

Standard Bidding Document

2nd Re-Tender for Electro-Medical Equipment/s of Cardiology Department,
under PSDP Project titled 'Strengthening and Up-Gradation of Cardiology
and other Departments during the FY 2025-26 & 2026-27
(Goods)

National

Single Stage-Two Envelope



October 13, 2026

*FEDERAL GOVERNMENT POLYCLINIC ISLAMABAD (FGPC-PGMI) (Federal Government Polyclinic (FGPC-PGMI)), Joint Executive Director
FEDERAL GOVERNMENT POLYCLINIC (PGMI) G-6/2, ISLAMABAD, Islamabad Capital Territory
Phone: +92-333-550-9275, Email: jed@fgpc.gov.pk*

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REQUEST FOR BIDS

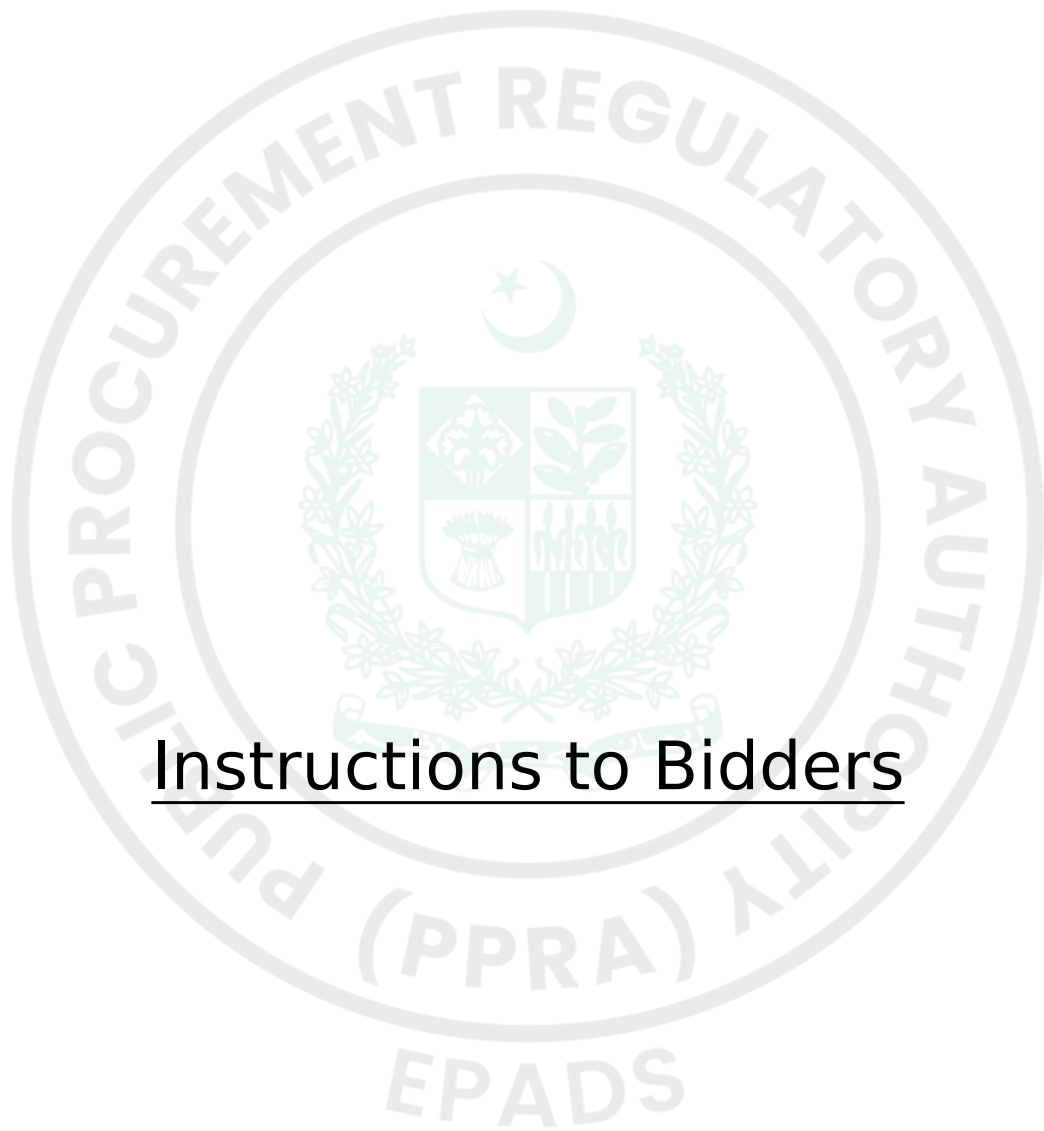
PROCUREMENT OF GOODS

1. The **FEDERAL GOVERNMENT POLYCLINIC ISLAMABAD (FGPC-PGMI)** (**Federal Government Polyclinic (FGPC-PGMI)**) has reserved Funds for the procurement planned for FY **2026-27**. The **FEDERAL GOVERNMENT POLYCLINIC ISLAMABAD (FGPC-PGMI)** (**Federal Government Polyclinic (FGPC-PGMI)**) intends to apply part of the proceeds of this Fund to cover eligible payments under the contract for the "**2nd Re-Tender for Electro-Medical Equipment/s of Cardiology Department, under PSDP Project titled 'Strengthening and Up-Gradation of Cardiology and other Departments during the FY 2025-26 & 2026-27'**" with the reference of "**P148647**"
2. The **FEDERAL GOVERNMENT POLYCLINIC ISLAMABAD (FGPC-PGMI)** (**Federal Government Polyclinic (FGPC-PGMI)**) invites sealed Bids from eligible Bidders for procurement of goods described in the bidding documents on **EPADS v2.0**.
3. **Single Stage-Two Envelope** will be used by adopting **Least Cost Based Selection (LCBS)** Technique for the subject procurement, in line with the Public Procurement Rules, 2026 and any Regulations, Regulatory Guides, Procurement Guidelines or Instructions issued by the Authority from time to time.
4. All Bids must be accompanied by a Bid Security amount described in Bid Security Section in Bidding Document in the form of **Pay Order, Call at Deposit** or all bids must be accompanied by bid securing declaration in the format specified in the Bidding documents
5. E-Bidding documents, containing detailed terms & conditions, specifications and requirements etc. are available on **e-Pak Acquisition and Disposal System (EPADS)** at **<https://epads.gov.pk/opportunities/federal/procurements/148647>** for all the interested bidders registered on **EPADS v2.0**. Bidders are required to get themselves registered on **EPADS v2.0** to participate in Bidding process.

6. The e-bids, prepared in accordance with the instructions in the e-Bidding Documents, must be submitted through **EPADS v2.0** on or before **Monday, November 2, 2026 09:30 AM**. E-bids will be opened using **EPADS v2.0** on the same day at **Monday, November 2, 2026 10:00 AM**. Manual submission of Bids shall not be entertained. Those vendors who have not yet registered on the new version of **EPADS v2.0**, may register themselves on <https://vendors.epads.gov.pk/>. A tutorial to explain the registration process is available at <https://www.youtube.com/watch?v=MNW6T38v7tc>

In terms of Rule 65 of Public Procurement Rules, 2026 Grievance Redressal Committee (GRC) is notified for the subject procurement and notification copy is available on the procuring agency's website and on Authority's website at (www.ppra.gov.pk).

FEDERAL GOVERNMENT POLYCLINIC ISLAMABAD (FGPC-PGMI) (Federal
Government Polyclinic (FGPC-PGMI)), Joint Executive Director
FEDERAL GOVERNMENT POLYCLINIC (PGMI) G-6/2, ISLAMABAD, Islamabad
Capital Territory
+92-333-550-9275
jed@fgpc.gov.pk



Instructions to Bidders

A. Introduction

1.Scope of Bids

1.1 The Procuring Agency (PA), as indicated in the **Bids Data Sheet (BDS)** invites Bids **through EPADS v2.0** for the provision of Goods as specified in the BDS and **in Section V - Evaluation Criteria, Specifications & Schedule of Requirements**. The name, identification, and number of items/deliverables are provided in the **BDS**. The successful Bidders will be expected to provide the goods within the specified period and timeline(s) as stated in the **BDS**.

2. Source of Funds

2.1 Source of funds is referred in Clause-2 I of Invitation for Bids.

3. Eligible Bidders

3.1 A Bidder may be natural person, company or firm or public or semi-public agency of Pakistan or any foreign country, or any combination of them with a formal existing agreement (on Judicial Papers) in the form of a joint venture, consortium, or association. In the case of a joint venture, consortium, or association, all members shall be jointly and severally liable for the execution of the Contract in accordance with the terms and conditions of the Contract. The joint venture, consortium, or association shall nominate a Lead Member as nominated in the BDS, who shall have the authority to conduct all business for and on behalf of any and all the members of the joint venture, consortium, or association during the Bidding process, and in case of award of contract, during the execution of the contract.

Verifiable copy of the agreement that forms a joint venture, consortium or association shall be required to be submitted as part of the Bid.

The appointment of Lead Member in the joint venture, consortium, or association shall be confirmed by submission of a valid Power of Attorney to the Procuring Agency.

Any bid submitted by the joint venture, consortium or association shall indicate the part of proposed contract to be performed by each party and each party shall be evaluated (or post qualified if required) with respect to its contribution only, and the responsibilities of each party shall not be substantially altered without prior written approval of the Procuring Agency and in line with any

instructions issued by the Authority.

(The limit on the number of members of JV or Consortium or Association may be prescribed in BDS, in accordance with the guidelines issued by the PPRA).

3.2 The invitation for Bids is open to all prospective suppliers, manufacturers, or authorized agents / dealers subject to any provisions of incorporation or licensing by the respective national incorporating agency or statutory body established for that particular trade or business. Procuring agencies shall specify the registration/licensing requirements for the foreign bidders keeping in view the requirement of that business.

3.3 A Bidder shall not have a conflict of interest. All Bidders found to have a conflict of interest shall be disqualified. A Bidder may be considered to have a conflict of interest with one or more parties in this Bidding process, if they:

1. are associated or have been associated in the past, directly or indirectly with a firm or any of its affiliates which have been engaged by the Procuring Agency to provide consulting services for the preparation of the design, specifications and other documents to be used for the procurement of the Goods to be purchased under this Invitation for Bids.
2. have controlling shareholders in common; or
3. receive or have received any direct or indirect subsidy from any of them; or
4. have the same legal representative for purposes of this Bid; or
5. have a relationship with each other, directly or through common third parties, that puts them in a position to have access to information about or influence on the Bids of another Bidder, or influence the decisions of the Procuring Agency regarding this Bidding process; or
6. Submit more than one Bid in this Bidding process.

3.4 A Bidder may be ineligible if –

1. he is declared bankrupt or, in the case of company or firm, insolvent;
2. payments in favor of the Bidder is suspended in accordance with the judgment of a court of law other than a judgment declaring bankruptcy and resulting (in accordance with the national laws) in the total or partial loss of the right to administer and dispose of its property;

3. the Bidder is convicted, by a final judgment, of any offence involving professional conduct;

4. the Bidder is blacklisted locally or by international organizations and hence debarred due to involvement in corrupt and fraudulent practices, or submission of false or misleading information or performance failure or involved in disrupting procurement process or due to breach of bid Securing Declaration.

3.5 As and when required, bidders shall provide to the Procuring Agency evidence of their eligibility, proof of compliance with the necessary legal requirements to carry out the contract effectively.

3.6 Bidders shall submit Bids relating to the nature, conditions and modalities of sub-contracting wherever the sub-contracting of any elements of the contract amounting to more than ten (10) percent of the Bid price is envisaged.

4. Eligible Goods and Related Services

4.1 All goods and related services to be supplied under the contract shall have their origin in eligible source countries, and all expenditures made under the contract will be limited to such goods and services. For purpose of this Bid, ineligible countries are the countries declared ineligible by the Federal Government.

5. One Bid per Bidder

5.1 A bidder shall submit only one Bid, in the same bidding process, either individually as a Bidder or as a member in a joint venture or any similar arrangement.

5.2 The Bidder shall not engage a subcontractor for any portion of the contract if the value of such subcontracting exceeds thirty percent (30%) of the total contract amount.

6. Cost of Bidding

6.1 Any cost incurred by the bidder relating to the preparation and submission of its Bid shall be borne by the bidder, and the Procuring Agency shall in no case be responsible or liable for those costs, regardless of the conduct or outcome of the bidding process.

B. Bidding Documents

7. Contents of Bidding Document

7.1 The Goods required, Bidding procedures, and terms and conditions of the contract are prescribed in the Bidding Documents. In addition to the Invitation for Bids, the Bidding documents which should be read in conjunction with any addenda issued in accordance with **ITB 10.1** include:

Section I -Invitation to Bids

Section II Instructions to Bidders (ITB)

Section III Bid Data Sheet (BDS)

Section IV Evaluation Criteria, Specifications, Schedule of Requirements

Section V Bid Forms

Section VI General Conditions of Contract (GCC)

Section VII Special Conditions of Contract (SCC)

Section VIII Contract Forms

Section IX Integrity Pact

Section X Declaration of Beneficial Ownership Information.

7.2 The Bidder is expected to examine all instructions, forms, terms and specifications in the Bidding documents. Failure to furnish all the information required in the Bidding documents through EPADS v2.0 will be at the Bidder's risk and may result in the rejection of its Bids.

8. Clarification of Bidding documents

8.1 A prospective Bidder requiring any clarification of the Bidding documents may notify the Procuring Agency through EPADS v2.0.

8.2 The Procuring Agency will within two (02) working days after receiving the request for clarification, respond to any request for clarification through EPADS v2.0 provided that such request is received not later than three (03) days, prior to the deadline for the submission of Bids as prescribed in **ITB 22**

8.3 The Procuring Agency's response will be forwarded to all identified Prospective Bidders through EPADS v2.0, including a description of the inquiry,

but without identifying its source.

8.4 Should the Procuring Agency deem it necessary to amend the Bidding document as a result of a clarification, it shall do so following the procedure under **ITB 9**.

8.5 If indicated **in the BDS**, the Bidder's designated representative is invited at the Bidder's cost to attend a pre-Bid meeting at the place, date and time mentioned **in the BDS**. During this pre-Bid meeting, prospective Bidders may request clarification of the schedule of requirement, the Evaluation Criteria or any other aspects of the Bidding document.

8.6 Minutes of the pre-Bid meeting, if applicable, including the text of the questions asked by Bidders, including those during the meeting (without identifying the source) and the responses given, together with any responses prepared after the meeting will be uploaded on EPADS v2.0. Any modification to the Bidding documents that may become necessary as a result of the pre-Bid meeting shall be made by the Procuring Agency exclusively through the use of an Addendum pursuant to ITB 9. Non-attendance at the pre-Bid meeting will not be a cause for disqualification of a Bidder.

9. Amendment of Bidding documents

9.1 Before the deadline for submission of Bids, the Procuring Agency for any reason, whether at its own initiative or in response to a clarification requested by a prospective Bidder or Pre-Bid meeting may modify the Bidding documents by issuing addenda through EPADS v2.0.

9.2 The Procuring Agency shall promptly publish the addendum through EPADS v2.0.

Any addendum issued including the notice of any extension of the deadline shall also be communicated through EPADS v2.0 to all the bidders who have already submitted their bids. Such bidders shall have the right to withdraw their already submitted bid and re-submit the revised bid prior to the original or extended bid submission deadline.

9.3 To give prospective Bidders reasonable time in which to take an addendum/corrigendum into account in preparing their Bids, the Procuring Agency may, at its discretion, extend the deadline for the submission of Bids through EPADS v2.0:

Provided that the Procuring Agency shall extend the deadline for submission of Bids, if such an addendum is issued within last three (03) days of the Bids submission deadline.

C. Preparation of Bids

10. Language of Bid

10.1 The Bid prepared by the bidder, as well as all correspondence and documents relating to the Bids exchanged by the Bidder and the Procuring Agency shall be written in the English language unless otherwise specified in the BDS. Supporting documents and printed literature furnished by the Bidder may be in another language provided they are accompanied by an accurate translation of the relevant pages in the English language unless otherwise specified in the **BDS**, in which case, for purposes of interpretation of the Bidder, the translation shall govern.

11. Documents and samples Constituting the Bid

11.1 The Bid prepared by the Bidder shall constitute the documents required in the **BDS**.

Details of sample(s) where applicable and requested in the BDS.

Documentary evidence established in accordance with ITB 13 that the Bidder is eligible and/or qualified for the subject bidding process;

Documentary evidence established in accordance with **ITB 03** that the Bidder has been authorized by the manufacturer to deliver the goods into Pakistan, where required and where the supplier is not the manufacturer of those goods;

Documentary evidence established in accordance with ITB 12 that the goods and related services to be supplied by the Bidder are eligible goods and services, and conform to the Bidding Documents;

Bid security or Bid Securing Declaration furnished in accordance with **ITB ---**.

12. Documents Establishing Eligibility of the Goods and Conformity to Bidding documents

12.1 To establish the conformity of the bidder to the Bidding document, the Bidder shall furnish as part of its Bids the documentary evidence that Goods

provided conform to the technical specifications and standards.

13. Documents Establishing Eligibility and Qualification of the Bidder

13.1 Pursuant to **ITB 10**, the Bidder shall furnish, as part of its Bid, all those documents establishing the Bidder's eligibility to participate in the Bidding process and/or its qualification to perform the contract if its Bid is accepted.

14. Form of Bid

14.1 The Bidder shall fill the Form of Bid furnished in the Bidding document. The Bid Form must be completed without any alterations to its format and no substitute shall be accepted.

15. Bid Prices

15.1 The Bid Prices quoted by the Bidder in the Form of Bid and in the Price Schedules shall conform to the requirements specified below or exclusively mentioned hereafter in the Bidding documents.

15.2 All items in the Schedule of Requirement must be listed and priced separately in the Price Schedule(s). If a Price Schedule shows items listed but not priced and neither explicitly denied, their prices shall be construed to be included in the prices of other items.

15.3 In the event that any item(s) in the Price Schedule are not quoted, and such omission is determined by the Procuring Agency not to materially affect the responsiveness of the Bid, the Procuring Agency may, for the sole purpose of bid evaluation and comparison, assign a price to the unquoted item(s) in accordance with the following methodology:

(a) The assigned price shall be the average of the prices quoted for the same item(s) by all substantially responsive bidders; or

(b) In cases where no comparable quotations are available, the Procuring Agency shall apply the corresponding item price from its duly approved cost estimate for evaluation purposes.

The prices so assigned shall be utilized exclusively for bid evaluation and comparison, and shall not, under any circumstances, be incorporated into or form part of the Contract Price.

15.4 The Bid price to be quoted in the Form of Bid in accordance with **ITB 13.1** shall be the total price of the Bid.

15.5 The Bidder shall indicate on the appropriate Price Schedule, the unit prices (where applicable) and total Bid price of the Goods it proposes to provide under the contract.

15.6 Prices quoted by the Bidder shall be fixed during the Bidder's performance of the contract and not subject to variation on any account. A Bid submitted with an adjustable price will be treated as non-responsive and shall be rejected.

16. Bids Currencies

16.1 Prices shall be quoted in Pakistani Rupees unless otherwise specified in the BDS in accordance with Rule 42 (2) of the Public Procurement Rules, 2026.

17. Bids Validity Period

17.1 Such extensions may be more than one time but the total period of such extension shall not exceed the original bid validity period.

17.2 Bids shall remain valid for the period specified in the **BDS** after the Bid submission deadline prescribed by the Procuring Agency. A Bid valid for a shorter period shall be rejected by the Procuring Agency as non-responsive. The period of Bid validity will be determined from the complementary Bid securing instrument, i.e. the expiry period of Bid Security or Bids Securing Declaration as the case may be.

The procuring agency shall ordinarily be under an obligation to process and evaluate the bid and to issue letter of award within the stipulated bid validity period.

17.3 Under exceptional circumstances, prior to the expiration of the initial Bid validity period, the Procuring Agency may request the Bidders' consent to an extension of the period of validity of their Bids through EPADS v2.0. The Bid Security provided under **ITB 17** shall also be suitably extended. A Bidder may refuse the request without forfeiting its Bid security or causing to be executed its Bid Securing Declaration. A Bidder agreeing to the request will not be required nor permitted to modify its Bid, but will be required to extend the validity of its Bid Security or Bid Securing Declaration for the period of the extension.

18. Bid Security or Bid Securing Declaration

18.1 The Bidder shall furnish as part of its Bid, a Bid Security in accordance with Rule 36 of the Public Procurement Rules, 2026.

18.2 The original Bid Security shall be enclosed within the sealed envelope and to be submitted physically before closing time for submission of bids. Whereas, scanned copy of bid security shall be uploaded electronically through EPADS v2.0 before closing hours for submission of bids.

18.3 The Bidder who failed to submit the original Bids security before the submission deadline shall be disqualified straightaway.

18.4 The Bid Security or Bid Securing Declaration is required to protect the Procuring Agency against the risk of Bidder's conduct which would warrant the security's forfeiture, pursuant to **ITB 18.7**.

18.5 The Bid Security shall be denominated in the local currency or in another freely convertible currency, and it shall be in the form specified in the BDS which shall be in any of the following:

- a) a bank guarantee, a bank draft, an irrevocable letter of credit issued by a Scheduled bank in the form provided in the Bidding Documents or another form acceptable to the Procuring Agency;
- b) a cashier's or certified cheque; or
- c) another security if indicated in the BDS.

The Bid Security shall be valid for twenty-eight (28) days beyond the end of the validity of the Bid. This shall also apply if the period for Bid Validity is extended. In either case, the form must include the complete name of the Bidder.

18.6 The Bid Security shall be payable promptly upon written demand by the Procuring Agency in case any of the conditions listed in **ITB 18.7** are invoked.

18.7 Unsuccessful Bidders' Bid Security will be discharged or returned as promptly as possible, however in no case later than thirty (30) days after the expiration of the period of Bid Validity prescribed by the procuring agency pursuant to **ITB 17**. The Procuring Agency shall make no claim to the amount of the Bid Security, and shall promptly return the Bid Security documents, after whichever of the following that occurs earliest:

- a. the expiry of the Bid Security;

- b. the entry into force of a procurement contract and the provision of a Performance Guarantee, for the performance of the contract if such a guarantee, is required by the Bid documents;
- c. the rejection by the Procuring Agency of all Bids;
- d. cancellation of procurement proceedings.
- e. the withdrawal of the Bid prior to the deadline for the submission of Bids, unless the Bid document stipulates that no such withdrawal is permitted.

18.8 The successful Bidder's Bid Security will be discharged upon the Bidder signing the contract, or furnishing the Performance Guarantee.

18.9 The Bid Security may be forfeited or the Bid Securing Declaration executed:

- a. if a Bidder:
- b. withdraws its Bid during the period of Bid Validity as specified by the Procuring Agency, and referred by the Bidder on the Form of Bids except as provided for in **ITB 16.2**; or
- c. does not accept the correction of errors; or
- d. in the case of a successful Bidder, if the Bidder fails:
- e. to sign the contract; or
- f. to furnish Performance Guarantee.

19. Withdrawal, Substitution, and Modification of Bid

19.1 Before Bid submission deadline, any Bidder may withdraw, substitute, or modify its Bid after it has been submitted through EPADS v2.0.

20. Format and Signing of Bid

20.1 The Bidder shall prepare and submit Bids with due diligence after carefully reading all the terms and condition before bid submission deadline through EPADS v2.0.

D. Submission of Bids

21. Submission of Bids through EPADS v2.0 before bid submission deadline.

21.1 The Technical and Financial Bids, shall be submitted electronically through EPADS, before bid submission deadline. The bid submission option shall be automatically disabled once the deadline is over.

22. Deadline for Submission of Bids

22.1 Bids shall be received by the Procuring Agency through EPADS v2.0 before bid submission deadline.

22.2 The Procuring Agency may, under exceptional circumstances, extend the deadline for the submission of Bids, after recording reasons in writing and in an equal opportunity manner.

In such case, all rights and obligations of the Procuring Agency and the Bidders that were previously governed by the original deadline shall thereafter be subject to the revised deadline.

E. Opening and Evaluation of Bids

23. Opening of Bids

23.1 The respective Bid Evaluation Committee of the Procuring Agency shall open all Bids through the EPADS v2.0, on the date and time specified in the Bid Data Sheet (BDS).

23.2 The respective Bid Evaluation Committee **shall generate minutes through EPADS v2.0 containing brief details of bid opening process.** The record of the Bid opening shall include, as a minimum: the name of the Bidder, the Bid price if applicable, and the presence or absence of a Bid Security or Bid Securing Declaration.

23.3 The procuring agency shall live broadcast the opening of bids on national media or on their website or digital channels, if the volume of procurement exceeds five hundred million rupees in case of goods and services and one thousand million rupees in case of works.

23.4 In case the date of opening of bid has been declared as public holiday or the procuring agency fails to open bid due to any EPADS v2.0 related issues, the submission and opening of bids shall be shifted to the next working day at the same time.

23.5 In case of Single Stage One Envelope Procedure, the Bidders names, the Bid prices, the total amount of each Bid and, the presence or absence of Bid Security, Bid Securing Declaration and such other details as the Procuring Agency may consider appropriate, will be announced by the Bid respective Evaluation Committee.

23.6 No bid will be rejected at the time of bid opening except for bids whose original bid security has not been provided to the procuring agency before bid submission deadline.

24. Clarification of Bids

24.1 To assist in the examination, evaluation and comparison of Bids of the Bidders, the respective Bid Evaluation Committee may, ask any Bidder for a clarification of its Bid including breakdown of prices.

24.2 The request for clarification and the response shall be sought through EPADS v2.0 before **three days prior to the deadline for submission of bids.** No change in the prices or substance of the Bids shall be sought, offered, or permitted.

24.3 The alteration or modification in the bids which in any way affect the following parameters will be considered as a change in the substance of a Bid:

1. evaluation & qualification criteria;
2. required scope of work or specifications;
3. all securities requirements;
4. tax requirements;
5. terms and conditions of Bidding documents.
6. change in the ranking of the Bidder

24.4 From the time of bid(s) opening to the time of Contract award if any Bidder wishes to contact the Procuring Agency on any matter related to the Bids it should do so through **EPADS v2.0**.

25. Preliminary Examination of Bids

25.1 Prior to the detailed evaluation of Bids, the respective Bid Evaluation Committee will determine whether each Bid:

1. meets the eligibility criteria defined in **ITB 3**;
2. has been prepared as per the format and contents defined by the Procuring Agency in the Bidding documents;
3. is accompanied by the required securities; and
4. is substantially responsive to the requirements of the Bidding documents.

25.2 The respective Bid Evaluation Committee's determination of a Bid's responsiveness will be based on the contents of the Bid itself.

25.3 A substantially responsive Bid is one which conforms to all the terms, conditions, and specifications of the Bidding documents, without material deviation or reservation. A material deviation or reservation is one that: -

1. affects in any substantial way the scope, quality, or performance of the Goods;
2. limits in any substantial way, inconsistent with the Bidding documents, the Procuring Agency's rights or the Bidders obligations under the Contract; or
3. if rectified, would affect unfairly the competitive position of other Bidders presenting substantially responsive Bids.

25.4 If a Bid is not substantially responsive, it will be rejected by the respective Bid Evaluation Committee and may not subsequently be evaluated for complete technical responsiveness.

26. Examination of Terms and Conditions; Technical Evaluation

26.1 The respective Bid Evaluation Committee shall examine the Bids to confirm that all terms and conditions specified in the **GCC** and the **SCC** have been accepted by the Bidder without any material deviation or reservation.

26.2 The respective Bid Evaluation Committee shall evaluate the technical aspects of the Bids submitted, to confirm that all requirements specified in Schedule of Requirements and Technical Specifications of the Bidding documents have been met without material deviation or reservation.

26.3 If after the examination of the terms and conditions and the technical evaluation, the respective Bid Evaluation Committee determines that the Bid is not substantially responsive in accordance with **ITB**, it shall reject the Bid.

27. Correction of Errors

27.1 Bids determined to be substantially responsive will be checked for any arithmetic errors. Errors will be corrected as follows: -

1. if there is a discrepancy between unit price and the total price that is obtained by multiplying the unit price and quantity, the unit price shall prevail, and the total price shall be corrected, unless in the opinion of the respective Evaluation Committee there is an obvious misplacement of the decimal point in the unit price, in which the total price as quoted shall govern and the unit price shall be corrected;
2. if there is an error in a total corresponding to the addition or subtraction of sub-totals, the sub-totals shall prevail and the total shall be corrected; and
3. where there is a discrepancy between the amounts in figures and in words, the amount in words will govern.
4. Where there is discrepancy between grand total of price schedule and amount mentioned on the Form of Bid, the amount referred in Price Schedule shall be treated as correct subject to elimination of other errors.

27.2 The amount stated in the Bid will be adjusted by the respective Bid Evaluation Committee in accordance with the above procedure for the correction of errors and, with the concurrence of the Bidder, shall be considered as binding upon the Bidder. If the Bidder does not accept the corrected amount, its Bid will then be rejected, and the Bid Security may be forfeited or the Bids Securing Declaration may be executed.

28. Conversion to Single Currency

28.1 For the purposes of comparison of bids quoted in different currencies, the price shall be converted into a single currency in accordance with Rule 42(2) of Public Procurement Rules, 2026.

29. Evaluation of Bids

29.1 The Bids, quotations, or proposals shall be evaluated by the respective evaluation committees as per evaluation criteria prescribed in the Bidding Documents in accordance with Rule 11, 12 and 13 of the Public Procurement Rules, 2026.

29.2 While evaluating the bids, the procuring agency shall consider the following:

- i. Bids will be evaluated on lump sum bid basis or items wise or Lot wise (contracts), as specified in the BDS; and the Bid Price;
- ii. correction of arithmetic errors in accordance with ITB 27;

30. Domestic Preference

30.1 The respective Bid Evaluation Committee shall evaluate and compare bids, allow for preference to domestic bidders, while competing with the international bidders in accordance with the policies of the Federal Government.

The percentage of preference, to be accorded shall be clearly mentioned in the bidding documents under the bid evaluation criteria.

31. Determination of Successful Bid

31.1 Selection technique will be adopted for determining the Successful Bid in accordance with the criteria referred in the BDS or prescribed in the separate section titled as Evaluation Criteria.

31.2 Least Cost Based Selection (LCBS)

After meeting the requirements of eligibility, qualification and substantial responsiveness, the bid in compliance with all the mandatory (technical) specifications/requirements and/or requisite quality threshold (if any), and having lowest evaluated cost (or financial proposal) shall be considered Successful Bid.

31.3 Quality and Cost Based Selection (QCBS)

In such combination, there shall be some specific weightage of both the technical features and financial aspects of the proposal. The financial marks shall be awarded on the basis of inverse proportion calculations. The successful bid shall be declared, on the basis of combined evaluation.

31.4 Quality Based Selection (QBS)

After meeting the requirements of eligibility, qualification and substantial responsiveness the bid in compliance with all the mandatory (technical) specifications/requirements and attaining highest marks in the Technical Evaluation considering all other qualitative and/or quantitative parameters (or point rated criteria) for technical proposal(s) such as working methodology, implementation plan, resource allocation, additional functionalities, risk

management approach, knowledge transfer techniques, post implementation methodology etc. shall be treated as highest ranked bid. Later on, the financial proposal of highest ranked bidder shall be opened, however, in case of failure to proceed further with such a bidder, the procuring agency may resort to second highest bidder and so on.

31.5 The Procuring Agency may adopt the Quality & Cost Based Selection Technique due to the following two reasons:

1. Where the Procuring Agency knows about the main features, usage and output of the products; however not clear about the complete features, technical specifications and functionalities of the goods to be procured and requires the bidders to submit their proposals defining those features, specifications and functionalities; or
2. Where the Procuring Agency, in addition to the mandatory requirements and mandatory technical specifications, requires parameters specified in Evaluation Criteria to be evaluated while determining the quality of the goods.

31.6 In such cases, the Procuring Agency may allocate certain weightage to these factors as a part of Evaluation Criteria, and may determine the ranking of the bidders on the basis of combined evaluation in accordance with provisions of Rule 2(1)(xxviii) of the Public Procurement Rules, 2026.

32. Abnormally Low Financial Bids

32.1 An abnormally low bid is a bid that, in light of the Procuring Agency's cost estimate, market conditions, or other received bids, appears to be unrealistically low and may not be sustainable for successful contract execution.

32.2 Where the Bid price is considered to be abnormally low, the Procuring Agency shall perform price analysis either during determination of Successful Bids or as a part of the post-qualification process.

32.3 The Procuring Agency may reject an Abnormally low financial bid in accordance with Rule 43 of the Public Procurement Rules, 2026.

32.4 In order to identify the Abnormally Low Bids (ALB) following approaches can be considered to minimize the scope of subjectivity:

1. Comparing the Bid price with the cost estimate;

2. Comparing the Bid price with the Bid(s) offered by other Bidders submitting substantially responsive Bids; and

3. Comparing the Bids price with prices paid in similar contracts in the recent past either government- or development partner-funded.

32.5 The respective Bid Evaluation Committee will determine to its satisfaction whether the Bidder that is selected as having submitted the successful bid is qualified to perform the contract satisfactorily.

32.6 The determination will take into account the Bidder's financial, technical, and production capabilities. It will be based upon an examination of the documentary evidence of the Bidder's qualifications submitted by the Bidder, as well as such other information as the Procuring Agency deems necessary and appropriate. Factors not included in these Bidding documents shall not be used in the evaluation of the Bidders' qualifications.

32.7 Respective Bid Evaluation Committee may seek "Certificate for Independent Price Determination" from the Bidder and the results of reference checks may be used in determining an award of contract.

Explanation: The Certificate shall be furnished by the Bidder. The Bidder shall certify that the price is determined keeping in view of all the essential aspects such as raw material, its processing, value addition, optimization of resources due to economy of scale, transportation, insurance and margin of profit etc.

32.8 An affirmative determination will be a prerequisite for award of the contract to the Bidder. A negative determination will result in rejection of the Bidder's Bid, in which event the respective Bid Evaluation Committee will proceed to the next ranked Bidder to make a similar determination of that Bidder's capabilities to perform satisfactorily.

F. Award of Contract

33. Criteria of Award

33.1 The Procuring Agency will award the Contract to the Bidder whose Bids has been determined to be substantially responsive to the Bidding documents and who has been declared as Successful Bidder.

34. Negotiations

34.1 In accordance with Rule 52A of Public Procurement Rules, 2026, without changing the cost and scope of work or services, the procuring agency may

negotiate with the successful bidder (with a view to streamline the work or task execution, at the time of contract finalization) on methodology, work plan, staffing and special conditions of the contract.

There shall be no limitation on the successful bidder offering voluntary reduction in bid price without affecting the quality or scope of the procurement.

35. Procuring Agency's Right to reject all bids

35.1 The Procuring Agency reserves the right to reject all bids or proposals at any time prior to the issuance of the Letter of Award, without incurring any liability, in accordance with Rule 45 of the Public Procurement Rules, 2026.

36. Cancellation of Procurement

36.1 The procuring agency reserves the right to cancel the procurement proceedings in accordance with Rule 46 of the Public Procurement Rules, 2026.

37. Procuring Agency's Right to Vary Quantities at the Time of Award

37.1 The Procuring Agency reserves the right at the time of contract award to increase or decrease the **quantity of** Goods originally specified in these Bidding documents provided this does not exceed **by** 15%, without any change in unit price or other terms and conditions of the Bids and Bidding documents.

38. Notification of Award

38.1 Prior to the award of contract, the procuring agency shall announce and publish the result of bid evaluation on EPADS v2.0 in accordance with Rule 48 of the Public Procurement Rules, 2026.

The Bidder whose Bids has been accepted will be notified of the award by the Procuring Agency prior to expiration of the Bids/Bid Validity period. The Letter of Award will state the sum that the Procuring Agency will pay the successful Bidder in consideration for the delivery of Goods as prescribed by the Contract (hereinafter and in the Contract called the "Contract Price).

38.2 The Letter of award will constitute the formation of the Contract, subject to the Bidder furnishing the Performance Guarantee and signing of the contract.

39. Signing of Contract

39.1 Promptly after issuance of Letter of award, Procuring Agency shall send the successful Bidder the draft Contract, incorporating all terms and conditions as agreed by the parties to the contract.

39.2 Immediately after the Redressal of grievance by the GRC (if any), and **after fulfillment of all conditions precedent** of the Contract Form, the successful Bidder and the Procuring Agency shall sign the Contract.

40. Corrupt & Fraudulent Practices

40.1 Procuring Agencies (including beneficiaries of Government funded projects and procurement) as well as Bidders/Contractors under Government financed contracts, observe the highest standard of ethics during the procurement and execution of such contracts, and will avoid engaging in any corrupt and fraudulent practices.

F. Grievance Redressal & Complaint Review Mechanism

41. Constitution of Grievance Redressal Committee

41.1 The Grievance Redressal Committee shall address the grievance, if any submitted by any party, including the bidder, in accordance with Rule 65 of the Public Procurement Rules, 2026 read with Redressal of Grievances Regulations, 2021.

In case if any party or the bidder is not satisfied with the decision of the GRC or if it fails to decide within ten days, the bidder or the party may file an appeal before the Appellate Committee of the Authority in accordance with Rule 65(7) of the Public Procurement Rules, 2026 to be read with Redressal of Grievances Regulations, 2021.

G. Mechanism of Blacklisting

42. Mechanism of Blacklisting

42.1 The Procuring Agency shall initiate blacklisting proceedings against any bidder, supplier, or contractor in accordance with the Rule 25 of the Public Procurement Rules, 2026.

42.2 The blacklisted/debarred bidder may file the review petition before the Authority in accordance with Rule 25 (6) of the Public Procurement Rules, 2026 to be read with Procedure of filing and disposal of Review Petitions Regulations, 2021.

43. Contract Management

43.1 For each procurement, the Procuring Agency shall designate a Contract Manager, who shall be duly responsible for the efficient management and administration of the contract in strict compliance with the provisions set forth under Rules 58, 59, and 60 of the Public Procurement Rules, 2026.

44. Inspection of goods and evaluation of services.

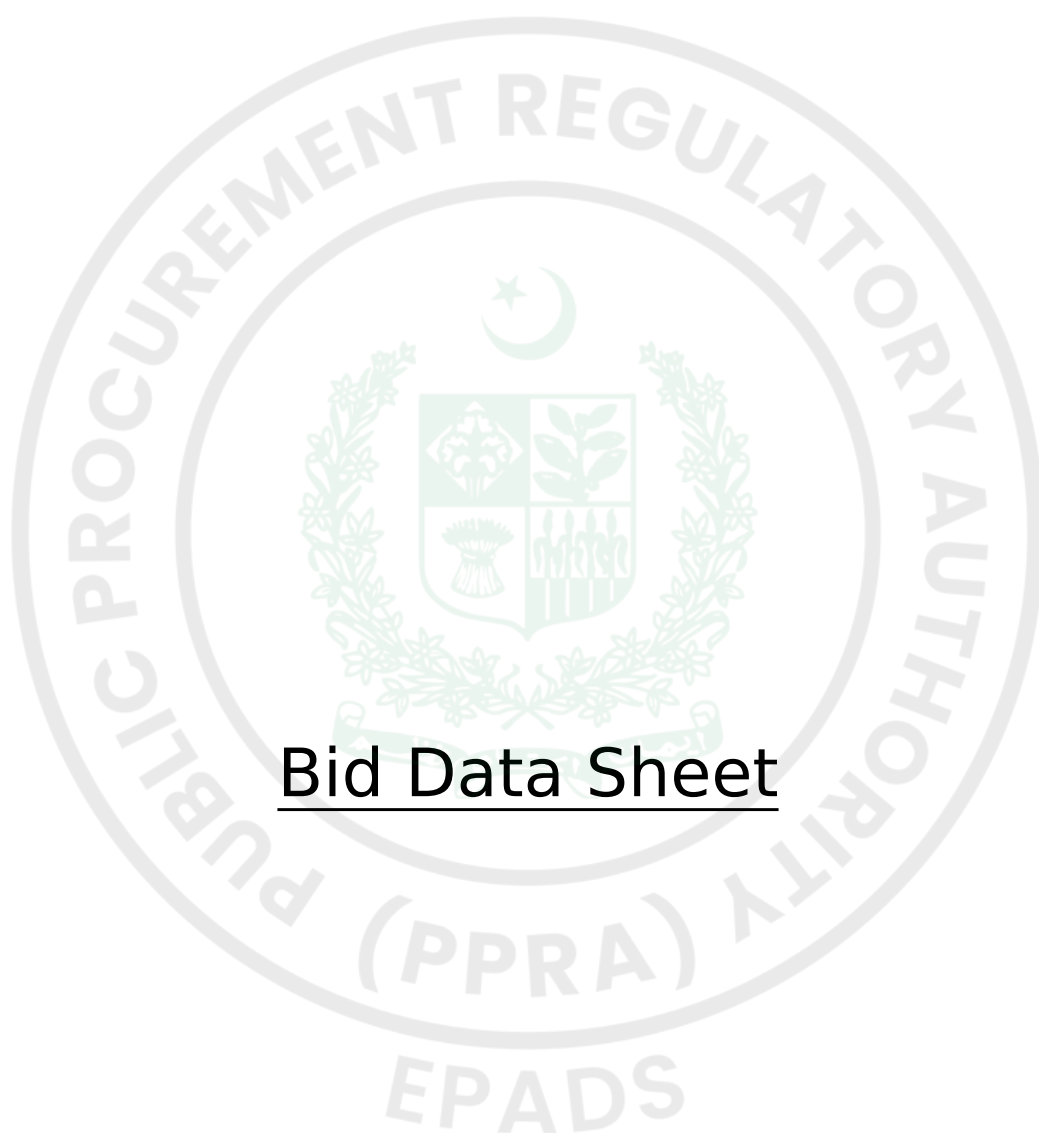
44.1 The Inspection Committee or the Firm shall have the right to conduct inspections of the goods in accordance with Rule 61 of the Public Procurement Rules, 2026.

45. Socio-economic development.

45.1 In accordance with Rule 63 of the Public Procurement Rules, 2026, preference may be granted to small and medium enterprises and marginalized groups in accordance with the policies notified by the Federal Government.

46. Environmental objectives

46.1 As per Rule 64 of the Public Procurement Rules, 2026, the procuring agency may seek to procure goods with a reduced environmental impact throughout their life cycle when compared to goods services with the same primary function that may otherwise be procured.



Bid Data Sheet

Bids Data Sheet (BDS)

The following specific data for the procurement of Goods to be procured shall complement, supplement, or amend the provisions in the Instructions to Bidders (ITB). Whenever there is a conflict, the provisions herein shall prevail over those in ITB.

BDS Clause Number

ITB Number

Amendments of, and Supplements to, Clauses in the Instruction to Bidders

A. Introduction

BDS Clause Number 1

ITB Number 1.1

Name of Procuring Agency: **FEDERAL GOVERNMENT POLYCLINIC ISLAMABAD (FGPC-PGMI) (Federal Government Polyclinic (FGPC-PGMI))**

The subject of procurement is: **2nd Re-Tender for Electro-Medical Equipment/s of Cardiology Department, under PSDP Project titled 'Strengthening and Up-Gradation of Cardiology and other Departments during the FY 2025-26 & 2026-27**

Period for Provision of Goods:

Extension Period (if applicable):

Expected commencement date: **Wednesday, March 31, 2027**

BDS Clause Number 2

ITB Number 2.1

Source of funds [Enter source of funds]

Financial year for the operations of the Procuring Agency: **2026-27**

Name and identification number of the Contract: **P148647**

Identification Number:

BDS Clause Number 3

ITB Clause Number 3.1

Eligible Bidder [Enter eligibility of bidder]

JV/Consortium or Association Allowed: **No**

Number of JV/Consortium Members: **Nil**

see section of eligibility criteria.

B. Bidding Documents

BDS Clause Number 4

ITB Clause Number 4.1

Eligible goods and related services [Enter requirements for eligible goods and related services, if any]

ITB Number 8.1

The Bidders may seek clarifications through **EPADS v2.0** : Clarification Date: Friday, October 30, 2026

Pre-Bid Meeting: Thursday, October 22, 2026 09:30 AM

Venue: FEDERAL GOVERNMENT POLYCLINIC (PGMI) G-6/2, ISLAMABAD

ITB Number 8.2

The Bidders may seek clarifications through **EPADS v2.0** : Clarification Date: Friday, October 30, 2026

Pre-Bid Meeting: Thursday, October 22, 2026 09:30 AM

Venue: FEDERAL GOVERNMENT POLYCLINIC (PGMI) G-6/2, ISLAMABAD

ITB Number 8.3

The Bidders may seek clarifications through **EPADS v2.0** : Clarification Date: Friday, October 30, 2026

Pre-Bid Meeting: Thursday, October 22, 2026 09:30 AM

Venue: FEDERAL GOVERNMENT POLYCLINIC (PGMI) G-6/2, ISLAMABAD

ITB Number 8.5

Pre-bid meeting will be/will not be held [Tick the appropriate] If yes write down the time, and date of the pre-Bid meeting:

[The meeting should take place not later than one weeks before the deadline for Bid submission.]

C. Preparation of Bids

BDS Clause Number 5

ITB Number 10.1

The Language of all correspondences and documents related to the Bids shall be in: **English**

List of documents required along with the bid:

1. Technical brochures/Literature brand country of origin of quoted item/material must be attached with the bid where applicable
2. Quality Certification as per specification of the each item should be attached. In case of local instrument, CE or ISO13485
3. Under taking on company/bidder letterhead that bidder will provide sample and arrange demonstration of the quoted equipment if needed by the committee
4. The firms must provide a letter of Warranty including all parts of complete Angiography System machine for five years from the manufacturer starting from the date of installation. However, firm will quote further 10 years post warranty should be quoted which will be applicable after expiry of standard warranty period. The local vendor's warranty will not be acceptable

BDS Clause Number 6

ITB Number 11.1

Documents required / not required with bid submission [enter list of documents] Samples required / not required with the bid [Enter the names and number of samples]

Items/Lots and there related documents:

See section items and Lots

BDS Clause Number 7

ITB Number 12.1

Items / Lots Specifications:

see section of items specifications.

BDS Clause Number 8

ITB Number 15.6

The price shall be **Fixed**.

BDS Clause Number 9

ITB Number 16.1

Currency of the Bids shall be : **PKR**

BDS Clause Number 10

ITB Number 17.1

The Bids/Bid Validity period shall be: **120 Days**

BDS Clause Number 11

ITB Number 18.1

The amount of Bid Security shall be as defined in Bid Security Section for items and lots given in **BDS 6**

Instrument of Bid Security shall be in the Name of : **FEDERAL GOVERNMENT POLYCLINIC ISLAMABAD (FGPC-PGMI)**

The Bid Security shall be in the form of: **Pay Order, Call at Deposit**

ITB Number 18.2

The address for submission of original bid security is [Insert the address for bid security submission]

D. Submission of Bids

BDS Clause Number 12

ITB Number 20.1

Bid shall be submitted online on EPADS v2.0 whereas hard copy of the bid security should be submitted to the following;

FEDERAL GOVERNMENT POLYCLINIC (PGMI) G-6/2, ISLAMABAD, Islamabad Capital Territory before bid submission deadline.

Bids that are not submitted on EPADS v2.0 shall be disqualified.

The deadline for Bids submission is: **Monday, November 2, 2026 09:30 AM**

Day : **Monday**

Date: **Monday, November 2, 2026**

Time : **10:00 AM**

E. Opening and Evaluation of Bids

BDS Clause Number 13

ITB Number 23.1

The Bids opening shall take place on **EPADS v2.0**.

Day : **Monday**

Date: **Monday, November 2, 2026**

Time : **10:00 AM**

ITB Number 29.2

The procuring agency shall consider the following:

lump sum bid basis; or

items wise; or

Lot wise (contracts),

ITB Number 30.1

BDS Clause Number 14

ITB Number 31.1

Domestic Preference [applicable / not applicable]

ITB Number 31.2

Selection technique adopted will be:

The Procuring Agency shall use any of the following evaluation techniques for determination of successful bid.

Least Cost Base Selection (LCBS)

Quality and Cost Based Selection (QCBS)

Quality Based Selection (QBS)

[Insert the Evaluation Technique]

Selection technique adopted will be: **Least Cost Based Selection (LCBS)**
see Evaluation Criteria

F. Review of Procurement Decisions

BDS Clause Number 15

ITB Number 41.1

Grievance against this procurement shall be submitted online on EPADS v2.0.

Arbitrator shall be appointed by mutual consent of the both parties.

ITB Number 42.2

The Address of PPRA Grievance Redressal Appellate Committee to submit a copy of grievance:

Grievance Redressal Appellate Committee,
Public Procurement Regulatory Authority
1st Floor, G-5/2, Islamabad, Pakistan
Tel: +92-51-9202254



Eligibility Criteria

Bidder's Type	Required Registration
Individual / Individual Consultant	NADRA CITIZENSHIP (CNIC/NICOP)
Sole Proprietorship	FBR (NTN)
Partnership Firm	FBR (GSTN)
Company (Private Limited)	DRAP
Company (Public Limited)	
Company (Holding Company)	
Company (Limited by Guarantee)	

Eligibility Criteria	Document
Legal name, registered address, ownership status, telephone, email and names of directors, partners or proprietor. Attach SECP (if applicable) /registration of firms or other applicable registration.	Yes
Name, Designation & specimen signature of owner/authorized person of the firm along with CNIC copies.	Yes
Authorization from Manufacturer or through its Sole Agent. In case of Distributor/sole agent the bidder must provide valid sole authorization, all quoted product/s issued by the manufacturer or its exclusive sole agent. If manufacturer is directly applying, he will attach proof of its manufacturing. If the bidder participates as general order supplier for general items, then proof of registration as general order supplier should be attached.	Yes

After-Sales services Infrastructure The bidder must attach undertaking on legally Stamp paper / E-Stamp attested by Notary public or Oath Commissioner regarding After Sale services Infrastructure for maintenance/service during the warranty period & or as per specifications/as appropriate for each item	Yes
Declaration of Main equipments and its Accessories The bidder must clearly declare on legally Stamp paper / E-Stamp duly attested by Notary public or Oath Commissioner the make, model, and country of origin and manufacturing of all quoted medical equipments alongwith its accessories. (If the origin of accessories different from the main equipment, then should be mentioned separately)	Yes
Firm must have at least one (01) PEC Registered biomedical Engineer with three (03) years' relevant experience for Bio/electro-medical equipments (attached PEC registration/ Valid PEC card/ online proof of registration)	Yes
Valid DRAP establishment license for the manufacturer or importer or Distributor. DRAP Valid registration, enlistment or market authorization for each quoted medical device, as applicable. Where exemption allowed by DRAP, documentary evidence of exemption shall be provided	Yes
The bidder will provide account maintenance certificate along with bank statement of the concerned account registered with FBR, for the last three fiscal years duly verified.	Yes
The bidder will provide Income Tax/Sale Tax Return statement issued by FBR for the last three (03) financial year.	Yes
The bidder/vendor should have minimum 03 years' experience for supply of Bio/electro medical equipment to public sector hospitals/Tertiary care Hospitals independently or through International Agencies and NGO's (attach documentary proof).	Yes
The bidder must submit an affidavit on legally notarized stamp paper/ E-stamp duly attested by Notary public or Oath Commissioner declaring that the firm has not been blacklisted/Debarred by any Federal or Provincial Government department, or any public sector organization within Pakistan.	Yes

• The bidder must attach undertaking on legally notarized stamp paper/ E-stamp duly attested by Notary public or Oath Commissioner that it has no litigation(s) or arbitrary cases against the bidder by any department, is not insolvent, in receivership, bankrupt or being penalized and its activities or affairs are not suspended or being administered under any Act, by a court or by a judicial officer.	Yes
• The bidder must attach undertaking on legally notarized stamp paper/ E-stamp duly attested by Notary public or Oath Commissioner that none of its quoted product/items supplied in public sector Institutions has been declared spurious, adulterated & misbranded during the last three(03 years) till the closing date of this tender.	Yes
The bidder must attach undertaking on legally notarized stamp paper/ E-stamp duly attested by Notary public or Oath Commissioner that item/s for which the firm intends to apply shall be made freely available for delivering the supplies and will be supplied at FGPC by the bidder at its own expense. All freight/damages including installation will be the responsibility of the vendor.	Yes
The bidder must attach undertaking on legally Stamp paper / E-Stamp duly attested by Notary public or Oath Commissioner that the quoted item/medical equipment's/Instruments should be genuine /original parts manufactured and warrantied as per bidding documents.	Yes
The bidder must attach undertaking on legally Stamp paper / E-Stamp duly attested by Notary public or Oath Commissioner that all documentation submitted with the bid is valid, authentic and genuine. No facts have been hidden and no forgery/false declaration has been made. If any such discrepancy is found at any stage, the bidder will be fully responsible for such miscommunication/ concealment of facts and will be liable for disciplinary action under PPRA Rules and tender TORs.	Yes
The bidder must attach undertaking on legally Stamp paper / E-Stamp duly attested by Notary public or Oath Commissioner that the offered discount in the tender is not less than the offered discount in any other government hospital on same TOR's & same fiscal year. In case of any discrepancy found at any stage, the vendor will be bound to refund the excess amount through challan in government treasury or excess amount will be deducted from the outstanding bills/CDR.	Yes

The bidder will attach an undertaking on legally Stamp paper / E-Stamp duly attested by Notary public or Oath Commissioner regarding acceptance of tender documents including terms & conditions and specifications.

Yes



Evaluation Criteria

Eligible bidder(s) with substantially responsive bid(s) offering **Least Cost Based Selection (LCBS)** shall be consider for the award of contract(s).

Least Cost Based Selection (LCBS)

Weightage

Technical Evaluation %	
100	
Technical Marks	100
Passing Marks	70
Technical Evaluation Criteria	

PEC Registered Engineers (Quantitative) (Doc Required) (Documentary evidence of employment & experience should be attached) (Quantitative)(Doc Required) 03 Biomedical Registered Engineers with 03 years relevant experience & valid PEC registration/Card/Online verified documents (on the pay roll of firm for the last six month) (15) 02 Biomedical Registered Engineers with 03 years' experience & valid PEC registration/Card/Online verified documents (on the pay roll of firm for the last six month) (10) 01 Biomedical Registered Engineers with 03 years' experience & valid PEC registration/Card/Online verified documents (on the pay roll of firm for the last six month) (5)	15
Availability of proper Office & workshop with proper tools/technical staff for service/maintenance/repair of relevant medical equipment's (Attached evidence of ownership/ lease/ authorization address, dated inventory, serial numbers tools/equipment, attached). A valid calibration certificate/workshop authorization, list of available tools and equipment, shall also be provided on duly attested legal stamp paper/e-stamp, duly attested by an Oath Commissioner or Notary Public) (Quantitative)(Doc Required) Evidence of established office and workshop within twin cities i.e Islamabad/ Rawalpindi plus in other cities of Pakistan (15) Only Islamabad and Rawalpindi (10) Other than Islamabad and Rawalpindi (5)	15

Number of contracts/installation of quoted / relevant Electro medical/bio-medical equipment in public sector/ Tertiary health care Hospital during last 5 years. Each contract must involve supply, installation and commissioning of the same category or a pre-defined comparable category of biomedical/electro medical equipment. (Evidence: purchase order or contract plus delivery, installation, commissioning or acceptance record (Quantitative))(Doc Required) 05 or above (20) 4 (16) 3 (12) 2 (8) 1 (4)	20
Vendor Past Performance : Number of Satisfactory report/certificate issued by the public sector/tertiary care hospitals regarding provision of performance report regarding services/performance of quoted equipment. The certificate should be verified by the administrative authority of the hospital. (Quantitative)(Doc Required) 05 or above (20) 4 (16) 3 (12) 2 (8) 1 (4)	20

Financial Capability ; Assessed based on annual business turnover (supported by financial statements for the last fiscal year): (Attach documentary evidence of FBR Tax return statement/ concerned bank statement/ certificate) (Quantitative) (Doc Required) >200 million (20) >100 million to 200 :million (15) >50 million to 100 million (10) 10 million to 50 million (5)	20
The bidder is registered in FBR/EPADS as Manufacturer or sole agent or Authorized distributor or general order supplier (Quantitative)(Doc Required) Manufacturer or sole agent (10) Authorized distributor or general order supplier (5)	10

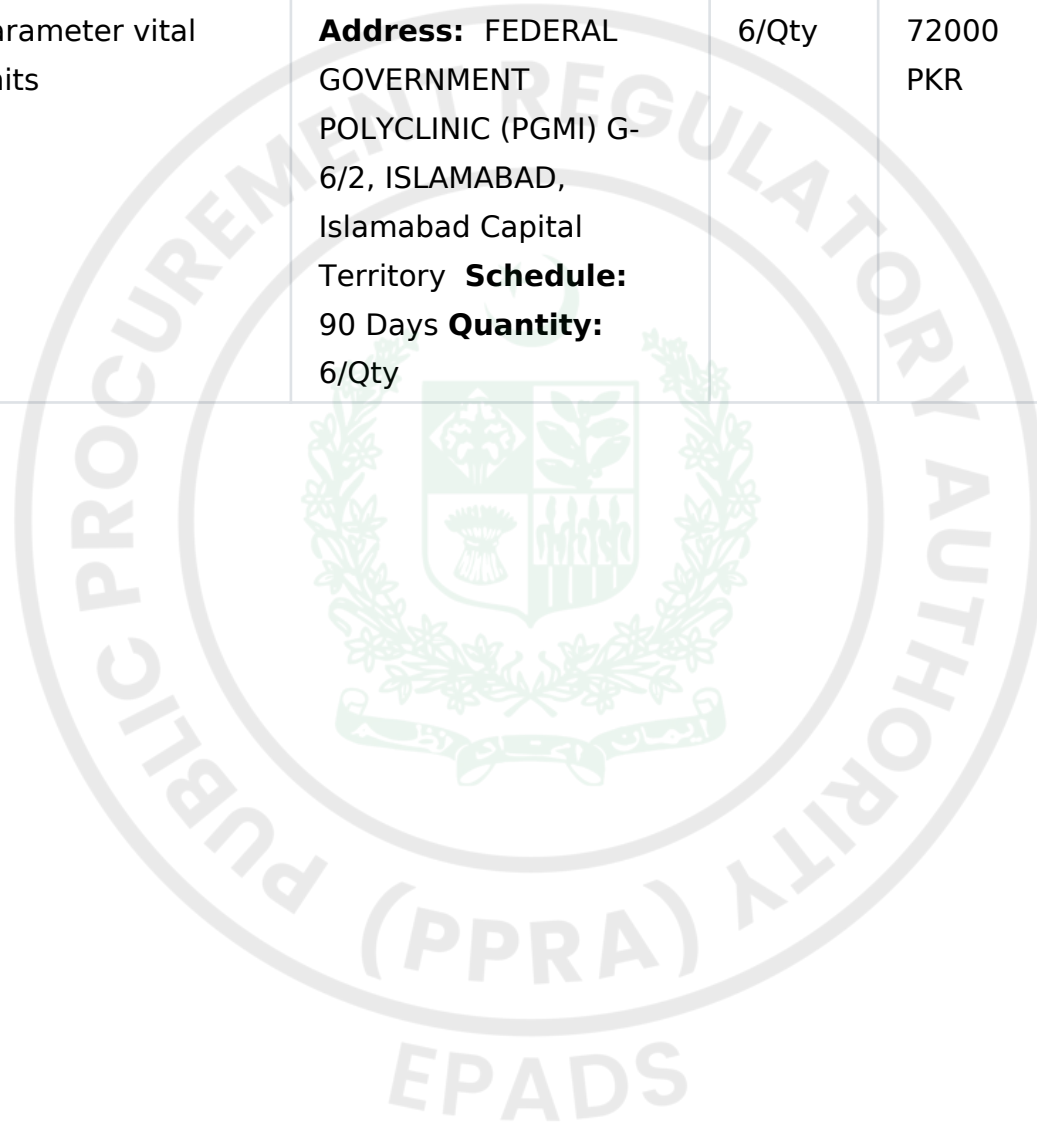
Items/Lots

Items Without Lots :

Item	UNSPSC	Delivery Schedule	Quantity	Bid Security	Manufacturer / Dealer Authorization	Warranty
Angiography System (Cath Lab)	Cardiac catheterization lab equipment	Address: FEDERAL GOVERNMENT POLYCLINIC (PGMI) G-6/2, ISLAMABAD, Islamabad Capital Territory Schedule: 120 Days Quantity: 1/Qty	1/Qty	7049000 PKR	Any	5 Years
Defibrillator machines	Mobile medical services automated external defibrillators AED or hard paddles	Address: FEDERAL GOVERNMENT POLYCLINIC (PGMI) G-6/2, ISLAMABAD, Islamabad Capital Territory Schedule: 90 Days Quantity: 6/Qty	6/Qty	108000 PKR	Any	3 Years

Item	UNSPSC	Delivery Schedule	Quantity	Bid Security	Manufacturer / Dealer Authorization	Warranty
ElectroHydraulic Moveabel Bed	Patient bed or table scales for general use	Address: FEDERAL GOVERNMENT POLYCLINIC (PGMI) G-6/2, ISLAMABAD, Islamabad Capital Territory Schedule: 90 Days Quantity: 20/Qty	20/Qty	40000 PKR	Any	3 Years
ECG machine	Electrocardiography EKG graphic recorders	Address: FEDERAL GOVERNMENT POLYCLINIC (PGMI) G-6/2, ISLAMABAD, Islamabad Capital Territory Schedule: 90 Days Quantity: 10/Qty	10/Qty	80000 PKR	Any	3 Years

Item	UNSPSC	Delivery Schedule	Quantity	Bid Security	Manufacturer / Dealer Authorization	Warranty
Cardiac Monitors	Multiparameter vital sign units	Address: FEDERAL GOVERNMENT POLYCLINIC (PGMI) G-6/2, ISLAMABAD, Islamabad Capital Territory Schedule: 90 Days Quantity: 6/Qty	6/Qty	72000 PKR	Any	3 Years



Related Services of Goods:

Yes

Item	UNSPSC	Related Services
Angiography System (Cath Lab)	Cardiac catheterization lab equipment	as per specific terms and conditions of the bid documents and the specification of the item
Defibrillator machines	Mobile medical services automated external defibrillators AED or hard paddles	as per specific terms and conditions of the bid documents and the specification of the item
ElectroHydraulic Moveabel Bed	Patient bed or table scales for general use	as per specific terms and conditions of the bid documents and the specification of the item
ECG machine	Electrocardiography EKG graphic recorders	as per specific terms and conditions of the bid documents and the specification of the item
Cardiac Monitors	Multiparameter vital sign units	as per specific terms and conditions of the bid documents and the specification of the item

Items/Lot Specification

Items Without Lots :

Item: Angiography System (Cath Lab)

UNSPSC: Cardiac catheterization lab equipment

Specifications / Requirementss:

S.no:	Equipment name	Equipment description	Quantity
1	Cath Lab Machine	1. Angiography System (CEILING MOUNTED)	1
		2. Fluoroscopy unit	1
		3. Hemodynamic monitoring System	1
		4. Cath Lab Table	1
		5. Power Injectors.	1
		6. Intravascular ultrasound (IVUS).	1
		7. Fraction flow reserve (FFR) equipment	1
		8. Intra-Aortic Balloon Pump (IABP)	1
		9.IVL intravascular Lithotripsy balloon with consol (1) and catheters (10)	1
		10. Rota blator equipment (Rota Pro)	1

S.no:	Equipment name	Equipment description	Quantity
2	Radiation Protection Gear	Lead aprons, shields, and other radiation protection gear for staff and patients.	10
3	Anesthesia cart	Anesthesia delivery system or managing patient sedation during procedure	2
4	Sterile draping and supplies	Sterile drapes and other supplies for maintaining aseptic conditions.	10
5	Emergency Crash Cart	Trolley with medications and equipment for managing cardiac emergencies.	2
6	Cardiac Monitors	Continuous vitals monitoring system for tracking patient's cardiac status.	6
7	Echocardiographic machine	standard as per given specification	1
8	Portable Echocardiographic machine	standard as per given specification	1
9	Storage and documentation system	System for storing and documenting patient data and images along-with Soft wares for installed systems for reporting	2
10	Instrument sterilization equipment	Autoclaves and sterilization equipment for maintaining sterile instruments along with installation with area development for autoclave in cath lab	1
11	Emergency oxygen supply	To established the new oxygen supply connections with existing oxygen supply system of the FGPC.	1
12	I.T Equipment for Cath-Lab along with Software and accessories	Coronary and Peripheral Angiography Provision of 02 work stations of C.T.	L/S
13	Software for CT Angio	CT coronary and peripheral Angio software and Hardware compatible with already installed CT-Scanner machine at FGPC, Islamabad.	L/S
CEILING MOUNTED ANGIOGRAPHY MACHINE			

S.no:	Equipment name	Equipment description	Quantity
		Quality Certification: FDA 510K (mandatory)with any one of the following certificates.(CE,MHLW)	
		C. The firms must quote their latest and leading models from the above-mentioned origins with the proven past performance nationally and internationally. The firm must possess its related back up support services including trained engineers, workshop facilities, spare parts availability and repair/calibration tools etc. Firm must have PEC registration. The date of launch of quoted model must not be older than 5-6 years however version of the quoted model should be latest version.	
		D. The quoting firm must possess ISO certificate for service operations and should have proper infrastructure to handle and execute the complete package with previous experience.	
		E. The quoting firm must have installed at least 10-units of same equipment & model in Pakistan and must bring satisfactory recommendation letters from at least 10 local users along with installation certificates including govt sector.	
		F. The firm must be a nationwide sole distributor at least for the five consecutive years and should have sole agency from manufacturer and also must have an established track record government supply of over 5-years.	
		G. The most important criterion is the capability to provide quick and efficient after sales service at site. The hospital reserves the right to inspect workshop facilities of the vendor at any time to ascertain technical delivery capability. Bidders with inadequate facilities will not be considered.	
		H. Foreign training of four (04) hospital personnel is mandatory (Clinicians/Technicians/Biomedical Engineer). The firm shall arrange the training for the hospital personnel, including all expenses related to the training which shall be borne by the firm.	

S.no:	Equipment name	Equipment description	Quantity
		I. The firms must provide a letter of Warranty including all parts of complete machine for five years from the manufacturer starting from the date of installation. However, firm will quote further 10 years post warranty should be quoted which will be applicable after expiry of standard warranty period. The local vendor's warranty will not be acceptable	
Clinical Specialty:Cardiology			
	Generic Name:Single Plane Angiography Machine (Ceiling Mounted)		
	Clinical Purpose: Single Plane Angiography Machine is intended to perform diagnostic and therapeutic procedures.		
	TECHNICAL SPECIFICATIONS		
	Fully digital flat single plane cardiac angiography / cardiac catheterization system for interventional cardiac procedures alongwith peripheral and neuro procedures. Capable of head-to-toe coverage without repositioning of patient. Companies shall offer their latest systems and applications with best resolution and minimum possible radiation dose.		
	Ceiling Mounted Positioning Arm:		
	1. The position placement of the arm ceiling reflects on the capacity and capability of the equipment.		
	Geometry:		
	2. C-Arm or G - Arm Geometry (Better or More)		
		3. Motorized digital flat panel system with smart collision detector (Better or More)	
		4. Rotation Range (Degrees): ROA / LAO +/- 105° (Better or More)	
		5. Cranial / Caudal: Minimum +/-45° (Better or More)	

S.no:	Equipment name	Equipment description	Quantity
		6. Rotational Angiography Speed: 50° / sec or more in LAO to RAO or vice versa; 18° / sec or more for AP Cranial to AP Caudal or vice versa	
		7. Source to Image Distance (SID) Range: 95 - 105 cm or wider (PA /Lateral) (Better or More)	
		8. Iso-Centric Height: Variable / fixed	
		9. Auto Positioning: Programmable auto positioning of selected angulations, (50 or more) programmable positions (Better or More)	
		10. All rotational angles should be digitally displayed on the monitor	
		11. Motorized / manual parking & rotation of the positioning arm, capable of head-to-toe coverage without patient repositioning	
		Digital Flat Panel Detector:	
		30 x 30 cm or larger or Diagonal 40 cm (+ / - 4 cm) (Better or More)	
		Further Detector Specifications:	
		12. ASi / CsI / aSe Type	
		13. Image matrix of 1024 x 1024 pixels at minimum 16 bits depth for detector (Better or More)	
		14. Standard Cardiology Field of View (FOV) sizes: Three formats (Better or More)	
		15. Built in temperature stabilizer	

S.no:	Equipment name	Equipment description	Quantity
		16. Integrated Collision Protection / Collision Detection Technology 18. All other standard accessories according to this digital flat panel	
		17. Minimum Pixel Size: 200 µm or better/less	
		18. Removable grid for Paediatric applications	
		19. Minimum Spatial Resolution: 2.5 lp / mm (Better or More)	
		20. Minimum Detector / Detective Quantum Efficiency (DQE): More than 75 % (Better or More)	
		Patient Support Table:	
		21. Floor mounted, 8-ways horizontal floating top catheterization table with up / down, vertical, longitudinal, and transverse movements with pivoting.	
		Further Table Specifications:	
		22 Minimum table length 280 cm with unobstructed image coverage(Better or More)	
		23. Longitudinal Travel: 1100 mm with tabletop movement (Better or More)	
		24. Lateral Travel: 280 mm (Better or More)	
		Vertical Travel: 90 – 105 cm from the floor (Better or More)	
		20. CPR should be possible in any table position	
		21. Table top should be able to accept patients' weight of upto 200kg plus 100kg for	
		resuscitation (Better or More)	

S.no:	Equipment name	Equipment description	Quantity
		22. Tabletop should be metal free (X-ray density 2mm Al equivalent),(Better or More)	
		23. Complete accessories including arm holder, hand grip, arm support/arm rest and positioning aids	
		X-Rays Generator:	
		Microprocessor based high frequency using fiber optics for data communication between each imaging system	
		24. Dedicated X-Rays generator of 100 KW (Better or More)	
		25. Radiographic mA up to 1,000 (Better or More)	
		26. Radiographic kV 40 / 50 – 125kVp (Better or More)	
		27. Serial filming exposures with shortest exposure of 1 ms, with automatic KV and mA control for optimum image quality (Better or More)	
		28. Capability of digital radiography and fluoroscopy	
		29. Capability of doing multiple rate digital pulsed fluoroscopy and cine minimum at range of 3.75 to 30fps (Better or More)	
		30. Capability to change frame rate during the same study	
		X-Rays Tube:	
		The bidder will mention the models in their technical offer.	
		31. Tube (Liquid Metal Bearing Technology)	

S.no:	Equipment name	Equipment description	Quantity
		32. Dual / Triple focus with Anode Heat Storage Capacity: 3.8 MHU (actual) (Better or More)	
		33. Anode Small Focal Point: 0.3 – 0.5 or better or less	
		34. Anode Large Focal Point: 0.8 – 1.0 mm or better or less	
		35. Dose management with Fluoro filters range of 0.1 / 0.2 mm and 0.6 / 1.0 mm Cu and Real Time Automatic Filter selection based on patient bodyweight (Better or More)	
		36. State of art cooling system	
		Radiation Reduction and Monitoring Systems:	
		Dose management and reduction / optimization to be done without image quality compromise.	
		37. Grid pulsed / physical radiation reduction system / flat emitter technology / similar technology tubes with radiation dose management and with auto adjusting Fluoro filters	
		38. Dedicated X-Rays radiation dose monitoring and dose reduction software features of the latest variety provided by the manufacturer	
		39. Angulation dependent automatic exposure control including KV, mA, pulse width & focal spot adjustments	
		40. Automatic calculation and optimization of exposure data based on fluoroscopic data	
		41. Radiation free patient positioning without active Fluoro	

S.no:	Equipment name	Equipment description	Quantity
		42. Radiation free collimation	
		43. DAP threshold intimation / warning system	
		44. DICOM structured report containing patient/procedure/dose data	
		High Resolution Digital Imaging and Acquisition / Fluoroscopy System with Real Time Image Processing:	
		45. Acquisition, storage, and display all at minimum 16 bits (Better or More)	
		46. Parallel processing capability / multitasking facility	
		47. Real time filtering and roadmap function	
		48. Magnetic disk capacity for storage of minimum 100,000 images in 1024 x 1024 x 16 bit on the magnetic disk of main console (Better or More)	
		49. Minimum Cine Length: 10 seconds in 1024 matrix @ minimum 30fps (Better or More)	
		50. Images storage and retrieval from archive disk for possible manipulation and quantification using available software packages (compatible format / bits)	
		51. Acquisition at 1024 x 1024 @ 30fps and 512 x 512 @ 60fps for pediatrics should both be available (Better or More)	
		Monitoring System:	

S.no:	Equipment name	Equipment description	Quantity
		a) Medical grade large display monitor in examination room, minimum 55" diagonal in size 8 Mega Pixels resolution along with Two 19" or larger medical grade monitors for backup, one for live image, second monitor for reference image (Better or More)	
		52. Two LCD / LED medical grade monitors for live images and road mapping in the control room, minimum 19" inch diagonal or larger size (Better or More)	
		53. All 19" or larger Monitors should be Medical Graded, complied with International Standards for medical monitors (Better or More)	
		Controls:	
		54. Integrated table side touch screen console to adjust system parameters, frame rates, Haemodynamic system controls, quantitative analysis, stent enhancement soft-wares, Digital Subtraction Angiography (DSA), Mask- Unmask (Road-mapping) and Rotational Angiography	
		55. All controls of digital imaging system, measurements, post processing shall be in the control room as well as the examination room.	
		56. The control panel can be mounted at any side of the patent table	
		Recording / Archiving & Communication System:	
		57. Recording / archiving system should be DICOM-3 compatible	
		58. Digital images should be stored as backup on CDs / DVDs with incorporated original DICOM software	
		59. DICOM (Send / store, commitment, retrieve /query)	

S.no:	Equipment name	Equipment description	Quantity
		60. PACS compatibility /integration	
		61. Compatibility to display Intra-Vascular Ultrasound, OCT, and FFR system images on main screen	
		62. Compatibility to display of ultrasound / echocardiography system images on main screen	
		63. Integrated bi-directional intercom system for Cath lab to control room (from Original Manufacturer).	
		Software Packages:	
		Complete analysis package for following cardiac applications should be included:	
		64. Dynamic pre and post PTCA / Valvotomy comparison with one image live and other reference	
		65. Automatic loops replay after acquisition or fluoroscopy	
		66. Dynamic real time pan /zoom	
		67. Dynamic real time digital imaging processing like edge enhancement or gamma correction, noise reduction (spatial filtration)	
		68. Online image density (gray scale) correction	
		69. Simultaneous display of fluoroscopy and reference images	

S.no:	Equipment name	Equipment description	Quantity
		70. Facility to review previous studies in the examination room from the patients' old CD	
		71. Automatic positioning of the c-arm corresponding to reference image	
		72. Store Fluoro facility to store last fluoro scopy run	
		Dedicated Adult Features:	
		73. Digital subtraction angiography package with manual / real time pixel shift for compensation of patient movement during acquisition	
		74. Image masking with real-time overlay of live Fluoro	
		75. Real time live stent enhancement with integrated table side control with touch panel (older stent enhancement on frozen image is not acceptable)	
		76. Rotational angiography software	
		77. Quantitative coronary analysis software	
		Physiological Haemodynamic Monitoring System:	
		Original from manufacture of Angiography system.	
		78. Integrated Haemodynamic System with full table side control having bi-directional communication with the machine for automatic / synchronized acquisition, transfer and display of patient Haemodynamic parameters and demographic data	

S.no:	Equipment name	Equipment description	Quantity
		79. Multichannel (16 channels or more) to record at least 4 channels IBP, Cardiac output with thermo dilution method, Surface ECG in any configuration and simultaneous 12 lead ECG, NIBP and SpO2 measurements with Electrophysiology capability and hardware for 64 Unipolar channels/32 Bipolar Channel (Better or More)	
		Haemodynamic System Features:	
		80. Complete software for all Paediatric & adult, right / left heart, Angio / Valvular Haemodynamic calculations such as gradients, valve areas, shunts including annotations and 12 channels ECG	
		81. Live Waveforms display on main monitor in the examination room	
		82. Digital display of all the parameters like IBP, Heart rate, cardiac output parameters	
		83. Possibility to print the waveforms simultaneously while acquiring the data in the background	
		84. Capability to store the waveforms on the hard disk of the physiological recording system	
		85. Two Monitors in the control room, one for live waveforms and the other for control and data entry	
		86. Facility for freezing Haemodynamic data and simultaneous recovery of recent data/ compare stored data with current wave form.	
		Workstation:	

S.no:	Equipment name	Equipment description	Quantity
		Hardware must be as per original manufacturer's recommendations and Software should be from original manufacturer of Angiography System. Online workstation to review studies directly and the facility to review studies with lossless compression and original image quality as on console.	
		87. DICOM - 3 compatible	
		88. Capable of universal DICOM support	
		89. Original DICOM handling software from the manufacturer with permanent license	
		90. Edge enhancement/adjustable view speeds/post processing capabilities	
		91. High-Definition Medical Graded LCD / LED monitor of minimum 19 inches size (Better or More)	
		92. CD / DVD writer and CD / DVD ROM Drive	
		93. Image storage capacity as per manufacturers' approved hardware	
		94. SCSI or equivalent controller as per manufacturer's approved hardware	
		95. Black & white laser/ink jet printer	
		96. Dedicated application for TAVI planning on workstation/console.	
		Radiation Protection Equipment / Accessories:	
		Must be FDA / CE / MHLW approved.	
		97. Ceiling suspended tilt table lead glass for radiation protection of operator's head & neck regions and upper body parts	

S.no:	Equipment name	Equipment description	Quantity
		98. Table mounted adjustable lower body radiation shields / flaps on both sides	
		99. Lead glass window size 3 x 1.5 meter or larger lead equivalent 2.0 mm or better	
		100. Lead aprons (vest and skirt), double side with different sizes, light weight, lead equivalent front 0.5 mm and back 0.35 mm with belts (FDA / CE approved) (Qty.05)	
		101. Wall Mounted / Trolley mounted hangers of S.S 304L for lead aprons (To be supplied Locally)	
		102. Thyroid shields with lead caps.(FDA / CE approved) (Qty.05)	
		103. Head Cap (Lead) Qty-05	
		104. Leg Pads (Qty-05)	
		105. Lead Goggles: Light Weight (FDA / CE approved) (Qty.05)	
		106. Patient Protection Lead Sheet: Light weight having 30 inches in size approximately	
		Other Accessories:	
		107. Ceiling suspended shadow-less light	
		108. Programmable Contrast Media Power Injector (Medrad, Angiomat, Medtrone, Nemoto, Guerbet) with 100 disposable syringes	
		109. Operational & Service Manual	

S.no:	Equipment name	Equipment description	Quantity
		110. Brand New and Latest Version 160 KVA or more true online sine wave Double conversion UPS for whole system with a minimum back up time of 10 minutes including room lights, microprocessor based IGBT technology. Display and alarms of parameters. Three phase line voltage of 220 50Hz with all necessary standard parts including dry batteries with establishing on site electricity connection with Cath-Lab as per site requirement.	
		111. Diesel Generator Set 160 KVA Brand New from well-known brand (USA/Europe/Japan) along with complete installation as per requirement of hospital and its installation including base/Foundation and sound proof (60-65dB(A)) canopy OR lower measured at a distance of 7 Meters, with establishing on site electricity connection with Cath-Lab as per site requirement.	
		Quality and Safety Standards:	
		For Angiography FDA 510K (mandatory)with any one of the following certificates.(CE,MHLW)	
		OPTIONAL TO BE QUOTED IN STADNARD	
		Artificial Intelligence Deep Learning server for real time image processing to allow high quality noiseless images with two-time higher contrast to noise ratio as compared to noised images.	
		AI Deep Learning processing to allow for fluoro quality equivalent to DA run for usage of 2x times diluted contrast without any difference in image quality.	
	O	CIVIL WORK AND SITE RENOVATION:	

S.no:	Equipment name	Equipment description	Quantity
	P	Lead Lining and civil works of Cath room, i.e DB, Earthling, flooring, ACs, Ceiling and control room. . The Cath Lab installation will be on turnkey basis and BOQ as per given drawing (Site Drawing is provided in the section of Annexure) will be submitted alongwith technical offer. As per given drawing all civil/electric work, LED lining and establishing electric connection of cath lab with main power supply room of FGPC along with required cabel/s will be the responsibility of vendor. if any vendor wants to visit the site, firm official can come to FGPC during working hours from Monday to Saturday.	
		POWER REQUIREMENTS:	
		Three Phase with line voltage of 220V, 50Hz.	
		ACCESSORIES	
		Automatic CPR Machine with synchronization of Defibrillator for Cath Lab x 02	
		Intravascular Ultrasound Imaging System (IVUS)	
		Intravascular Ultrasound Imaging System to include Imaging Catheter and Ultrasound Imaging Console. 40MHz and 60MHz transducer frequency to provide more options for physicians during complex PCI procedures. 10mm/s pullback speed, 100FPS high framerate, and 150mm pullback distance to finish the vessel inspection in 15 seconds without any loss of details and to improve the reproducibility of tissue and optimizing workflow (Better or More)	
		Maximum 100FPS or better High Frame Rate (Better or More)	

S.no:	Equipment name	Equipment description	Quantity
		To enhance the temporal resolution of images and more precise display	
		Catheter Recognition	
		Identification of the catheter without contact	
		Hydrophilic Coating	
		To enhance crush ability and push ability	
		Intravascular Ultrasound Console	
		System should be equipped with double processors and large capacity SSD, to enable superior performance in processing high framerate images smoothly. It should provide more information to help physicians in assessing interventional therapy.	
		It should be integrated with advanced data analysis and processing algorithms.	
		Detection of the hierarchical structure of coronary vascular, and analyze the plaque area also the stenosis rate.	
		· Intelligent Image Analysis Platform	
		· Easy user interface	
		· DICOM/AVI/BMP/JPG, etc.	
		System to be supplied with 5-Qty. 40MHz and 5-Qty. 60MHz transducers (Better or More)	

S.no:	Equipment name	Equipment description	Quantity
		FFR system with separate 12 inches touch screen for FFR measurement ultrathin microcatheter 0.0205 or thinner in lesion profile which can be used with 0.014" guidewire for easy pullback and post-PCI measurements without reviewing. iFR/iFFR/CRR or similar to calculate without adenosine 1000 hours of recording facility to be built in. System complete with 5 microcatheters. (Better or More)	
		Should be compatible with offered Angiography System and can be from any global franchised manufacturer like China, Singapore and Taiwan etc.	
		Rota Blatter (Rota Pro)	
		Vibrant Digital Display – Enhanced feedback and deceleration indicator	
		Streamlined Connections – Quick and easy setup	
		IV Pole Clamp – Installation flexibility	
		IVL intravascular Lithotripsy balloon with consol (1) and catheters (10)	
		Tablet type Echo Machine with 5 years warranty and built Artificial Intelligence system with life time subscription, Trolley should be provided with machine as well.	2
		Intra-Aortic Balloon Pump	

S.no:	Equipment name	Equipment description	Quantity
		Self-contained Fiber optic based intra arotic ballon pump having mobile console with ECG amplifier with possible selection 5 leads arterial blood pressure amplifier. Discriminative triggering circuit to command balloon actions on patient's ECG arterial blood pressure curve or internal simulator 80 BPM. Colour graphic displays at least of 10" for display of arterial and pressure heart rate balloon volume used and alarm conditions with trouble shooting procedures. Wave form displays for ECG, arterial pressure and balloon pressure on three channel memory type oscilloscopes. Fall safe system. V Pacing switch. Progressive viewing sequence. Integrated battery power supply to take patient to catheterization labs, operating theatre or other hospital: 60-minute autonomy. CO2 / helium tank wrench. 5 lead ECG cable, male connector pressure, transducer adopter, chart recorder. (Better or More)	
		220 V, 50 Hz, Ac.	
		One spare set of patient cable.	
		System should be complete to display all the parameters with all accessories and catheters 12 Fr, 10, 5 Fr, 9.5 Fr (Disposable balloon catheters of varying sizes for use with the pump 25 nos.) (Better or More)	
		PORTABLE ECHOCARDIOGRAPHY SYSTEM	
		Quality Certification: FDA 510K (mandatory)with any one of the following certificates.(CE,MHLW)	

S.no:	Equipment name	Equipment description	Quantity
		Portable Digital Echocardiography System for Cardiology, Vascular, Operating theatre, point of care with frequency range of 2-14 MHz, DICOM compatible. (Better or More)	
		• Modes: B, M, Color Flow Mapping, Power Doppler, PW, CW	
		• Linear, Convex and Phased Array Technology Transducers should be compatible.	
		• Compact Portable Ultrasound System (5 kg, with battery)	
		• 12" IPS LCD Monitor Touch Screen. (Better or More)	
		• Tissue Harmonic Imaging.	
		• Scanning Depth: 38cm (Better or More)	
		• 1 Probe Connector on-board	
		• Rechargeable Built-in Battery with 60-minutes backup time. (Better or More)	
		• Dedicated Needle Visualization software.	
		• Automatic Image Optimization.	
		• Operating System should Windows 7 or Latest	
		• Hard Drive Integrated 500GB SSD; for image storage and software upgradation	
		• Input / Output: 3 x USB, 1 x SD card, 1 x LAN Ethernet, 1 x DVI-D	
		• Probes:	

S.no:	Equipment name	Equipment description	Quantity
		Multi Frequency Linear Probe (4 – 14 MHz) (Better or More)	
		Phased Array Sector Probe (2 – 4 MHz) (Better or More)	
		• Original pole cart	
		• 3 Connectors on Pole cart	
		ECHOCARDIOGRAPHY MACHINE	
		Quality Certification: FDA 510K (mandatory)with any one of the following certificates.(CE,MHLW)	
1		Digital Echocardiography unit for wide range of premium performance application of cardiovascular imaging in pediatrics and adult. System should have compatibility to handle up to 22MHz multifrequency probes. (Better or More)	
2		DISPLAY:	
		23” High-Definition LCD/LED Monitor with resolution 1920 x 1080, tilt able and swiveable type (Better or More)	
3		OPERATION MODES	
		B, 2D, M-Mode, Power Doppler, HPRF, Spectral Doppler, Color Doppler, THI, D-THI, TDI, Duplex and Triplex Doppler, PW Doppler, CW Doppler Steerable and ECG Gating.	
4		CONTROL PANEL	
A		Alphanumeric keyboard with trackball.	

S.no:	Equipment name	Equipment description	Quantity
B		12-Inches Touch Control Panel Screen.	
C		Direct access to system functions through dedicated keys.	
D		Adjustable transmit focusing control.	
E		TGC / STC: 8-step slide controls	
F		4-Active Transducer Connector for Transthoracic probes.	
5		CALIPERS / MEASUREMENTS:	
		Measurements for: Distance, angle, Stenosis %, area, circumferences, volume, slope, time, heart rate and acceleration. LV (left ventricular function) measurements, LA (left atrial volume) measurements, AV (aortic valve) measurements, MV (mitral valve) measurements, PV (pulmonary valve) measurements, PISA measurements, LV MASS measurements, Vascular measurements (CCA, ECA, ICA, VA, SA), Flex-M/Anatomical M-Mode, Auto EF LA & LV measurements, Auto IMT should also be provided.	
6		APPLICATION:	
		Cardiac, Peripheral, Pediatric, Adult Cephalic, and Trans esophageal with software for measurements.	
7		FRAME RATE:	
		2000fps or more. (Better or More)	
8		CINE MEMORY:	
		Cine Memory: 900MB or more. (Better or More)	

S.no:	Equipment name	Equipment description	Quantity
9		SYSTEM SCANNING DEPTH:	
		48cm (Better or More)	
10		SYSTEM FREQUENCY RANGE:	
		2 - 22 MHz (Better or More)	
11		STORAGE DEVICE:	
		Built-in CD / DVD Drive. 1000GB HDD. (Better or More)	
12		SYSTEM DYNAMIC RANGE:	
		Dynamic range minimum 300 dB (Better or More)	
13		COMMUNICATION SOFTWARE:	
		DICOM communication software for Storage (server / media), print, Verification (export/import), MPPS, MWM, Query / Retrieve.	
14		PORTS:	
		Video Output, 4 or more USB Ports, Networking. Provision for 3D export output for 3D printing. (Better or More)	
15		IMAGING MODES / TECHNIQUES:	
		- Tissue harmonic Imaging, Color Angio, Tissue Doppler Imaging.	
		- Exercise and pharmacological stress echo examinations. Data Acquisition & Review Mode. Stress echo module for storing and reporting stress echo.	
		- 2D Tissue Strain Imaging / Wall motion tracking along with polar map and graphic display,	

S.no:	Equipment name	Equipment description	Quantity
		local & whole myocardial wall motion parameter curve display.	
		- MPI (Myocardial Performance Index) value should be calculated from the time change curve in TDI.	
		- Artificial Intelligence for measurement of Auto EF, Auto E/A, Auto TR, LVOT, Auto Ao.	
		- Reference Imaging for Ultrasound to Ultrasound, Ultrasound to CT and Ultrasound to MRI for Image Comparison.	
16		STANDARD TRANSDUCERS:	
A		Multi frequency Linear Probe to cover 5.0 – 11.0 MHz for vascular. (Better or More)	1
B		Multi frequency Phased array single crystal sector probe to cover 2.0 – 5.0MHz for Adults. (Better or More)	1
C		Multi frequency Phased array sector probe to cover 3.0 – 8.0MHz for peads. (Better or More)	1
D		Multiplane TEE Transducer (3.0 – 6.0 MHz) for Adults (Better or More)	1
		18 MHz high frequency Linear Probe. (Better or More)	1
17		STANDARD ACCESSORIES	
		a. Compatible UPS.	
		b. Digital B/W Thermal Printer.	
		AUTOMATIC CPR MACHINE WITH DEFIBRILLATOR	

S.no:	Equipment name	Equipment description	Quantity
		Quality Certification: FDA 510K (mandatory)with any one of the following certificates.(CE,MHLW)	
		SPECIFICATIONS	
		· The system shall have clinical benefits documented in comparative, human studies that	
		are published in peer-reviewed medical journals that demonstrate the improvements	
		to the following clinical parameters.	
		a. Vital signs (EtCO2, SpO2, blood pressure, etc.).	
		b. Coronary perfusion pressure (CPP).	
		c. Return of spontaneous circulation (ROSC).	
		· The system shall insure the integrity of delivered compressions by:	
		a. Automatically fitting itself to patients to prevent under compression.	
		b. Adjusting force delivery to accommodate patient chest stiffness up to the 95th percentile.	
		· The system shall reduce the potential for patient injury by:	
		a. Automatically fitting itself to patients to prevent over compression.	
		b. Applying pressures to the patients' chest that does not exceed 6 pound per square inch (Better or More)	
		of contact surface.	

S.no:	Equipment name	Equipment description	Quantity
		c. Automatically halts compressions, and alerts user, when patient is in an unsafe position	
		· The system shall be designed for moving patients with standard extrication equipment	
		as follows:	
		a. Can be secured directly to stretchers.	
		b. Can be secured directly to, and operated with, patients on backboards.	
		c. Operable on patients with soft stretchers (i.e. Reeves).	
		d. Operable on a patient at a 45-degree angle in any axis.	
		· The system shall operate from rechargeable batteries.	
		· System weight, including battery shall not exceed 25 pounds. (Better or More)	
		· A carrying case shall be available for the system that meets the following requirements:	
		a. Constructed of rugged easy to clean and sanitize material.	
		B. Comes standard with straps that gives responders the option of carrying over the	
		shoulder, or as a backpack.	
		c. Includes secure pouches for spare battery, disposable, and accessories requirements:	

S.no:	Equipment name	Equipment description	Quantity
		· The system shall have demonstrated improved survival to discharge in a prospectively	
		randomized multicenter trial.	

Item: Defibrillator machines

UNSPSC: Mobile medical services automated external defibrillators AED or hard paddles

Specifications / Requirementss:

Sr#	Description
	Department: Cardiology
	Name: DEFIBRILLATOR
	Quantity:
1	Brand
2	Make & Model
3	Quality Certification: FDA 510K (mandatory)with any one of the following certificates.(CE,MHLW)
4	Warranty: Three years with all spare parts, during warranty period firm should maintain equipment by doing Planned Preventive Maintenance (PPM) as per principal recommendation and attending service call within 02 hours whenever called by end user. If any spare part required during installation or PPM will be responsibility of the firm.
	TECHNICAL SPECIFICATIONS:

Sr#	Description
	☐ Biphasic transthoracic (external) defibrillator with LCD colour display Synchronized output with ECG.
	☐ Should have Control of energy, charging/delivering on main panel and paddle.
	☐ Energy delivery on control panel for internal defibrillation.
	☐ The energy range should be adjustable for peds and adults up to 200Joules.
	☐ Charging Time for full energy should be less than 07 sec.
	☐ Screen Size of approx. 6.5-inch colored or better.
	☐ Display of HR, ECG through paddles and Lead I,II & III patient cable.
	☐ Built in recorder for printing of full summery on standard 50mm paper or better.
	☐ Alarms for High and low Heart rate, low battery warning.
	☐ Built-in Rechargeable battery for maximum 100 shocks at max energy with shelf life of 3-years.
	☐ Monitoring time should be at least 3 hours or better
	☐ Should have Auto tester / self-check.
	☐ External Paddles with integrated Adult & Peads.
	☐ AED Facility with disposable pad having shelf life for at least 2 years.
	☐ Pacing facility with technology to use less current by using a 40 msec constant current and the pacing current should be 0 - 140mA.
	☐ AC 220 / 50hz operated.
	☐ Complete with standard accessories, including reusable type cables
	(Mandatort to Quote)

Sr#	Description
	☐ SPO2 (Pulse Oximeter)
	☐ NIBP (Non-Invasive Blood Pressure)
	☐ ETCO2 (Capnography)
	☐ Reusable Autoclavable Internal Paddles

Item: ElectroHydraulic Moveabel Bed

UNSPSC: Patient bed or table scales for general use

Specifications / Requirementss:

ElectroHydraulic Moveabel Bed			
Sr.#	Description	Compliance status Yes/No	Page #
	Department : Cardiology		
	Name: ElectroHydraulic Moveable Bed		
	Quantity: 30		
	Description		
1	Brand		
2	Make & Model		

ElectroHydraulic Moveabel Bed

3	Quality Certification: Dual certificate, as desired earlier in the bidding document under the heading of quality certification.		
4	Warranty: Three year with all spare parts, during warranty period firm should maintain equipment by doing PPM as per principal recommendation and attending service call within 02 hours whenever called by end user. If any spare part required during installation or PPM will be responsibility of the firm.		
5	Original Technical data sheet		
	TECHNICAL PARAMETERS		
	Technical Specification		
1)	Should have positions Back Raise, Knee Raise, Hi-Low Adjustment, Auto Contour, Trendelenburg, Reverse Trendelenburg and Chair Position and Flat Position with X-ray imaging base, low height indicator, Lock button, Electric CPR Button, and Power Indicator and battery Back-Up.		
	Movements:		
	Stress Saver Raise		
	(Stress on the patient's body should be reduced during back raise) (Better or More)		
	Back Raise Angle 0-70° (Better or More)		
	Knee Raise Angle 0-25° (Better or More)		
	Hi-low adjustment from 350mm to 700mm (Mattress Base) (Better or More)		
	Trendelenburg/Reverse Trendelenburg position 0 to 12° (Better or More)		

ElectroHydraulic Moveabel Bed

	Chair position button and Flat position button (Better or More)		
	Controls		
	Side-rail integrated panel should be included switches for the bed operations that are performed.		
	Side-rail integrated panel should be available in both side-rails near to the headboard, on the right and left side.		
	The patient control panel should be available in both side-rails near to the headboard, on the right and left side.		
	The angle of the back section base and the knee section base should be smoothly adjusted with no increments by using the patient control panel.		
	CPR Function		
	Should have Electric CPR Button & Manual CPR Lever		
	Should be quickly returns the patient to a horizontal position to execute CPR procedures.		
	Side Rails		
	Anti-Tempering Lock		
	Mattress Base		
	Low Bed Height		
	The low bed height of 350mm should ensure that patients should firmly place their heels on the floor (Better or More)		
	Angle Indicator		

ElectroHydraulic Moveabel Bed

	Head/Foot Board		
	Should be secured to prevent accidental removal.		
	The boards should be attached or detached.		
	Corner Bumpers		
	X-ray imaging using cassette		
	X-ray imaging using cassette should be possible in the back section area.		
	Adjusting the Angle of the Leg Section Base		
	The angle of the leg section base should be manually adjusted when the knee section base raised.		
	Restraint Belt mounting Holes		
	When using restraint belt, attach the restraint belt to the mounting holes below		
	Accessory Holders		
	The accessory holders at both sides of the hip section base and side-rails, on which items such as urine bags or other items should be hung.		
	Mattress Stopper		
	Should be on leg section base.		
	Option Attachment Holes		
	Option attachment holes should be at the four corners of the bed.		
	It should be installed IV pole or other accessories.		

ElectroHydraulic Moveabel Bed

Locking System		
The casters should be set to either of the positions below by operating the caster operation pedal on the both side of the foot end.		
Should have Lock Position, Free Position and Steering Position.		
Castors		
Should have 125 mm dia. single wheel castors.		
Bed Dimensions should be		
Bed Length 2195 mm or more without extension.		
Bed Width 985 mm or more		
Total height 755 mm to 1105 mm or more		
Mattress Base Height 350 mm to 700 mm or more		
Mattress		
Size: W: 860mm L: 2000mm		
Mattress Cover should be:		
MRSA antibacterial treatment.		
Water-proof treatment.		
Material Cover Polyester/ padding, urethane foam.		
Mattress must be manufactured by the same company manufacturing beds.		
Battery		

ElectroHydraulic Moveabel Bed

Should have Battery with low battery indicator.		
Safe working load 230 KG or more.		
Additional Accessories		
Bedside Table (Over Bed Table)		
Height Adjustment through Gas Spring		
Width 900mm x Depth 400mm x Height 745 to 1,130mm		
Castor: 40mm dia x 4pcs		
Bedside Cabinet:		
Pull-out Table (Detachable)		
Towel Rail at the both sides		
Detachable Drawer		
Inner Shelf		
40mm dia castor x 4pcs		
Dimensions: (W) 445mm x (D) 460mm x (H) 780mm		
Body power coated steel		
Table & Cabinet top & Drawer: ABS Plastic		
IV Pole		
Should be used for hanging intravenous drip bags and similar items.		

ElectroHydraulic Moveabel Bed

Should be adjust the length.

Item: ECG machine

UNSPSC: Electrocardiography EKG graphic recorders

Specifications / Requirementss:

Sr#	Description
	Department: Cardiology
	Name: ECGMACHINE 12-CHANNEL
	Quantity:
1	Brand
2	Make & Model
3	Quality certification: Any Dual Certification in the following certificates. (FDA,CE, MHLW)
4	Warranty: Three years with all spare parts, during warranty period firm should maintain equipment by doing Planned Preventive Maintenance (PPM) as per principal recommendation and attending service call within 02 hours whenever called by end user. If any spare part required during installation or PPM will be responsibility of the firm.
	TECHNICAL SPECIFICATIONS:
	☐ Twelve channel ECG on at least 12 inches Color Touch screen display or better.
	☐ Internal Storage upto 600 files or better

Sr#	Description
	☐ File formats. PDF, SCP, XML, DICOM, either natively or through compatible ECG management software
	☐ Thermal recorder for printing out, 3, 4, 6, 12 channels simultaneously.
	☐ Print Speed. 5 / 6.25 / 10 / 12.5 / 25 / 50 mm/s, user selectable
	☐ Heart rate range. 30 to 300bpm
	☐ Sensitivity / gain. 1.25, 2.5, 5, 10, 20, 10/5 mm/mV, AGC with automatic gain/auto setting or equivalent
	☐ Built in interpretation software.
	☐ Sample Frequency. 16,000Hz.
	☐ Defibrillator protection. accordance with applicable IEC 60601 requirements
	☐ Built in thermal recorder for 210 mm paper Z-fold paper.
	☐ Mains AC as well as battery backup of 5 hours Operations
	☐ Built-in AC interference, noise filter and baseline drift control.
	☐ Capability to interface with LAN/WLAN for data transfer.
	ACCESSORIES:
	☐ Complete with standard accessories.
	☐ Local mobile cart with imported wheels.

Item: Cardiac Monitors

UNSPSC: Multiparameter vital sign units

Specifications / Requirementssss:

Cardiac Monitor/Vital Sign Monitor
Description
Department:Cardiology
Name: Cardiac Monitor
Quantity: 06
Description
Brand
Make & Model
Quality Certification: Any Dual Certification in the following certificates. (FDA,CE, MHLW)
Warranty: Three years with all spare parts, during warranty period firm should maintain equipment by doing Planned Preventive Maintenance (PPM) as per principal recommendation and attending service call within 02 hours whenever called by end user. If any spare part required during installation or PPM will be responsibility of the firm.
Technical Specification
1. 17.3" high-definition touch screen design, easy and convenient to operate.(more or better)
2. Independent physiological and technical alarm function, convenient for medical staff to quickly know patient's condition.
3. Luminous "ON / OFF" button design, convenient to operate at night.
4. Modular design: the modules can be configured and combined flexibly and conveniently, expanded and upgraded at any time, meets the monitoring requirements of different departments on different parameters.

Cardiac Monitor/Vital Sign Monitor

5. Support dynamic hot-plugging parameter module: no need to restart the device, plug-and-play, change the monitoring parameters at any time as clinical requirements.
6. Adopt high-performance processor, stable, reliable and fast to process information.
7. Patient information can be rapidly entered, patient type (adult / neonate / pediatric) can be quickly switched, meets the requirements of different departments.
8. Abundant analysis functions: 23 arrhythmia, full-lead ST-segment and pacing analysis; multiple calculation functions (such as calculation of drug concentration, titration table, hemodynamics, ventilation, oxygenation, renal function, etc.).
9. With the functions of waveform freezing, holographic waveform storage and review.
10. Automatic data storage in case of power failure; achieves large-capacity data storage.
11. With HDMI expansion interface, the display screen can be connected according to clinical requirements.
12. Built-in large-capacity detachable lithium battery, ensures uninterrupted monitoring.
13. Fanless design, ensures no noise and low power consumption during working.
14. Standby mode, be applied to ICU, avoids affecting patient's rest, one-button switching, reduces the workload of medical staff.

Accessories:-

Standard accessories

NIBP CUFF 4 Nos

Spo2 Probe 4 Nos

ECG Lead 4 Nos

Temperature lead 4 Nos



Price Schedule

For Individual Items

#	Item Title	Quantity	Unit Price (PKR)	Total Price (PKR)	Delivery Location	Delivery Period / Year	Country of Origin
1							
2							

For Lots

#	Lot Title	Total Lot Price (PKR)	Country of Origin
1	[Lot 1 Title]		





General Conditions of Contract

A. General

1. Definitions

1.1 Unless the context otherwise requires, the following terms whenever used in this Contract shall have the same meaning and shall be interpreted as indicated

- a. “Applicable Law” means the laws and any other instruments having the force of law in the Government’s Country, or in such other country as may be specified in the Special Conditions of the Contract (SC), as they may be issued and in force from time to time;
- b. The “Arbitrator” is the person appointed with mutual consent of both the parties, to resolve contractual disputes as provided for in the General Conditions of the Contract GCC Clause 35 hereunder.
- c. “Authority” means Public Procurement Regulatory Authority.
- d. The “Commencement Date” is the date when the Supplier shall commence execution of the contract as specified in the SCC.
- e. “Completion date” means the date of completion of the Contract by the Bidder as certified by the Procuring Agency.
- f. “Procuring Agency” means:-
 - a. any Ministry, Division, Department or any Office of the Government;
 - b. any authority, corporation, body or organization established by or under a Law or which is owned or controlled by the Government;
- g. “The Contract” means an agreement enforceable by law;
- h. “The Contract Price” means the price payable to the Bidder under the Contract for the full and proper performance of its contractual obligations;
- i. “Ancillary Services” means those services ancillary to the provision of Goods, such as transportation and insurance, and any other incidental services, such as installation, commissioning, provision of technical assistance, training, and other such obligations of the Bidder covered under the Contract;
- j. “GCC” means the General Conditions of Contract contained in this section;
- k. “SCC” means the Special Conditions of Contract by which the GCC may be amended or supplemented;
- l. “Day” means calendar day unless indicated otherwise.
- m. “Defective Goods” are those goods which are below standards, requirements or specifications stated by the Contract.
- n. “Delivery” means the transfer of the goods from the supplier equipment, machinery, and /or other materials which the Supplier is required to supply to the Procuring Agency under Contract.

o. "Effective Date" means the date on which this Contract comes into force and effect.

p. "Force Majeure" means an unforeseeable event which is beyond reasonable control of either Party and which makes a Party's performance of its obligations under the Contract impossible or so impractical as to be considered impossible under the circumstances.

For the purposes of this Contract, "Force Majeure" means an event which is beyond the reasonable control of a Party, is not foreseeable, is unavoidable, and its origin is not due to negligence or lack of care on the part of a Party, and which makes a Party's performance of its obligations hereunder impossible or so impractical as reasonably to be considered impossible in the circumstances. and includes, but is not limited to, war, riots, civil disorder, earthquake, fire, explosion, storm, flood, epidemics, or other adverse weather conditions, strikes, lockouts or other industrial action (except where such strikes, lockouts or other industrial action are within the power of the Party invoking Force Majeure to prevent), confiscation or any other action by Government agencies.

q. "The Bidder" means the individual or corporate body whose Bids to provide the Goods has been accepted by the Procuring Agency;

r. "The Project Site," where applicable, means the place or places named in Bids Data Sheet and technical Specifications;

s. "Government" means the Government of Pakistan;

t. "Subcontractor" means any entity to which the Bidder subcontracts any part of the Goods.

u. "The Services" means the work to be performed by the Bidder pursuant to this Contract and as prescribed in the Specifications and Schedule of Activities included in the Bidder's Bids;

v. "Party" means the Procuring Agency or the Bidder, as the case may be, and "Parties" means both of them;

w. "Foreign Currency" means any currency other than the currency of the country of the Procuring Agency;

x. "Completion Date" means the date of completion of the contract by the Bidder as certified by the Procuring Agency;

y. "In Writing" means communicated in written form with proof of receipt;

z. "Local Currency" means the currency of Pakistan;

a. "Origin" means the place where the Goods were mined, grown, or produced or from which the Services are supplied. Goods are produced when, through manufacturing, processing, or substantial and major assembly of components, a commercially recognized new produce results that is substantially different in basic characteristics or in purpose or utility from its components.

b. "Specification" means the Specification of the Goods and performance of incidental services in accordance with the relevant standards included in the Contract and any modification or addition made or approved by the Procuring Agency.

c. "Procuring Agency" means: -

any Ministry, Division, Department or any Office of the Federal Government;

any authority, corporation, body or organization established by or under a Federal Law or which is owned or controlled by the Federal Government;

2. Application and Interpretation

2.1 These General Conditions shall apply to the extent that they are not superseded by provisions of other parts of the Contract.

2.2 In interpreting these Conditions of Contract headings and marginal notes are used for convenience only and shall not affect their interpretations unless specifically stated; references to singular include the plural and vice versa; and masculine include the feminine. Words have their ordinary meaning under the language of the Contract unless specifically defined.

3. Applicable Law

3.1 The contract shall be governed and interpreted in accordance with the laws of Pakistan, unless otherwise specified in SCC.

4. Governing Language

4.1 The Contract as well as all correspondence and documents relating to the Contract exchanged between the Bidder and the Procuring Agency, shall be written in the **English language** unless otherwise stated in the **SCC**. Supporting documents and printed literature that are part of the Contract may be in another language provided these are accompanied by an accurate translation of the relevant passages in English, in which case, for purposes of interpretation of the Contract, this translation shall govern.

5. Notices

5.1 Any notice, request, or consent made pursuant to this Contract shall be in writing and shall be deemed to have been made when delivered in person to an authorized representative of the Party to whom the communication is addressed, or when sent by registered mail, telex, telegram, or facsimile to such Party at the address specified in the **SCC**.

6. Standard

6.1 The Goods supplied under this Contract shall conform to the standards mentioned in the Technical Specifications, and, when no applicable standard is mentioned, the American Standards (such as ACI, IEEE, ASME, etc.) or the Pakistani standards such as PSQCA Such standards shall be the latest issued by the concerned institution.

7. Delivery/Location

7.1 The Goods shall be delivered to such locations as the Procuring Agency may approve and as specified in SCC.

8. Authorized Representatives / Authority of Member in charge

8.1 Any action required or permitted to be taken, and any document required or permitted to be executed, under this Contract by the Procuring Agency or the Bidder may be taken or executed by the officials specified in the **SCC**.

B. Commencement, Completion, Modification, and Termination of Contract

8. Effectiveness of Contract

8.1 This Contract shall come into effect on the date the Contract is signed by both parties and such other later date as may be stated in the SCC.

9. Commencement of the Contract

9.1 The Bidder shall confirm availability of Key Experts and begin carrying out the Services not later than the number of days after the Effective Date specified in the SCC.

10. Program

10.1 Before commencement of the Contract, the Bidder shall submit to the Procuring Agency for approval a Program showing the general methods, arrangements, order and timing for all activities. The Services shall be carried out in accordance with the approved Program as updated.

11. Starting Date/Expiration Date

11.1 The Bidder shall start carrying out the Services Five (05) days after the date the Contract becomes effective, or at such other date as may be specified in the SCC.

11.2 Unless terminated earlier pursuant to Clause **GCC 16** hereof, this Contract shall expire at the end of such time period after the Effective Date as specified in the SCC.

12. Entire Agreement

12.1 This Contract contains all covenants, stipulations and provisions agreed by the Parties. No agent or representative of either Party has authority to make, and the Parties shall not be bound by or be liable for, any statement, representation, promise or agreement not set forth herein.

13. Modification

13.1 Any modification or variation of the terms and conditions of this Contract, including any modification or variation of the scope, may only be made by written agreement between the Parties. However, each Party shall give due consideration to any Bids for modification or variation made by the other Party.

13.2 In cases of any modifications or variations, the prior written consent of the Procuring Agency is required.

14. Force Majeure

14.1 Definition

For the purposes of this Contract, "Force Majeure" means an event which is beyond the reasonable control of a Party and which makes a Party's performance of its obligations under the Contract impossible or so impractical as to be considered impossible under the circumstances.

14.2 No Breach of Contract

The failure of a Party to fulfill any of its obligations under the contract shall not be considered to be a breach of, or default under, this Contract in so far as such inability arises from an event of Force Majeure, provided that the Party affected by such an event (a) has taken all reasonable precautions, due care and reasonable alternative measures in order to carry out the terms and conditions of this Contract, and (b) has informed the other Party as soon as possible about the occurrence of such an event.

14.3 Extension of Time

Any period within which a Party shall, pursuant to this Contract, complete any action or task, shall be extended for a period equal to the time during which such Party was unable to perform such action as a result of Force Majeure. However, such period shall not exceed sixty (60) days.

14.4 Payments

During the period of their inability to perform the Services as a result of an event of Force Majeure, the Bidder shall be entitled to continue to be paid under the terms of this Contract, as well as to be reimbursed for additional costs reasonably and necessarily incurred by them during such period for the purposes of the Services and in reactivating the Service after the end of such period.

15. Termination

15.1 By the Procuring Agency

The Procuring Agency may terminate this Contract in case of the occurrence of any of the events specified in paragraphs (a) through (e) of this Clause. In such an occurrence the Procuring Agency shall give at least thirty (30) calendar days' written notice of termination to the Bidder in case of the events referred to in (a) through (d); at least sixty (60) calendar days' written notice in case of the event referred to in (e);

- a. If the Bidder fails to remedy a failure in the performance of its obligations hereunder, as specified in a notice of suspension;
- b. If the Bidder becomes (or, if the Bidder consists of more than one entity, if any of its members becomes) insolvent or bankrupt or enter into any agreements with their creditors for relief of debt or take advantage of any law for the benefit of debtors or go into liquidation or receivership whether compulsory or voluntary;
- c. If the Bidder fails to comply with any final decision reached as a result of arbitration proceedings;
- d. If, as the result of Force Majeure, the Bidder is unable to perform a material portion of the Services for a period of not less than sixty (60) calendar days;
- e. If the Procuring Agency, in its sole discretion and for any reason whatsoever, decides to terminate this Contract;

15.2 By the Bidder

The Bidder may terminate this Contract, by not less than thirty (30) calendar days' written notice to the Procuring Agency, in case of the occurrence of any of the events specified in paragraphs (a) through (d) of this Clause.

- a. If the Procuring Agency fails to pay any money due to the Bidder pursuant to this Contract and not subject to dispute within forty-five (45) calendar days after receiving written notice from the Bidder that such payment is overdue.
- b. If, as the result of Force Majeure, the Bidder is unable to perform a material portion of the Services for a period of not less than sixty (60) calendar days.
- c. If the Procuring Agency fails to comply with any final decision reached as a result of arbitration.
- d. If the Procuring Agency is in material breach of its obligations pursuant to this Contract and has not remedied the same within forty-five (45) days (or such longer period as the Bidder may have subsequently approved in writing) following the receipt by the Procuring Agency of the Bidder's notice specifying such breach.

C. Obligations of the Bidder

16. General

16.1 Standard of Performance

1. The Bidder shall deliver the product and carry out the Services with all due diligence, efficiency and economy, in accordance with generally accepted professional standards and practices, and shall observe sound management practices, and employ appropriate technology and safe and effective equipment, machinery, materials and methods. The Bidder shall always act, in respect of any matter relating to this Contract or to the Services, as a faithful adviser to the Procuring Agency, and shall at all times support and safeguard the Procuring Agency's legitimate interests in any dealings with the third parties.

16.2 Law Applicable to Goods

The Bidder shall deliver the goods in accordance with the Contract and in accordance with the Law of Pakistan and shall take all practicable steps to ensure that any of its Experts and Sub-Bidders, comply with the Applicable Law.

17. Conflict of Interests

17.1 Bidder Not to Benefit from Commissions and Discounts.

The remuneration of the Bidder shall constitute the Bidder's sole remuneration in connection with this Contract or the Services, and the Bidder shall not accept for their own benefit any trade commission, discount, or similar payment in connection with activities pursuant to this Contract or to the Services or in the discharge of their obligations under the Contract, and the Bidder shall use their best efforts to ensure that the Personnel, any Subcontractors, and agents of either of them similarly shall not receive any such additional remuneration.

17.2 Bidder and Affiliates Not to be Otherwise Interested in Project

The Bidder agree that, during the term of this Contract and after its termination, the Bidder and its affiliates, as well as any Subcontractor and any of its affiliates, shall be disqualified from providing Goods for any project resulting from or closely related to the Services.

17.3 Prohibition of Conflicting Activities

Neither the Bidder nor its Subcontractors nor the Personnel shall engage, either directly or indirectly, in any of the following activities:

- a. during the term of this Contract, any business or professional activities in the Government's country which would conflict with the activities assigned to them under this Contract;
- b. during the term of this Contract, neither the Bidder nor their Subcontractors shall hire public employees in active duty or on any type of leave, to perform any activity under this Contract;

18. Confidentiality

18.1 Except with the prior written consent of the Procuring Agency, the Bidder and the Experts shall not at any time communicate to any person or entity any confidential information acquired in the course of the contract.

19. Insurance to be Taken Out by the Bidder

19.1 The Bidder(a) shall take out and maintain, and shall cause any Subcontractors to take out and maintain, at its (or the Subcontractors', as the case may be) own cost but on terms and conditions approved by the Procuring Agency, insurance against the risks, loss or damage, and for the coverage, as shall be specified in the SCC; and (b) at the Procuring Agency's request, shall provide evidence to the Procuring Agency showing

that such insurance has been taken out and maintained and that the current premiums have been paid.

20. Bidder's Actions Requiring Procuring Agency's Prior Approval

20.1 The Bidder shall obtain the Procuring Agency's prior approval in writing before taking any of the following actions:

- (a) appointing such members of the Personnel not provided by the Bidder;
- (b) changing the Program of activities; and
- (c) any other action that may be specified in the SCC.

21. Reporting Obligations

21.1 The Bidder shall submit to the Procuring Agency the reports and documents in the numbers, and within the periods as prescribed by the Procuring Agency.

22. Liquidated Damages

22.1 If the Supplier fails to deliver any or all of the Goods or to perform the Services within the period(s) specified in the Contract, the Procuring Agency shall, without prejudice to its other remedies under the Contract, deduct from the Contract Price, as liquidated damages, a sum equivalent to the percentage specified in **SCC** of the delivered price of the delayed Goods or unperformed Services for each week or part thereof of delay until actual delivery or performance, up to a maximum deduction of the performance security (or guarantee) specified in **SCC**. Once the said maximum is reached, the Procuring Agency may consider termination of the Contract pursuant to **GCC Clause 16**.

22.2 Correction for Over-payment

If the Intended Completion Date is extended after liquidated damages have been paid, the Procuring Agency shall correct any overpayment of liquidated damages by the Bidder by adjusting the next payment certificate. The Bidder shall be paid interest on the overpayment, calculated from the date of payment to the date of repayment, at the rates specified in SCC.

22.3 Lack of performance penalty

If the Bidder has not corrected a Defect within the time specified in the Procuring Agency's notice, a penalty for Lack of performance will be paid by the Bidder. The amount to be paid will be calculated as a percentage of the cost of having the Defect corrected, assessed as specified in the SCC.

23. Performance Guarantee

23.1 The Performance Security (or Guarantee) shall be provided to the Procuring Agency no later than the date specified in the Letter of Acceptance and shall be issued in an amount and form acceptable to the Procuring Agency as specified in SCC, and denominated in the types and proportions of the currencies in which the Contract Price is payable as specified in the SCC.

23.2 Within Seven (07) days from the issuance of acceptance letter from the Procuring Agency, the successful Bidder shall furnish the Performance Guarantee in shape of ----- at the discretion of the PA in the amount **specified in SCC**. In case the amount of Bids security is equal or greater than

23.3 The proceeds of the Performance Guarantee shall be payable to the Procuring agency as compensation for any loss resulting from the Supplier's failure to complete its obligations under the Contract.

23.4 The Performance Guarantee shall be denominated in the currency of the Contract, or in a freely convertible currency acceptable to the Procuring agency and shall be in the acceptable form as specified in SCC.

23.5 The Performance Guarantee will be discharged by the Procuring agency and returned to the Supplier not later than thirty (30) days following the date of completion of the Supplier's performance obligations under the Contract, including any warranty obligations, unless otherwise **specified in SCC**.

23.6 The bid security of the successful bidder shall not be substituted with the Performance Guarantee.

24. Fraud and Corruption

24.1 The Procuring Agency requires the Supplier to disclose any commissions or fees that may have been paid or are to be paid to agents or any other party with respect to the Bidding process or execution of the Contract. The information disclosed must include at least the name and address of the agent or other party, the amount and currency, and the purpose of the commission, gratuity or fee.

25. Sustainable Procurement

25.1 The Bidder shall conform to the sustainable procurement contractual provisions, if and as specified in the SCC.

D. Bidder's Personnel

26. Description of Personnel

26.1 The titles, agreed job descriptions, minimum qualifications, and estimated periods of engagement in the carrying out of the Services of the Bidder's Key Personnel. The Key Personnel listed by title as well as by name are hereby approved by the Procuring Agency.

27. Removal and/or Replacement of Personnel

27.1 Except as the Procuring Agency may otherwise agree, no changes shall be made in the Key Personnel. If, for any reason beyond the reasonable control of the Bidder, it becomes necessary to replace any of the Key Personnel, the Bidder shall provide as a replacement a person of equivalent or better qualifications.

27.2 If the Procuring Agency finds that any of the Personnel have (i) committed serious misconduct or have been charged with having committed a criminal action, or (ii) have reasonable cause to be dissatisfied with the performance of any of the Personnel, then the Bidder shall, at the Procuring Agency's written request specifying the grounds thereof, provide as a replacement a person with qualifications and experience acceptable to the Procuring Agency.

27.3 The Bidder shall have no claim for additional costs arising out of or incidental to any removal and/or replacement of Personnel.

E. Obligations of the Procuring Agency

28. Assistance and Exemptions

28.1 The Procuring Agency shall use its best efforts to ensure that the Government shall provide the Bidder such assistance and exemptions as specified in the SCC.

29. Change in the Applicable Law

29.1 If, after the date of this Contract, there is any change in the Applicable Law with respect to taxes and duties which increases or decreases the cost of the related Services rendered by the Bidder, then the remuneration and reimbursable expenses otherwise payable to the Bidder under this Contract shall be increased or decreased accordingly by agreement between the Parties, and corresponding adjustments shall be made to the amounts referred in the SCC.

30. Services and Facilities

30.1 The Procuring Agency shall make available to the Bidder and the Experts, for the purposes of the Services and free of any charge, the services, facilities and property described , at the times and in the manner specified in the SCC or terms of reference.

30.2 In case that such services, facilities and property shall not be made available to the Bidder, the Parties shall agree on (i) any time extension that it may be appropriate to grant to the Bidder for the performance of the Services, (ii) the manner in which the Bidder shall procure any such services, facilities and property from other sources, and (iii) the additional payments, if any, to be made to the Bidder as a result thereof.

F. Payments to the Bidder

31. Contract Price

31.1 The price payable shall be in Pakistani Rupees unless otherwise specified in the SCC. Prices charged by the Supplier for Goods delivered under the Contract shall not vary from the prices quoted by the Supplier in its Bid.

32. Terms and Conditions of Payment

32.1 Payments will be made to the Bidder according to the payment schedule stated in the SCC and as per actual invoice submitted by the Bidder.

32.2 Unless otherwise stated in the SCC, the advance payment shall be made against the provision by the Bidder of a bank guarantee for the same amount, and shall be valid for the period stated in the SCC. Any other payment shall be made after the conditions listed in the SCC for such payment have been met, and the Bidder have submitted an invoice to the Procuring Agency specifying the amount due.

33. Currency of Payment

33.1 Any payment under this Contract shall be made in the currency(ies) specified in the SCC.

G. Quality Control

34. Identifying Defects

34.1 The principle and modalities of Inspection of the Goods by the Procuring Agency shall be as indicated in the SCC. The Procuring Agency shall check the Bidder's performance and notify him of any Defects that are found. Such checking shall not affect the Bidder's responsibilities. The Procuring Agency may instruct the Bidder to search for a Defect and to uncover and test any service that the Procuring Agency considers may have a Defect. Defect Liability Period is as defined in the SCC.

35. Correction of Defects, and

Lack of Performance Penalty

35.1 The Procuring Agency shall give notice to the Bidder of any Defects before the end of the Contract. The Defects liability period shall be extended for as long as Defects remain to be corrected.

35.2 Every time notice a Defect is given, the Bidder shall correct the notified Defect within the length of time specified by the Procuring Agency's notice.

35.3 If the Bidder has not corrected a Defect within the time specified in the Procuring Agency's notice, the Procuring Agency will assess the cost of having the Defect corrected, the Bidder will pay this amount, and a Penalty for Lack of Performance.

36. Taxes and Duties

36.1 A Supplier shall be entirely responsible for all taxes, duties, fees, etc., incurred until delivery of the contracted Goods to the Procuring Agency.

H. Settlement of Disputes

37. Alternate Dispute Resolution

37.1 Upon the commencement of the Procurement Contract, any disputes arising between the contracting parties shall be resolved in accordance with Rule 66 of the Public Procurement Rules, 2026.

37.2 The procuring agency may refer the matter to the Chief Justice Islamabad High Court or Managing Director PPRA or the Secretary Ministry of Law & Justice for appointment of Arbitrator.

37.3 The fee for the Arbitrator shall be specified in Pak Rupees as determined by the appointing authority which shall be borne and shared equally by the contracting parties.

38. Change Orders

The Procuring Agency may at any time, by a written order given to the Supplier, make changes within the general scope of the Contract in any one or more of the following:

- a) Drawings, designs, or specifications, where Goods to be furnished under the Contract are to be specifically manufactured for the Procuring Agency;
- b) The method of shipment or packing;
- c) The place of delivery; and/or
- d) The Services to be provided by the Supplier.

39. Contract Amendments

39.1 Subject to GCC Clause 32, no variation in or modification of the terms of the Contract shall be made except by written amendment signed by the parties.

40. Sub-Contracts

40.1 The Supplier shall consult the Procuring Agency in the event of subcontracting under this contract if not already specified in the Bid. Subcontracting shall not alter the Supplier's obligations.



Special Conditions of Contract

SECTION VIII. SPECIAL CONDITIONS OF CONTRACT

The following Special Conditions of Contract shall supplement the General Conditions of Contract. Whenever there is a conflict, the provisions herein shall prevail over those in the Conditions of Contract. The corresponding clause number of the GCC is indicated in parentheses.

Number of GC Clause

Amendments of, and Supplements to, Clauses in the General Conditions of Contract

Number of GC Clause 1

Definitions

The Procuring Agency is: **FEDERAL GOVERNMENT POLYCLINIC ISLAMABAD (FGPC-PGMI) (Federal Government Polyclinic (FGPC-PGMI)), Joint Executive Director FEDERAL GOVERNMENT POLYCLINIC (PGMI) G-6/2, ISLAMABAD, Islamabad Capital Territory**

The Supplier is:

The title of the subject procurement is: **2nd Re-Tender for Electro-Medical Equipment/s of Cardiology Department, under PSDP Project titled 'Strengthening and Up-Gradation of Cardiology and other Departments during the FY 2025-26 & 2026-27**

Number of GC Clause 3

Applicable/Governing Law:

The Contract shall be interpreted in accordance with the laws of Islamic Republic of Pakistan

Number of GC Clause 4

Language:

The language of the Contract, all correspondence and communications to be given, and all other documentation to be prepared and supplied under the Contract shall be in **English**.

Number of GC Clause 5

Notices:

The addresses for the notices are:

The Authority:

**FEDERAL GOVERNMENT POLYCLINIC ISLAMABAD (FGPC-PGMI) (Federal Government Polyclinic (FGPC-PGMI)), Joint Executive Director
FEDERAL GOVERNMENT POLYCLINIC (PGMI) G-6/2, ISLAMABAD, Islamabad Capital Territory
+92-333-550-9275
jed@fgpc.gov.pk**

Contractor/ Bidder:

[Name, address and telephone number].

The Contractor/ Bidder's Representative(s)

[Name, address, telephone number and e-mail address]

Number of GC Clause 7.1

The Authorized Representatives are:

For the Procuring Agency:

FEDERAL GOVERNMENT POLYCLINIC ISLAMABAD (FGPC-PGMI) (Federal Government Polyclinic (FGPC-PGMI)), Joint Executive Director
FEDERAL GOVERNMENT POLYCLINIC (PGMI) G-6/2, ISLAMABAD, Islamabad Capital Territory
+92-333-550-9275
jed@fgpc.gov.pk

For the Bidder:

Name:

Designation:

Address:

Number of GC Clause 8

Effectiveness of the contract

Number of GC Clause 9

Commencement of Contract:

Number of GC Clause 11.2

Expiration of Contract:

Number of GC Clause 15

Termination

In the event of termination of the contract due to any reason as already defined in the General Conditions of Contract, the Bidder shall be responsible for providing to the Authority the Goods till the time of alternate arrangements.

Number of GC Clause 17

Conflict of Interest:

The Procuring Agency reserves the right to determine on a case-by-case basis whether the Bidder should be disqualified from providing goods or services due to a conflict of a nature described in Clause GCC 17.

Number of GC Clause 22

Liquidated Damages

If the Bidder fails to provide services as required under the contract or in case of any data loss/data breach or any incident compromising the data security or other such failures related to any services, the Bidder shall pay to the Procuring Agency as Liquidated Damages at a rate of **0.10% to 2.00%** of the Contract value, in accordance with the extent of performance failure & the cost of investigating such incidents as judged by the Authority.

Number of GC Clause 23

Performance Guarantee:

Instrument of Performance Guarantee shall be in the Name of : **FEDERAL GOVERNMENT POLYCLINIC ISLAMABAD (FGPC-PGMI)**

The amount of performance guarantee shall be **3.00%** of the contract price in acceptable form of **Pay Order, Call at Deposit**

Number of GC Clause 32

Payment terms:

Payment will be made to the Bidder against the procured Goods and services according to the actual invoice or running bills submitted by the Bidder against the services provided within the time given in the conditions of the contract.

Number of GC Clause 33

Currency of Payment:

All the payment to be released to the contractor/Bidder shall be in Pakistani Rupees.

Number of GC Clause 34

Identifying Defects:

The Authority reserves the right at any time to inspect the premises of the provider to inspect the goods and monitor the goods being provided.

Inspections & Tests Requirements

For being Brand New, bearing relevant reference numbers of the equipment (Certificate from supplier)

For Physical Fitness having No Damages (Certificate from supplier)

For the Country of Origin as quoted by the Supplier (Certificate from manufacturer)

For conformance to specifications and performance parameters, through Prior to delivery inspection (Inspection Report by Procurement Committee / Inspection Team)

For successful operation at site after complete installation, testing and commissioning of the equipment (Installation, Testing and Commissioning Report by Procurement Committee / Inspection Team)

Delivery & Documents

Copies of the Supplier's invoice showing Goods' description, quantity, unit price, and total amount;

Original and two copies of the usual transport document (for example, a negotiable bill of lading, a non-negotiable sea waybill, an inland waterway document, an air waybill, a railway consignment note, a road consignment note, or a multimodal transport document) which the buyer may require to take the goods;

Copies of the packing list identifying contents of each package;

Manufacturer's or Supplier's Valid Warranty Certificate;

Inspection Certificate issued by the Nominated Inspection Agency (if any), and the Supplier's Factory Inspection Report;

Certificate of Origin.

The above documents would be required even if the equipment has already been imported and is available with the supplier ex-stock

Number of GC Clause 37

Following is the guidance for Dispute Resolution

(a) For Contracts to be entered with foreign Contractor/ Service Provider:

All disputes arising in connection with the present Contract shall be finally settled under the Rules of Conciliation and Arbitration of the International Chamber of Commerce by one or more arbitrators appointed in accordance with said Rules.

(b) For Contracts to be entered with nationals of Pakistan:

i. If any dispute of any kind whatsoever shall arise between the Authority and the Bidder in connection with or arising out of the Contract, including without prejudice to the generality of foregoing, any question regarding its existence, validity, termination and the execution of the Contract – whether during developing phase or after their completion and whether before or after the termination, abandonment or breach of the Contract – the parties shall seek to resolve any such dispute or difference by mutual diligent negotiations in good faith within 14 (fourteen) days following a notice sent by one Party to the other Party in this regard.

ii. At future of negotiation the dispute shall be resolved through mediation and mediator shall be appointed with the mutual consent of the both parties.

iii. At the event of failure of mediation to resolve the dispute relating to this contract such dispute shall finally be resolved through binding Arbitration in accordance with GCC clause 38.

iv. The cost of the mediation and arbitration shall be shared by the parties in equal proportion however the both parties shall bear their own costs and lawyer's fees regarding their own participation in the mediation and arbitration. However, the Arbitrator may make an award of costs upon the conclusion of the arbitration making any party to the dispute liable to pay the costs of another party to the dispute.

v. Arbitration proceedings as mentioned in the above clause regarding resolution of disputes may be commenced prior to, during or after completion of the contract.

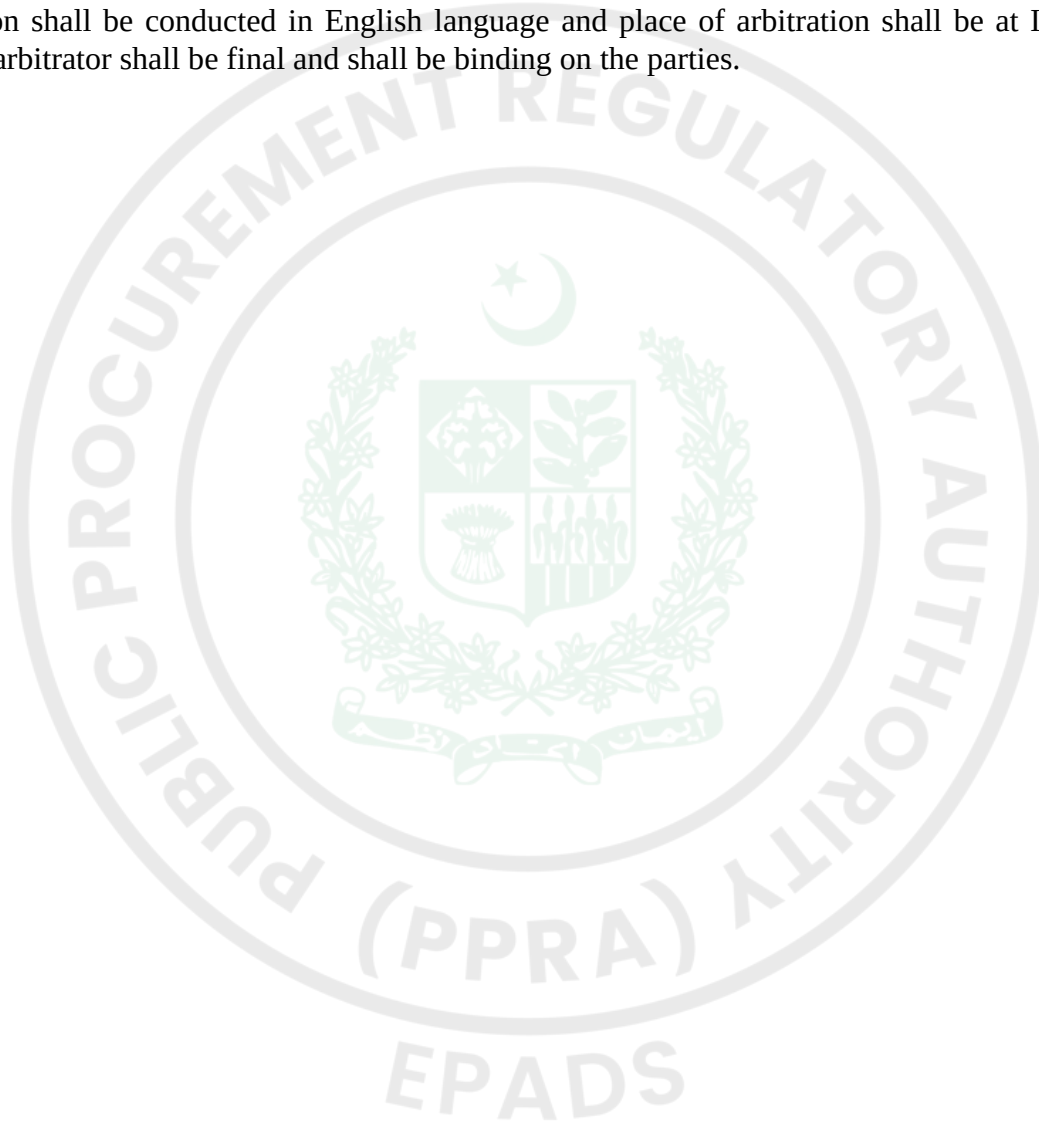
Notwithstanding any reference to the arbitration herein, the parties shall continue to perform their respective obligations under the Contract unless they otherwise agree that the Authority shall pay the Bidder any monies due to the Bidder.

Rules of procedure for arbitration proceedings:

Any dispute between the Authority and a Bidder who is a national of the Islamic Republic of Pakistan arising in connection with the present Contract shall be referred to adjudication or arbitration in accordance with the laws of the Islamic Republic of Pakistan including Arbitration Act 1940, however above provision shall prevail in referring the case to the Arbitrator.

Place of Arbitration and Award:

The arbitration shall be conducted in English language and place of arbitration shall be at Islamabad. The award of the arbitrator shall be final and shall be binding on the parties.





Bid Securing Declaration

Form 9: Bid Securing Declaration

Date: *[insert date (as day, month and year)]*

Bid No.: **P148647**

Alternative No.: **[insert identification No if this is a Bid for an alternative]**

To: **FEDERAL GOVERNMENT POLYCLINIC ISLAMABAD (FGPC-PGMI) (Federal Government Polyclinic (FGPC-PGMI)), Joint Executive Director FEDERAL GOVERNMENT POLYCLINIC (PGMI) G-6/2, ISLAMABAD, Islamabad Capital Territory**

We, the undersigned, declare that:

We understand that, according to your conditions, Bids must be supported by a Bid Securing Declaration.

We accept that we will be blacklisted and henceforth cross debarred for participating in respective category of public procurement proceedings for a period of (not more than) six months, if fail to abide with a bid securing declaration, however without indulging in corrupt and fraudulent practices, if we are in breach of our obligation(s) under the Bid conditions, because we:

1. have withdrawn or modified our Bid during the period of Bid Validity specified in the Form of Bid;
2. Disagreement to arithmetical correction made to the Bid price; or
3. having been notified of the acceptance of our Bid by the Procuring Agency during the period of Bid Validity, (i) failure to sign the contract if required by Procuring Agency to do so or (ii) fail or refuse to furnish the Performance Security or to comply with any other condition precedent to signing the contract specified in the Bidding Documents.

We understand this Bid Securing Declaration shall expire if we are not the successful

Bidder, upon the earlier of (i) our receipt of your notification to us of the name of the successful Bidder; or (ii) twenty-eight (28) days after the expiration of our Bid.

Signed: *[insert signature of person whose name and capacity are shown]* In the capacity of *[insert legal capacity of person signing the Bid Securing Declaration]*

Name: *[insert complete name of person signing the Bid Securing Declaration]*

Duly authorized to sign the Bid for and on behalf of: *[insert complete name of Bidder]*

Dated on day of , *[insert date of signing]*
Corporate Seal (where appropriate)



Contract Form

SECTION IX: CONTRACT FORMS

THIS AGREEMENT made the ____ day of _____ 20____ between **FEDERAL GOVERNMENT POLYCLINIC ISLAMABAD (FGPC-PGMI) (Federal Government Polyclinic (FGPC-PGMI)), Joint Executive Director FEDERAL GOVERNMENT POLYCLINIC (PGMI) G-6/2, ISLAMABAD, Islamabad Capital Territory**

(hereinafter called “the Procuring Agency”) of the one part and [name of Bidder] of [city and country of Bidder] (hereinafter called “the Bidder”) of the other part:

WHEREAS the Procuring Agency invited Bids for provision of goods, viz., **2nd Re-Tender for Electro-Medical Equipment/s of Cardiology Department, under PSDP Project titled ‘Strengthening and Up-Gradation of Cardiology and other Departments during the FY 2025-26 & 2026-27 (P148647)** and has accepted a Bids by the Bidder for the provision of Goods in the sum of [contract price in words and figures] (hereinafter called “the Contract Price”).

NOW THIS CONTRACT WITNESSETH AS FOLLOWS:

1. In this Contract words and expressions shall have the same meanings as are respectively assigned to them in the Conditions of Contract referred to.

2. The following documents shall be deemed to form and be read and construed as part of this Contract, In the event of any ambiguity or conflict between the Contract Documents listed below, the order of precedence shall be the order in which the Contract Documents are listed below:-

1. This form of Contract;
2. the Form of Bids and the Price Schedule submitted by the Bidder;
3. the Schedule of Requirements;
4. the Technical Specifications;
5. the Special Conditions of Contract;
6. the General Conditions of the Contract;
7. the Procuring Agency’s Letter of Acceptance; and
8. [add here: any other documents]

3. In consideration of the payments to be made by the Procuring Agency to the Bidder as hereinafter mentioned, the Bidder hereby covenants with the Procuring Agency to provide the Goods related services and to remedy defects therein in conformity in all respects with the provisions of the Contract.

4. The Procuring Agency hereby covenants to pay the Bidder in consideration of the provision of Goods and the remedying of defects therein, the Contract Price or such other sum as may become payable under the provisions of the contract at the times and in the manner prescribed by the contract.

IN WITNESS whereof the parties hereto have caused this Contract to be executed in accordance with their respective laws the day and year first above written.

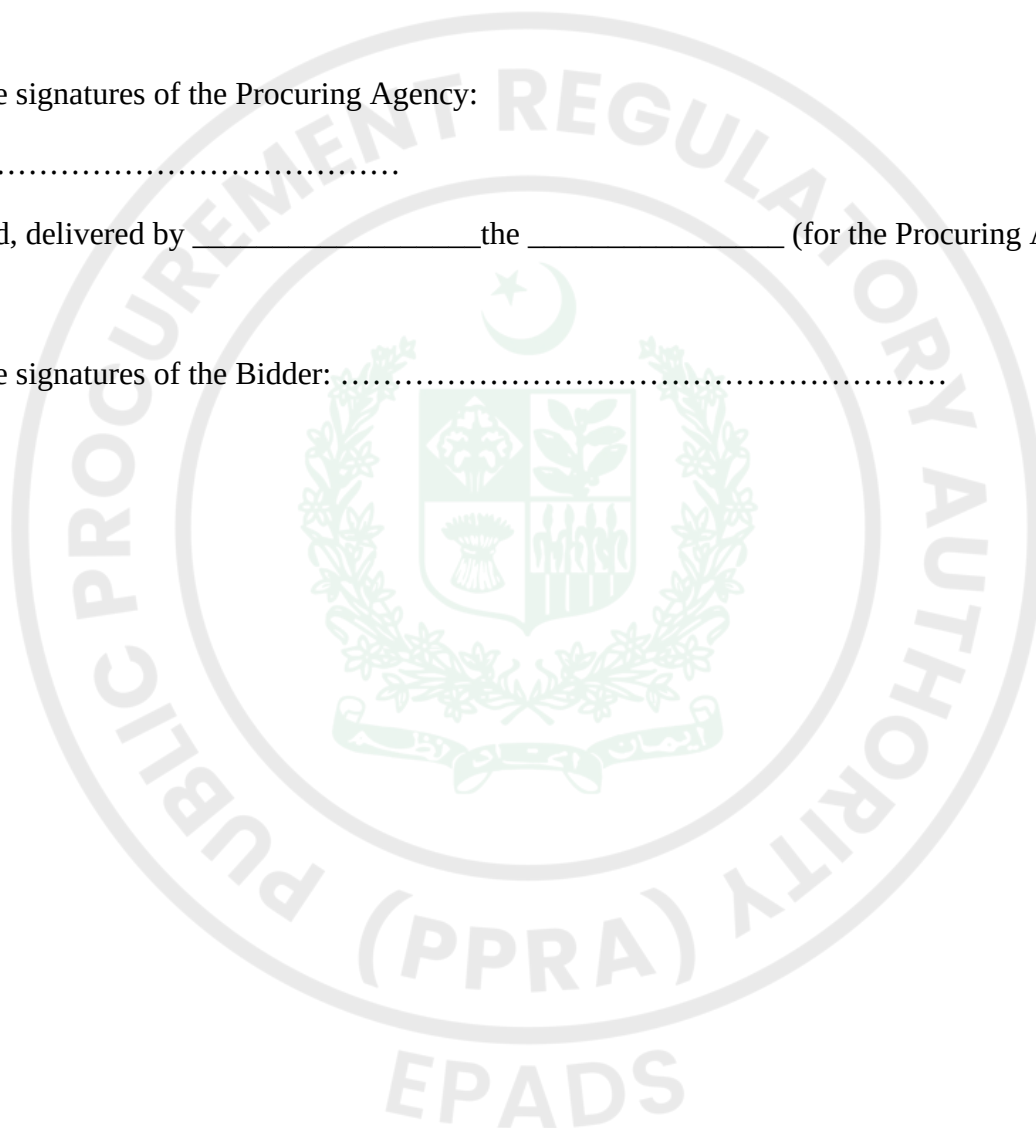
Signed, sealed, delivered by _____ the _____ (for the Procuring Agency)

Witness to the signatures of the Procuring Agency:

.....

Signed, sealed, delivered by _____ the _____ (for the Procuring Agency)

Witness to the signatures of the Bidder:





Integrity Pact

Integrity Pact

DECLARATION OF FEES, COMMISSION AND BROKERAGE ETC. PAYABLE BY THE SUPPLIERS OF GOODS, SERVICES & WORKS IN CONTRACTS WORTH RS.10.00 MILLION OR MORE

Contract

Number: Contract

Value: Contract Title:

Dated:

[Name of Supplier] hereby declares that it has not obtained or induced the procurement of any contract, right, interest, privilege or other obligation or benefit from Government of Pakistan or any administrative subdivision or agency thereof or any other entity owned or controlled by it (GoP) through any corrupt business practice.

Without limiting the generality of the foregoing [Name of Supplier] represents and warrants that it has fully declared the brokerage, commission, fee etc. paid or payable to anyone and not given or agreed to give and shall not give or agree to give to anyone within or outside Pakistan either directly or indirectly through any natural or juridical person, including its affiliate, agent, associate, broker, consultant, director, promoter, shareholder, sponsor or subsidiary, any commission, gratification, bribe, finder's fee or kickback, whether described as consultations fee or otherwise, with the object of obtaining or inducing the procurement of a contract, right, interest, privilege or other obligation or benefit in whatsoever form from GoP, except that which has been expressly declared pursuant hereto.

[Name of Supplier] certifies that it has made and will make full disclosure of all agreements and arrangements with all persons in respect of or related to the transaction with GoP and has not taken any action or will not take any action to circumvent the above declaration, representative or warranty.

[Name of Supplier] accepts full responsibility and strict liability for making and false declaration, not making full disclosure, misrepresenting fact or taking any action likely to defeat the purpose of this declaration, representation and warranty. It agrees that any contract, right interest, privilege or other obligation or benefit obtained or procured as aforesaid shall, without prejudice to any other right and remedies available to GoP under any law, contract or other instrument, be voidable at the option of GoP.

Notwithstanding any rights and remedies exercised by GoP in this regard, [Name of Supplier] agrees to indemnify GoP for any loss or damage incurred by it on account of its corrupt business practices and further pay compensation to GoP in an amount equivalent to ten times the sum of any commission, gratification, bribe, finder's fee or kickback given by [Name of Supplier] as aforesaid for the purpose of obtaining or inducing the procurement of any contract, right, interest, privilege or other obligation or benefit in whatsoever form from GoP.



Performance Guarantee Form

Performance Guarantee Form

To: **FEDERAL GOVERNMENT POLYCLINIC ISLAMABAD (FGPC-PGMI) (Federal Government Polyclinic (FGPC-PGMI)), Joint Executive Director FEDERAL GOVERNMENT POLYCLINIC (PGMI) G-6/2, ISLAMABAD, Islamabad Capital Territory**

WHEREAS *[name of Bidder]* (hereinafter called “the Bidder”) has undertaken, in pursuance of Contract No. *[reference number of the contract]* dated *[insert date]* for provision of Goods (hereinafter called “the Contract”).

AND WHEREAS it has been stipulated by you in the said Contract that the Bidder shall furnish you with a Bank Guarantee by a reputable bank for the sum specified therein as security for compliance with the Bidder’s performance obligations in accordance with the Contract.

AND WHEREAS we have agreed to give the Bidders guarantee:

THEREFORE, WE hereby affirm that we are Guarantors and responsible to you, on behalf of the Bidder, up to a total of *[amount of the guarantee in words and figures]*, and we undertake to pay you, upon your first written demand declaring the Bidder to be in default under the Contract and without cavil or argument, any sum or sums within the limits of *[amount of guarantee]* as aforesaid, without your needing to prove or to show grounds or reasons for your demand or the sum specified therein.

This guarantee is valid until the: *[insert date]*

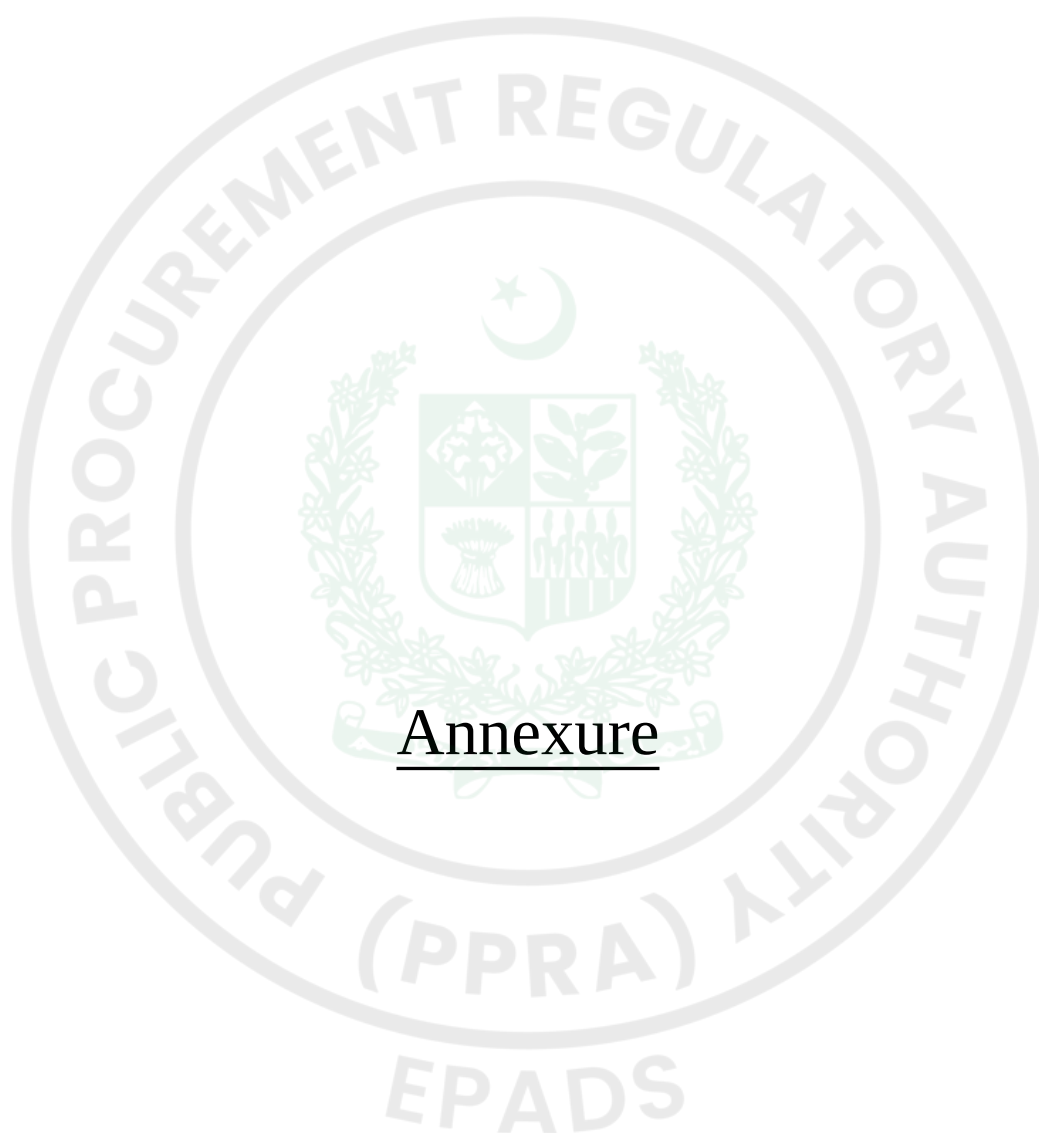
Signature and seal of the Guarantors

[name of bank or financial institution]

[address]

[date]





Annexure

instruction to bidder and general conditions

Information (Read-Only)

See Form Under Additional Forms and Documents: **instruction to bidder and general conditions** (page number: 118)

Specific Terms and conditions

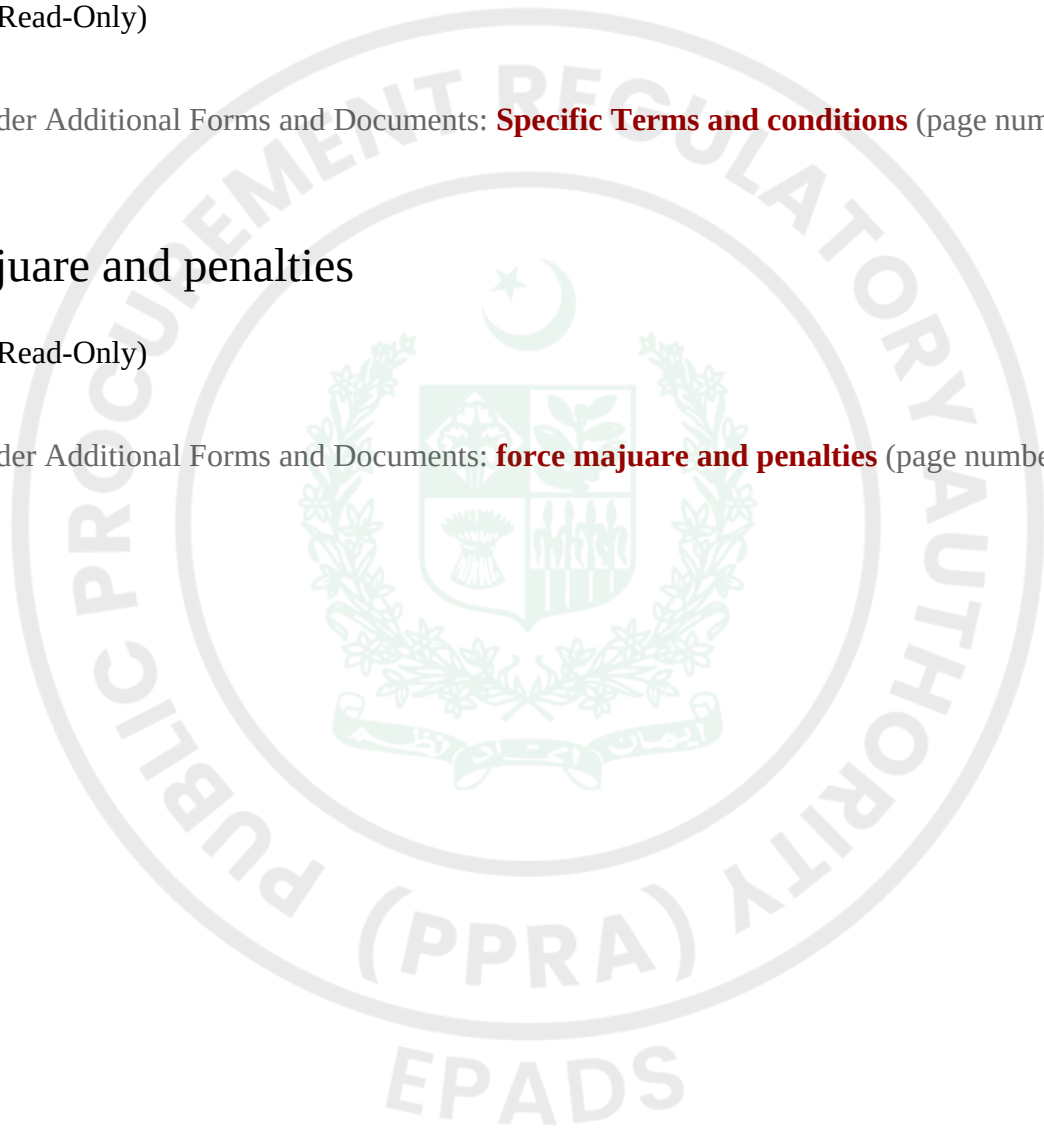
Information (Read-Only)

See Form Under Additional Forms and Documents: **Specific Terms and conditions** (page number: 119)

force majeure and penalties

Information (Read-Only)

See Form Under Additional Forms and Documents: **force majeure and penalties** (page number: 124)





Procurement Forms







Additional Forms and Documents

INSTRUCTIONS FOR FILLING OF THE CONTRACT DOCUMENTS.

1.	The Pre-Bid meeting will be held in the main auditorium of FGPC on scheduled date and time published in the advertisement/EPAD. Observations of those bidders will be considered who have uploaded their observations on EPAD.
2.	The participant firms are directed to submit their bids on-line through EPADS -V2.0 of PPRA
3.	The competent authority reserves the right to cancel tender process under 33(1) of PPRA Rules.
4.	All the documents attached must be attested/signed & stamped by owner/authorized person of firm.
5.	Participant bidders are directed to upload copies of judicial papers and Pay Order/CDR on EPADS-V2.0. Original Stamp paper along with Original CDR must be submitted to FGPC before or at the date and time of Bid opening by the bidder.

A. GENERAL CONDITIONS.

1.	This contract is under PSDP for the FY 2025-26 & 2026-2027.
2.	The successful bidder will be awarded contract after submission of 3% performance guarantee, integrity pact + contract agreement and safe guard certificate on legally Stamp paper / E-Stamp (at least Rs.300 or more) duly attested by Notary public or Oath Commissioner that he agrees to supply the items on approved rates within the stipulated period in accordance with tender TORs.
3.	The bidder must provide Technical brochures/Literature, brand, country of origin, country of manufacturing of quoted item/material with the bid.
4.	The Bidder/supplier will quote rates inclusive of all applicable taxes.
5.	All items will be received on FOR basis, FGPC Islamabad.
6.	Bid validity is 120 days from the date of opening of the tender.
7.	The deliveries must be made according to supply orders.
8.	The quantity mentioned in the bidding document is tentative/estimate in nature. However the same can be increased or decreased according to the need of the institution and available funds.
9.	In case of any failure in supply within prescribed period, the reminder for the period of 15 days will be issued. Failing to supply after reminder, penalty will be imposed according to tender TOR's.
10.	Before awarding contract, letter of intent will be issued to firms.
11.	The bidder should apply through online portal of PPRA, EPADS Version.2
12.	In case of any dispute the matter will be dealt according to PPRA rules.
13.	The Executive Director FGPC reserves the right to accept or reject any tender/all tenders under PPRA Rules.
14.	The owner of the firm or authorized person must sign/validate bid documents including undertakings.

. SPECIFIC TERMS & CONDITIONS AND RESPONSIBILITIES.

1.	The bidder must provide Technical Brochure/literatures and technical datasheet approved/provided by original manufacturer of quoted item/equipment along with its model and number which must be currently listed on the official website of the manufacturer.
2.	The product must comply with certifications specified in the specifications of each item or package. In case of local instruments or in instruments where certifications are not applicable, manufacturer ISO certification must be provided.
3.	The bid must comply with technical specifications of the quoted single item or complete package as published in the tender. Any deviation from the specifications of main equipment may result in straightaway rejection. However minor deviation in the specifications will be acceptable subject to the approval of procurement committee.
4.	The bidder will be responsible for onsite demonstration or on video demonstration of quoted item on their own expense, if desired by procurement committee.
5.	The bid/documents uploaded by the bidder on EPADS version.2 will be final and should be accepted only.
6.	Warranty period: The bidder must submit Undertaking/declaration on legally Stamp paper / E-Stamp duly attested by Notary public or Oath Commissioner regarding required warranty period as per bidding document of equipment and also declare the availability of parts and accessories for the period defined in bidding documents after installation of equipment. The bidder also undertake that they will be responsible for Free of cost replacement of parts and repair/service during warranty period.
7.	The warranty shall cover all parts, accessories, probe, sensor, cables, batteries, electronic boards, displays, software, labors, travel, transportation, calibration, preventive maintenance and corrective maintenance required to keep equipment fully functional. No charges shall be payable by procuring agency during the warranty period except for damage conclusively established as a resulting from misuse or external cause.
8.	The warranty shall automatically be extended by the total period during which the equipment remain nonfunctional or unavailable for clinical use due to fault covered under warranty.
9.	The bidder and its manufacturer shall provide an undertaking confirming the availability of genuine spare parts, accessories, consumable, software support and technical services for at least 10 years from the date of installation of equipment.
10	The supplier shall notify the procuring agency at least three months for major equipment's and one month for minor equipment before discontinuation of any model, spare parts, accessories, consumable, software platform or technical support. A compatible and approved alternative shall be offered without adversely affecting the installed equipment performance.
11	The payment will be made after installation and satisfactory report of the end-user/head of department duly verified by procuring agency.
12	The approved rates will be final and there will be no increase in rates and no escalation clause.
13	All products/items should be supplied in proper packing and in sealed position.
14	The freight charges will be borne by the suppliers.

15	Package should be worthy of transportation by Road, Sea, or mail as to ensure its contents being free from loss of damage due to faulty packing.
16	Commencement of warranty: The warranty for each item shall commence from the date of successful installation, commissioning, acceptance testing, completion of initial training, and issuance of the Final Acceptance Certificate by the authorized end-user, whichever occurs last.
17	Installation and commissioning: Installation and commissioning shall be performed by qualified, manufacturer-trained technical personnel. All installation materials, connectors, stabilizers, UPS units, mounting accessories, software, licenses, calibration tools, and other items required for safe and complete operation shall be included unless expressly excluded in the specifications.
18	training: The successful bidder shall provide structured operational, clinical-application, cleaning, basic troubleshooting, and user-maintenance training during a period of at least six weeks after installation at the premises of FGPC. Training shall cover all relevant shifts and nominated end-users without additional cost.
19	Training documentation: The supplier shall provide a training plan, attendance record, training certificates, competency checklist, and soft and hard copies of user manuals. Refresher training shall be provided at least once during each warranty year and whenever major software or hardware changes are introduced.
20	After-sales infrastructure: The bidder shall maintain, directly or through an authorized service arrangement, an adequately equipped after-sales service facility capable of meeting the response and repair times stated in the bidding documents. The Procuring Agency may inspect the facility before award or during the contract.
21	Personnel for each tender item: The bidder shall nominate for every quoted item or lot: a. At least one manufacturer-trained service engineer or qualified biomedical engineer; b. At least one suitably qualified backup engineer; and c. Where applicable, one clinical application specialist.
22	Evidence of personnel: Names, qualifications, experience, contact details, CVs, and valid manufacturer training certificates of the nominated personnel shall be submitted with the technical bid. A mere undertaking to arrange personnel after award shall not be sufficient.
23	Availability of personnel: The nominated service personnel shall remain available throughout installation, commissioning, training, and the entire warranty period. Replacement of nominated personnel shall require prior written approval of the Procuring Agency, and the substitute shall possess equivalent or higher qualifications.
24	Service contact: The supplier shall provide a twenty-four-hour complaint telephone number, email address, and escalation contact. Every complaint shall be assigned a unique service-call number and acknowledged electronically.

25	Response time: a. Telephone or remote response: within 30 minutes; b. On-site attendance for defibrillators, cardiac monitors, ICU equipment, and other critical items like Dialysis machine, ventilator, Anesthesia machine: within four hours; c. On-site attendance for non-critical equipment: within eight working hours; d. If the service centre is outside Islamabad or Rawalpindi, the bidder shall demonstrate how these response times will be achieved.
26	Repair time: Critical equipment shall be restored to full working condition within twenty-four hours and non-critical equipment within forty-eight hours after notification.
27	Standby equipment: If repair cannot be completed within the prescribed period, the supplier shall provide, at no cost, equivalent or better standby equipment within the next twenty-four hours for critical equipment and within seventy-two hours for other equipment.
28	Equipment uptime: The supplier shall maintain minimum operational uptime of 95% for equipment, calculated annually. Downtime attributable to hospital infrastructure, misuse, or force majeure shall be excluded only when properly documented and accepted by the Procuring Agency.
29	Planned preventive maintenance: Preventive maintenance shall be performed according to the manufacturer's recommended schedule or at least once every quarter or as per bidding documents, whichever provides greater protection. Maintenance shall be coordinated with the end-user to avoid disruption of patient care.
30	Calibration and safety testing: Preventive maintenance shall include calibration, performance verification, electrical-safety testing, functional checks, cleaning, lubrication, software checks, and replacement of parts due for routine replacement. Traceable calibration certificates and service reports shall be provided.
31	Service records: A service logbook shall be maintained for each item . Every service report shall record the complaint, date and time reported, response time, diagnosis, action taken, parts replaced, calibration status, downtime, and signatures of the service engineer and end-user duly verified by Bio-medical Department, FGPC
32	Genuine parts: Only new, genuine, manufacturer-approved parts and accessories shall be used. Repaired, salvaged, refurbished, locally substituted, or counterfeit parts shall not be accepted without prior written approval of the Procuring Agency.
33	Software and firmware: All software, licences, passwords, service access, firmware updates, security patches, and upgrades required for the contracted functionality shall be included during the warranty period. No equipment shall become partly or wholly unusable due to expiry of an undisclosed licence or subscription.
34	Cyber security and data protection: For network-enabled equipment, the supplier shall provide available security updates, vulnerability notifications, secure configuration guidance, data-backup procedures, and protection against unauthorized access throughout the warranty period.

35	Safety alerts and recalls: The supplier shall notify the Procuring Agency promptly of any manufacturer safety alert, field safety corrective action, or product recall and shall perform all required corrections or replacements without cost.
36	Acceptance testing: The end-user may verify quantity, configuration, accessories, performance, electrical safety, calibration, connectivity, and compliance with the contracted specifications. Defective, incomplete, substituted, or non-compliant equipment may be rejected.
37	Documents required before acceptance: The supplier shall provide the following for each item: a. Manufacturer's warranty certificate; b. Installation and commissioning report; c. Acceptance-testing and calibration certificates; d. User and service manuals; e. Complete inventory of supplied accessories; f. Recommended spare-parts and consumables list with part numbers; g. Preventive-maintenance schedule; h. Service and escalation contact matrix; and i. Training-completion records.
38	Supplier's continuing responsibility: Change or termination of dealership, distributorship, or manufacturer authorization shall not relieve the successful bidder of contractual, warranty, maintenance, spare-parts, or after-sales obligations.
39	Subcontracting: After-sales service shall not be subcontracted without prior written approval of the Procuring Agency. Approved subcontracting shall not reduce the successful bidder's responsibility or liability.
40	Failure to provide services: If the supplier fails to respond, repair, replace, or provide standby equipment within the stipulated time, the Procuring Agency may arrange repair or replacement from another source at the supplier's risk and cost, without prejudice to other contractual remedies.
41	Performance and warranty security: The performance guarantee or separate warranty security, in the amount and form specified in the Special Conditions of Contract, shall remain valid until completion of the relevant contractual and warranty obligations.
42	Service-level penalties: Failure to meet response time, repair time, preventive-maintenance, uptime, training, or standby-equipment requirements shall attract the lack-of-performance penalty or liquidated damages specified in the Special Conditions of Contract. Such deductions shall not relieve the supplier of its obligation to repair or replace the equipment.
43	Post-warranty support: The bidder shall submit separately itemized rates for comprehensive and non-comprehensive annual maintenance for at least five years after expiry of the warranty. These rates and their permitted annual adjustment mechanism shall be declared in the financial bid.
44	Inspection and verification: The Procuring Agency may verify the bidder's service facility, spare-parts inventory, tools, test equipment, manufacturer authorization, and availability of nominated personnel before award and at any time during contract performance (If required/necessary) .

45	Declaration of no conflict of interest in accordance with the bidding documents, including disclosure of related entities participating in the same procurement.
46	Compliance sheet of specifications/bidding documents for each quoted instrument will be given.
47.	The sole agent of Manufacturer will be responsible to provide valid authorization letter from manufacturer duly attested by the relevant Consulate general of Pakistan before supply of the approved product/item.



FORCE MAJEURE.

01.	For the purposes of this clause Force Majeure means an act of God or an event beyond the control of the supplier and not involving the supplier's fault or negligence directly or indirectly purporting to miss planning, miss management and/or lack of foresight to handle the situation. Such events may include but are not restricted to acts of the purchaser in its sovereign capacity, wars or revolutions, fires, floods, earthquakes, epidemics, quarantine restrictions and freight embargoes. If a Force Majeure situation arises, the supplier shall promptly notify the purchaser in writing with sufficient and valid evidence of such condition and the cause thereof. The Force Majeure Committee will examine the pros and cons of the case and all reasonable alternative means for completion of supply order under this Contract and will submit its recommendations to the competent authority. However, unless otherwise directed by the purchaser in writing, the supplier shall continue to perform its obligations under the Contract as far as is reasonably practical and shall seek reasonable alternative means for performance not prevented by the Force Majeure event.
	<u>Non-compliance of supply order.</u> If the supply order is not completed within stipulated period one or more of the following penalties can be applied against the firm according to the gravity of situation. - A penalty @ 0.1% per day against the amount of pending item (s) of the supply order shall be imposed maximum upto 2% of the supply order. - Risk purchase will be made at the cost of supplier and amount will be deducted from the bills of supplier. - The contract of the supplier will be treated as cancelled and the order for supply of item (s) will be placed to the next lowest. - Total or partial earnest money (CDR) will be forfeited. - The firm will be debarred for business at FGPC for that year. - The firm will be blacklisted. - In case of any complaint about the penalty(s) imposed on the firm, the firm can appeal against the decision in the Grievance Redressal Committee within 15 days of the issue of penalty(s) letter. After the lapse of this period no appeal will be entertained.