OBLONG DESIGN) – USFDA APPROVED	• Must be available in all approved sizes• Must be approved for pediatric and adult use• Self-centering, detachable device with delivery cable• Device and all accessories must be made up of biologically inert material
28	5.04	VASCULAR PLUG (MULTISEGMENTED AND OBLONG DESIGN) – CE MARKED	• Must be available in all approved sizes • Must be approved for pediatric and adult use • Self-centering, detachable device with delivery cable • Device and all accessories must be made up of biologically inert material
29	5.05	VASCULAR PLUGS FOR SMALL VESSELS LESS THAN 15 MM – USFDA APPROVED	• Must be available in all approved sizes • Must be approved for neonatal, pediatric and adult use • Self-centering, detachable device with delivery cable • Device and all accessories must be made up of biologically inert material • Compatible with 4Fr guiding catheter or 5Fr diagnostic catheter

30	5.06	VASCULAR PLUGS FOR SMALL VESSELS LESS THAN 15 MM – CE MARKED	• Must be available in all approved sizes • Must be approved for neonatal, pediatric and adult use • Self-centering, detachable device with delivery cable • Device and all accessories must be made up of biologically inert material • Compatible with 4Fr guiding catheter or 5Fr diagnostic catheter
31	5.07	VASCULAR PLUGS FOR VERY SMALL VESSELS LESS THAN 5 MM – USFDA APPROVED	• Must be available in all approved sizes • Must be approved for neonatal, pediatric and adult use • Self-centering, detachable device with delivery cable • Device and all accessories must be made up of biologically inert material • The smallest size devices should be deliverable by a microcatheter
32	5.08	VASCULAR PLUGS FOR VERY SMALL VESSELS LESS THAN 5 MM – CE MARKED	• Must be available in all approved sizes • Must be approved for neonatal, pediatric and adult use • Self-centering, detachable device with delivery cable • Device and all accessories must be made up of biologically inert material • The smallest size devices should be deliverable by a microcatheter
33	5.09	DEVICES FOR PARAVALVULAR LEAK CLOSURE WITHOUT DELIVERY SYSTEM	• Device designed and dedicated for paravalvular leak closure • Available in all approved dimensions and shapes to handle leaks of different morphologies • Device must be made up of biologically inert material • Device should be coated to prevent nickel leaching • Repositionable and retrievable device
	6.00	**ACCESSORIES FOR DEVICE CLOSURE**	
34	6.01	SIZING BALLOON FOR ASD DEVICE CLOSURE: OVAL SHAPED	• Compliant balloon material flexible shaft and soft tip • Radio-opaque markers bands • Available in all approved sizes • Over the wire: compatible with 0.035 wire
35	6.02	SIZING BALLOON FOR ASD DEVICE CLOSURE: CIRCULAR SHAPE	• Compliant balloon material flexible shaft and soft tip • Radio-opaque markers bands • Available in all approved sizes • Over the wire: compatible with 0.035 wire

36	6.03	SIZING PLATE FOR BALLOON SIZING OF ASD	• Measuring plate with circular openings range from 4 to 38 mm
	7.00	**LEFT ATRIAL APPENDAGE OCCLUDER**	-
37	7.01	LAA CLOSURE DEVICE WITHOUT DELIVERY SYSTEM – CE MARKED	• Must be available in all approved sizes• Must be approved for adult use• Self-centering, detachable device with delivery cable• Device and all accessories must be made up of biologically inert material• Device should be coated to prevent nickel leaching
38	7.02	LAA CLOSURE DEVICE WITHOUT DELIVERY SYSTEM – USFDA APPROVED	• Must be available in all approved sizes • Must be approved for adult use • Self-centering, detachable device with delivery cable • Device and all accessories must be made up of biologically inert material • Device should be coated to prevent nickel leaching
39	7.03	DELIVERY SYSTEM FOR LAA CLOSURE DEVICE	• Approved for adult's use • Should comprise of DUAL ANGELED delivery sheath with check valve mechanism • Soft radiopaque tip • Dilator with tapering tip should fit in the sheath locking mechanism • Loader, plastic vise and delivery cable should be included
	8.00	**BALLON DILATATION CATHETERS**	-
40	8.01	Mullin's Sheath for special Dilation - DCGI With US FDA /CE Approved	Should be in All Sizes-6F & above septal puncture needle.
41	8.02	Trans-Spetal puncture needle-DCGI/US FDA /CE Approved	
42	8.03	SPECIAL SWAN GANZ CATHETER - US FDA APPROVED	• 4 F catheter should allow the passage of at least 0.021 inch, 5 F – 0.025 inch, 6 F – 0.035 inch and 7 F, 8F – 0.038-inch hydrophilic guide wire • Must be available in 4F, 5F, 6F and 7F and 8 Fr. sizes • Catheter should be tapered at tip to ensure uniform diameter of the whole catheter • 10 cm marking along catheter body to confirm insertion depth

43	8.04	SPECIAL SWAN GANZ CATHETER - CE APPROVED	• 4 F catheter should allow the passage of at least 0.021 inch, 5 F – 0.025 inch, 6 F – 0.035 inch and 7 F, 8F – 0.038-inch hydrophilic guide wire • Must be available in 4F, 5F, 6F and 7F and 8 Fr. sizes • Catheter should be tapered at tip to ensure uniform diameter of the whole catheter • 10 cm marking along catheter body to confirm insertion depth
44	8.05	SPECIAL SWAN GANZ CATHETER - DCGI APPROVED	• 4 F catheter should allow the passage of at least 0.021 inch, 5 F – 0.025 inch, 6 F – 0.035 inch and 7 F, 8F – 0.038-inch hydrophilic guide wire • Must be available in 4F, 5F, 6F and 7F and 8 Fr. sizes • Catheter should be tapered at tip to ensure uniform diameter of the whole catheter • 10 cm marking along catheter body to confirm insertion depth
45	8.06	BERMAN CATHETER - DCGI With US FDA /CE Approved	· Must be available in 4F, 5F, 6F and 7F sizes · Should have 6-8 holes proximal to the balloon for dye injection · Catheter should be tapered at tip to ensure uniform diameter of the whole catheter · 10 cm marking along catheter body to confirm insertion depth
46	8.07	REVERSE BERMAN CATHETER - DCGI With US FDA /CE Approved	. Should be available in 4F, 5F, 6F & 7F sizes . Catheter should be tapered at tip to ensure uniform diameter of the whole catheter . Should have holes proximal to the balloon for dye injection . Should have a hole at the proximal tip to allow the passage over the wire
47	8.08	VALVOPLASTY BALLOON CATHETERS (sizes 10 – 22 mm) - US FDA APPROVED	· Approved for pediatric/adults use · Varying length and diameter · Nominal pressure < 4 ATM
48	8.09	VALVOPLASTY BALLOON CATHETERS (sizes 10 – 22 mm) - CE APPROVED	· Approved for pediatric/adults use · Varying length and diameter · Nominal pressure < 4 ATM
49	8.10	BALLOON CATHETERS (sizes 10 – 22 mm) - DCGI APPROVED	· Approved for pediatric/adults use · Varying length and diameter · Nominal pressure < 8 ATM
50	8.11	BALLOON IN BALLOON CATHETERS - US FDA APPROVED	· Quote in all sizes
51	8.12	BALLOON IN BALLOON CATHETERS - CE APPROVED	· Quote in all sizes
52	8.13	BALLOON IN BALLOON CATHETERS - DCGI APPROVED	· Quote in all sizes

53	8.14	STEERABLE SHEATH WITH DEFLECTABLE TIP (190-160DEGREE) - DCGI with US FDA / CE Approved	
54	8.15	SPECIAL BALLOON CATHETER COMPATIBLE WITH 4 FRENCH SHEATH - US FDA APPROVED	· Compatible with 0.018-inch guide wire · Varying length and diameter
55	8.16	SPECIAL BALLOON CATHETER COMPATIBLE WITH 4 FRENCH SHEATH - CE APPROVED	· Compatible with 0.018-inch guide wire · Varying length and diameter
56	8.17	SPECIAL BALLOON CATHETER COMPATIBLE WITH 4 FRENCH SHEATH – DCGI APPROVED	· Compatible with 0.018-inch guide wire · Varying length and diameter
57	8.18	SPECIAL PEDIATRIC VALVOPLASTY BALLOON CATHETERS COMPATIBLE WITH 4 FRENCH SHEATH - US FDA APPROVED	· Compatible with 0.025-inch guide wire · Varying length and diameter
58	8.19	SPECIAL PEDIATRIC VALVOPLASTY BALLOON CATHETERS COMPATIBLE WITH 4 FRENCH SHEATH – CE APPROVED	· Compatible with 0.025-inch guide wire · Varying length and diameter
59	8.20	SPECIAL PEDIATRIC VALVOPLASTY BALLOON CATHETERS COMPATIBLE WITH 4 FRENCH SHEATH – DCGI APPROVED	· Compatible with 0.025-inch guide wire · Varying length and diameter
60	8.21	BIB (BALLOON IN BALLOON) TYPE BALLOONS FOR COARCTATION DILATATION - US FDA/CE/DCGI APPROVED	· Monorail (rapid exchange) and/OR over-the-wire (OTW) balloons · Available in all sizes and lengths
61	8.22	Z MED TYPE BALLOON FOR AORTIC COARCTATION – US FDA/CE APPROVED	· Different sizes AND length · Over the wire / Rapid exchange, compatible with 0.014”/0.035” and 0.038” wire
62	8.23	C.P. STENT COVERED – CE APPROVED	· Should be balloon Mounted.
63	8.24	C.P. STENT COVERED – US FDA APPROVED	Should be balloon Mounted.
64	8.25	C.P. STENT COVERED – DCGI APPROVED	Should be balloon Mounted.
65	8.26	C.P. STENT UNCOVERED– CE APPROVED	· Balloon Mounted for Aortic Use.
66	8.27	C.P. STENT UNCOVERED – US FDA APPROVED	Balloon Mounted for Aortic Use.
67	8.28	C.P. STENT UNCOVERED– DCGI APPROVED	Balloon Mounted for Aortic Use.
68	8.29	Mitral valve Dilation Balloon (2 port Balloon catheter) with Accessories US-FDA approved	
69	8.30	Mitral valve Dilation Balloon (2 port Balloon catheter) without Accessories US-FDA approved	
70	8.31	Mitral valve Dilation Balloon (2 port Balloon catheter) with Accessories CE approved	

71	8.32	Mitral valve Dilation Balloon (2 port Balloon catheter) without Accessories CE approved	
72	8.33	Mitral valve Dilation Balloon (2 port Balloon catheter) with Accessories DCGI approved	
73	8.34	Mitral valve Dilation Balloon (2 port Balloon catheter) without Accessories DCGI approved	
74	8.35	Mitral valve Dilation Balloon (3 port Balloon catheter) with Accessories US-FDA approved	
75	8.36	Mitral valve Dilation Balloon (3 port Balloon catheter) without Accessories US-FDA approved	
76	8.37	Mitral valve Dilation Balloon (3 port Balloon catheter) with Accessories CE approved	
77	8.38	Mitral valve Dilation Balloon (3 port Balloon catheter) without Accessories CE approved	
78	8.39	Mitral valve Dilation Balloon (3 port Balloon catheter) with Accessories DCGI approved	
79	8.40	Mitral valve Dilation Balloon (3 port Balloon catheter) without Accessories DCGI approved	
80	8.41	Delivery catheter system for balloon-expandable valve for TAVR- US-FDA approved	
81	8.42	Delivery catheter system for balloon-expandable valve for TAVR- CE approved	
82	8.43	Delivery catheter system for balloon-expandable valve for TAVR- DCGI approved	
83	8.44	Delivery catheter system for Self expanding valve Intra annular for TAVR-US-FDA approved	
84	8.45	Delivery catheter system for Self expanding valve Intra annular for TAVR-CE approved	
85	8.46	Delivery catheter system for Self expanding valve Intra annular for TAVR-DCGI approved	

86	8.47	Delivery catheter system for Self expanding valve Supraannular for TAVR- US-FDA approved	
87	8.48	Delivery catheter system for Self expanding valve Supra annular for TAVR- CE approved	
88	8.49	Delivery catheter system for Self expanding valve Supra annular for TAVR- DCGI approved	
89	8.50	Loading system for balloon-expandable valve for TAVR- US-FDA approved	
90	8.51	Loading system for balloon-expandable valve for TAVR- CE approved	
91	8.52	Loading system for balloon-expandable valve for TAVR- DCGI approved	
92	8.53	Loading system for Self expanding valve Intra annular for TAVR- US-FDA approved	
93	8.54	Loading system for Self expanding valve Intra annular for TAVR- CE approved	
94	8.55	Loading system for Self expanding valve Intra annular for TAVR- DCGI approved	
95	8.56	Loading system for Self expanding valve Supraannular for TAVR- US-FDA approved	
96	8.57	Loading system for Self expanding valve Supra annular for TAVR- CE approved	
97	8.58	Loading system for Self expanding valve Supra annular for TAVR- DCGI approved	
98	8.59	TAVR Valve – DCGI/CE Approved	Balloon Expandable Transcatheter Heart Valve system, Unique hybrid-honey comb cell design (only 3 rows of hexagons) Open cells- 53%, Close cells 47%, available in large size matrix (conventional, intermediate & extra-large) available sizes: Dia-20.0, 21.5, 23.0, 24.5, 26.0, 27.5, 29.0, 30.5, 32.0 mm, comes with 14 Fr Python Introducer Sheath compatible with all sizes, Navigator THV Delivery System, Mammoth Predilation Balloon, Transcatheter crimping tool.

99	8.60	TAVR Valve – DCGI/ CE Approved	Balloon Expandable Transcatheter Heart Valve system, Unique hybrid-honey comb cell design (only 3 rows of hexagons) Open cells- 53%, Close cells 47%, available in large size matrix (conventional, intermediate & extra-large) available sizes: Dia-20.0, 21.5, 23.0, 24.5, 26.0, 27.5, 29.0, 30.5, 32.0 mm, comes with 14 Fr Python Introducer Sheath compatible with all sizes, Navigator THV Delivery System, Mammoth Predilation Balloon, Transcatheter crimping tool.
100	8.61	TAVR Valve –USFDA Approved	Percutaneous Transcatheter Self-expanding intra-annular bovine tissue heart valve for Aortic Heart position available in sizes 23mm-29mm with low profile Delivery System. Valves should have Anti-calcification treatment. Polyethylene fabric material with dynamic cuff that synchronizes with the cardiac cycle. Compatible with 14F & 15F low profile delivery system.
101	8.62	TAVR Valve –USFDA Approved	Percutaneous Transcatheter Self-expanding intra-annular bovine tissue heart valve for Aortic Heart position available in sizes 23mm-29mm with low profile Delivery System. Valves should have Anti-calcification treatment. Polyethylene fabric material with dynamic cuff that synchronizes with the cardiac cycle. Compatible with 14F & 15F low profile delivery systemShould have marker with Vision technology for depth assesment
102	8.63	TMVR Valve - DCGI approved	
103	8.64	Transcatheter Edge to Edge Repair (TEER) System	• Made up of Cobalt Chromium covered with Polyester fabric & Nitinol graspers, • Clip-Arms Widths: Regular: 3 & 4mm, Wide: 5 & 6mm, Extra-wide: 7mm, and Clip arm length: Short (S): 9mm, Long (L): 12mm, • comes with MyClip Guide Catheter (MGC), Delivery System (MDS), Accessories - Ground-Plate, Podium, Anti-Skid Silicone Mat, Bracket. • 2 years of Shelf Life.
104	8.65	Predilation Balloon for TAVR- CE Approved	
105	8.66	Predilation Balloon for TAVR- DCGI approved	
106	8.67	Post Dilation Balloon suitable for TAVR-USFDA approved	

107	8.68	Post Dilation Balloon suitable for TAVR- CE approved	
108	8.69	Post Dilation Balloon suitable for TAVR- DCGI approved	
109	8.70	Wire for TAVI procedure DCGI with US FDA / CE Approved	
110	8.71	Sheath for TAVI procedure DCGI with US FDA / CE Approved	
111	8.72	Transcatheter Aortic valve Balloon expandable type US FDA approved	
112	8.73	Transcatheter Aortic valve Balloon expandable type CE approved	
113	8.74	Transcatheter Aortic valve Balloon expandable type DCGI approved	
114	8.75	Transcather Aortic valve Self expanding type intra annular valve position US FDA approved	
115	8.76	Transcather Aortic valve Self expanding type intra annular valve position CE approved	
116	8.77	Transcather Aortic valve Self expanding type intra annular valve position DCGI approved	
117	8.78	Transcather Aortic valve Self expanding type supra-annular valve position US FDA approved	
118	8.79	Transcather Aortic valve Self expanding type supra-annular valve position CE approved	
119	8.80	Transcather Aortic valve Self expanding type supra-annular valve position DCGI approved	
120	8.81	Transcather Aortic valve Implantation (TAVI) with dry tissue technology, preloaded delivery system, all sizes DCGI/CE APPROVED/USFDA	
121	8.82	Pulmonary Valvuloplasty Balloon (VACS II) (USFDA/CE/DCGI APPROVED)	• Low profile valvuloplasty catheter with maximum steribility and trackability with alternative tip, radiopaque gold markers, short inflation and deflation duration sizes with • Balloon diameter 4–30 mm • Balloon length 20–60 mm Rated Burst Pressure 6–1.5 atm

122	8.83	Aortic Valvuloplasty Balloon (VACS III) (USFDA/CE/DCGI APPROVED)	• High pressure valvuloplasty balloon catheter with maximum steribility and trackability with alternative tip, radiopaque gold markers for TAVI, aortiv valves stenosis. • Balloon diameter 5–30 mm • Balloon length 20–60 mm • Rated Burst Pressure 15–4 atm
123	8.84	Vascular closure device with Bio absorbable PGA suture that provide a secure seal* - USFDA Approved	• Bioabsorbable collagen designed, • low profile Bioabsorbable Anchor (50:50 blend glycolide and lactide polyer) • Must have bovine collagen plug. Must have 9 hole V-twist pattern suture weave. • Must be available in 6F and 8F sizes.
124	8.85	TAVI Guidewire - (USFDA/CE APPROVED)	Outer Diameter 0.035" (0.889 mm), Length 275 cm, Core Material Stainless Steel, Spring Coil Material Stainless Steel, Spring Coil Coating LUBRIGREEN™ PTFE, **Curves** Extra small (16cm x 3.2cm x 2.9cm), Small (16cm x 4.2cm x 4.2cm), Large (18cm x 5.0cm x 4.9cm)
125	**8.86**	Pre Balloon mounted covered stent-Perpheral - USFDA/CE APPROVED	Pre Balloon mounted covered stent, Peripheral stent with high radial force and low profile. Cobalt-chrome platform with micro-porous ePTFE membarne 6Fr / 7Fr introducer compatible, diameters (5mm, 6mm, 7mm, 8mm with overexpansion upto 10mm and 9mm, 10mm with over expansion upto 12mm) all lengths (17mm, 27mm, 37mm, 57mm) over 0.035 guidewire
126	**8.87**	Pre Balloon mounted covered stent-Aortic - (USFDA/CE APPROVED)	Pre balloon mounted Aortic stent grafts 0.035 wire caompatible, Sizes - Diameter 12mm to 24mm, Length 19mm t0 59 mm, Aortic stent graft with low foreshortening with high radial force. Cobalt chromium stent is covered with 200 um ePTFE tubing Peripheral stent with high radial force and low profile. Cobalt-chrome platform with micro-porous ePTFE membrane
127	8.88	PTA Balloon - (USFDA/CE APPROVED)	Peripheral PTA High Pressure Balloon, Diameter in mm: 3.00, 3.50, 4.00, 5.00, 6.00, 7.00, 8.00, 9.00 10.00; Balloon length in mm: 20, 30, 40, 60, 80, 100, 120, 150, 180, 200;Guidewire compatibility: 0.035.
128	8.89	Adult Valvuloplasty Balloon) - (USFDA/CE APPROVED)	Semi Compliant Double lumen balloon Catheter for Adult Valvuloplasty. Diameter 12 mm to 40mm & length 40 mm to 60 mm. Total shaft length 110cm. 0.038 compatible

129	8.90	Pediatric Valvuloplasty Balloon – (USFDA/CE APPROVED)	Semi Compliant Double lumen balloon Catheter for Pediatric Valvuloplasty. Diameter 8mm to 15 mm & length 20 mm to 40 mm. Total shaft length 70cm. 0.035 compatible
130	8.91	Self-Expanding Endovascular Stent Graft for repair of Aneurysm involving the ARCH/Thoracic/Abdominal Aorta	• Stainless Steel stent Graft device covered by woven polyster graft material with one,two , three or multiple components • preloaded into 18,20,22,23 F,Flexible hydrophilic coated • USFDA/ CE Approved
131	8.92	Perclose Prostyle	* Vascular Closure Device for 5-22F * polypropylene monofilament suture mediated vessel closure * with depth markers on device and Knot Pusher/Suture trimmer * comes with a Snare Knot Pusher * With no re - access restriction * Immediate re-access advantage * Flexibility to pre-close and close over the wire * (USFDA Approved) * For small hole and large hole closure both
132	8.93	Proglide	• Vascular Closure Device for 5–22F • Polypropylene monofilament suture mediated vessel closure • With no re-access restriction • Immediate re-access advantage • Flexibility to pre-close and close over the wire • USFDA Approved • For small hole and large hole closure both.
133	8.94	Vascular closure device with Bio absorbable PGA suture that provide a secure seal* - USFDA Approved	• Bioabsorbable collagen designed, • low profile Bioabsorbable Anchor (50:50 blend glycolide and lactide polyer) • Must have bovine collagen plug. Must have 9 hole V-twist pattern suture weave. • Must be available in 6F and 8F sizes.

134	8.95	Sequre Snare System Retrieval Device	• Titanium Nitride Coated Tungsten Loop provides excellent grip and bio-compatibilty• Enhanced fracture resistance and radio opacity• Excellent shape supporting and distal shaft control ability• Flexible Nitinol shaft provides Kink Resistance and 1:1 Torque control• 90°angle between Loop and Shaft makes manipulation and retrieval more simple• Radiopaque marker at distal tip of catheter provides excellent Fluoroscopic visualization• A wide range of usage for clinical procedures• Available Sizes 5mm-35mm• USFDA,CE,DCGI
135	8.96	Vascular Closure Device – DGCI Approved	• Bio-absorbable Collagen (Bovine Type-I) based vascular closure device with bio-absorbable anchor (PDLG) and PGA suture. Require no exchange of introducer sheath. Device Size- 6Fr and 8Fr; complete degradation- 90 days. Device Total length (mm)- 205 ±10 and device ffective length (mm)- 155 ±10
136	8.97	Flexible steerable introducer sheath. – CE Approved	• It should hydrophilic coated. It should available in sizes of 5fr-16fr and 55cm-90cm. It should have ergonomic rotating knob.
137	8.98	Transcatheter Aortic Heart Valve system	• size 23-34mm with Self Expanding, supra annular, Recapturable Porcine Pericardium leaflets • Nitinol Frame valve with external skirt & compatible delivery catheter and loading system • USFDA Approved
138	9.01	SAPIEN 3 Valve - Special Balloon Expading Trileaflet Bovine Pericadial Tissue Vave with Delivery System for TAVR	* Should be balloon - expandable transcatheter Aortic Valve System with delivery accessories * Should have a Cobalt Chromium frame and an inner and outer PET skirt to minimize paravalvular leak * Must be available in sizes of 20,23,26,29 mm * Valve must be crimped away from the Balloon * Should have been treated with Thermafix process for calcification USFDA Approved

139	9.02	SAPIEN 3 Ultra Resilia Valve - Special Balloon Expading Trileaflet Resilia Technology Bovine Pericadial Tissue Vave with Delivery System for TAVR	* Should be balloon - expandable transcatheter Aortic Valve System with delivery accessories * Should have a Cobalt Chromium frame and an inner and outer PET skirt to minimize paravalvular leak * Must be available in sizes of 20,23,26,29 mm * Valve must be crimped away from the Balloon * Should use "Dry Tissue " preservative technique, allowing it to be stored dry - USFDA Approved