

Standard Bidding Document

SUPPLY OF BLOOD BAGS, ANTI SERAS, KITS AND DISPOSABLE
ITEMS, TO BE USED IN BLOOD BANK AT PIMS ON ANNUAL
CONTRACT BASIS, FOR THE FY 2026-2027
(Goods)

National

Single Stage-Two Envelope



July 04, 2026

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REQUEST FOR BIDS

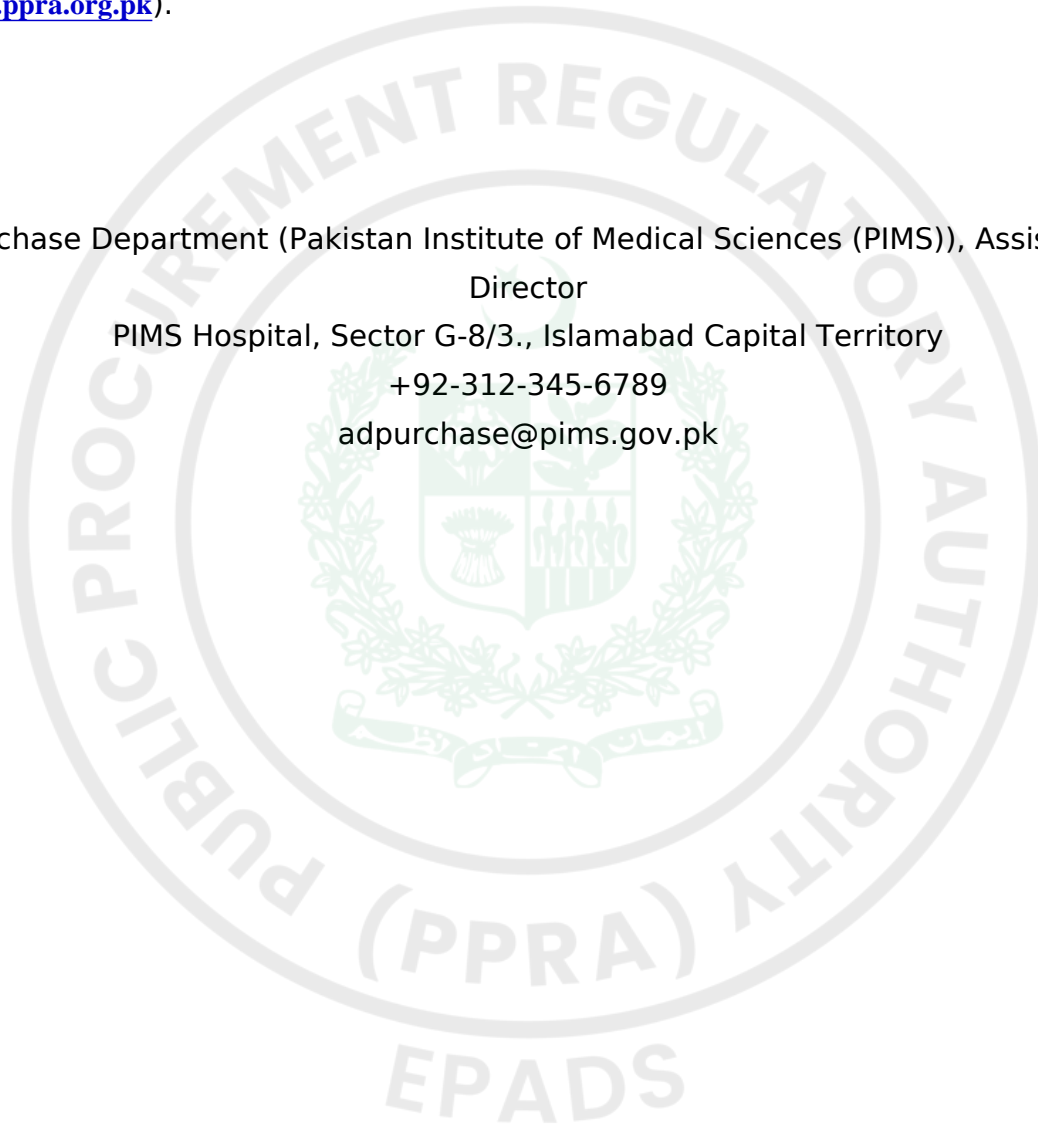
PROCUREMENT OF GOODS

1. The **Purchase Department (Pakistan Institute of Medical Sciences (PIMS))** has reserved Funds for the procurement planned for FY **2026-27**. The **Purchase Department (Pakistan Institute of Medical Sciences (PIMS))** intends to apply part of the proceeds of this Fund to cover eligible payments under the contract for the "**SUPPLY OF BLOOD BAGS, ANTI SERAS, KITS AND DISPOSABLE ITEMS, TO BE USED IN BLOOD BANK AT PIMS ON ANNUAL CONTRACT BASIS, FOR THE FY 2026-2027**" with the reference of "**P45305**"
2. The **Purchase Department (Pakistan Institute of Medical Sciences (PIMS))** 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, 2004 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 amounting 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/45305>** 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 **Thursday, July 23, 2026 11:00 AM**. E-bids will be opened using **EPADS v2.0** on the

same day at **Thursday, July 23, 2026 11:30 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 48 of Public Procurement Rules, 2004 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.org.pk).

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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 for 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-1 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.

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

3.3 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.

3.4 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).

3.5 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.

3.6 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.

3.7 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 performance failure or due to breach of Bid securing declaration.

3.8 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.

3.9 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 9.1** 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

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 his 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 three (3) 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 Copies of 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**.

9.3 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.

9.4 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.

1. Documentary evidence established in accordance with ITB that the Bidder is eligible and/or qualified for the subject bidding process;
2. Documentary evidence establish 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;
3. Documentary evidence establish that the goods and related services to be supplied by the Bidder are eligible goods and services, and conform to the Bidding Documents;
4. Bid security or Bid Securing Declaration furnished in accordance with **ITB 18**.

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 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 Bids

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

15. Bids Prices

15.1 The Bids 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 Items not listed in the Price Schedule shall be assumed not to be included in the Bid, and provided that the Bid is still substantially responsive in their absence or due to their nominal nature, the corresponding average price of the respective item(s) of the remaining substantially responsive Bidder(s) shall be construed to be the price of those missing item(s).

15.4 The Bid price to be quoted in the Form of Bid in accordance with **ITB 14.1** 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 30 (2) of the Public Procurement Rules, 2004.

17. Bids Validity Period

17.1 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.

17.2 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 only once through **EPADS v2.0**, for the period not more than the period of initial bid validity. The Bid Security provided under **ITB 18** 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 25 of the Public Procurement Rules, 2004.

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, and it shall be a Bank Draft in the name of the Procuring Agency and valid for twenty-eight (28) days beyond the end of the validity of the Bid. This shall also apply if the period

for Bids/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** 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 Bids 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 document, 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. the withdrawal of the Bids prior to the deadline for the submission of Bids, unless the Bids documents stipulate that no such withdrawal is permitted.

18.8 The successful Bidder's Bids 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 17.2**; 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. Bids requested to be withdrawn, shall be returned unopened to the Bidders 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

21.1 The Technical and Financial Bids if required to submitted, shall be submitted on **EPADS v2.0**.

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 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 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 fail 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 on 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 Evaluation Committee.

24. Clarification of Bids

24.1 To assist in the examination, evaluation and comparison of Bids of the Bidders, the Procuring Agency 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 Bids:

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 Bids 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 Procuring Agency 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 Procuring Agency'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.

25.3 If a Bids is not substantially responsive, it will be rejected by the Procuring Agency and may not subsequently be evaluated for complete technical responsiveness.

26. Examination of Terms and Conditions; Technical Evaluation

26.1 The Procuring Agency 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 Procuring Agency 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 Procuring Agency determines that the Bid is not substantially responsive in accordance with **ITB 25.2**, 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 prices 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 Procuring Agency 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 Bids, 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 Procuring Agency 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 To facilitate evaluation and comparison, the Procuring Agency will convert all Bids prices expressed in the amounts in various currencies in which the Bids prices are payable. For the purposes of comparison of bids quoted in different currencies, the price shall be converted into a single currency specified in the bidding documents. The rate of exchange shall be the selling rate prevailing on the date of opening of financial bids specified in the bidding documents, in accordance with weighted average customer exchange rates list issued by the State Bank of Pakistan on that day.

29. Evaluation of Bids

29.1 The Bids, quotations, or proposals shall be evaluated by the respective evaluation committees as per evaluation criteria described in the Bidding Documents in accordance with Rule 29 and 30 of the Public Procurement Rules, 2004.

1. 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.

2. 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.

3. 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.

29.2 In case of tie of bids, the bidders shall be provided an opportunity to offer their best and final monetary offer through EPADS v2.0. However, in no case the rates shall be higher than the original financial bids.

30. Domestic Preference

30.1 The procuring agency shall evaluate and compare bids, allow for preference to domestic bidders, while competing with the international bidders in accordance with the policies of 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 In case where the Procuring Agency adopts the Cost Based Evaluation Technique and, the Bid with the lowest evaluated price from amongst those which are eligible, compliant and substantially responsive shall be the Successful Bid.

31.3 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.

31.4 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)(h) of the Public Procurement Rules, 2004.

32. Abnormally Low Financial Bids

32.1 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.

32.2 The Procuring Agency may reject an Abnormally low financial bids.

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

1. Comparing the Bids price with the cost estimate;
2. Comparing the Bids price with the Bids 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.

32.4 The Procuring Agency will determine to its satisfaction whether the Bidder that is selected as having submitted the successful bid is qualified to perform the contract satisfactorily.

32.5 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.

32.6 Procuring Agency 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.

32.7 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 Bids, in which event the Procuring Agency 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 Most Advantageous Bidder.

34. Negotiations

34.1 The procuring agency shall not engage in negotiations with respect to scope and price with the bidder except when the procuring agency conducts a procurement using direct **or negotiated** contracting or a request for proposals with evaluation based on quality alone.

34.2 The procuring agency may negotiate with the most advantageous bid with a view to streamline the work or task execution, at the time of contract finalization on methodology, work plan, staffing, finalizing payment arrangements, delivery arrangements, minor amendments to the special conditions of the contract.

35. Procuring Agency 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 33 of the Public Procurement Rules, 2004.

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

36.1 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.

37. Notification of Award

37.1 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 35

of the Public Procurement Rules, 2004.

37.2 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).

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

38. Signing of Contract

38.1 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.

38.2 Immediately after the Redressal of grievance by the GRC (if any), mandatory standstill period in accordance with Rule 35 of the Public Procurement Rules, 2004 and **after fulfillment of all condition's precedent** of the Contract Form, the successful Bidder and the Procuring Agency shall sign the Contract.

39. Corrupt & Fraudulent Practices

39.1 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 to engage in any corrupt and fraudulent practices.

F. Grievance Redressal & Complaint Review Mechanism

40. Constitution of Grievance Redressal

40.1 The Grievance Redressal Committee shall address the grievance, if any submitted by any party, including the bidder, in accordance with Rule 48 of the Public Procurement Rules, 2004 to be read with Redressal of Grievances Regulations, 2021.

40.2 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 48 of the Public Procurement Rules, 2004 to be read with Redressal of Grievances Regulations, 2021.

G. Mechanism of Blacklisting

41. Mechanism of Blacklisting

41.1 The Procuring Agency shall initiate blacklisting proceedings against any bidder, supplier, or contractor in accordance with the Mechanism for Blacklisting Regulations, 2024, read with Rule 19 of the Public Procurement Rules, 2004.

41.2 The blacklisted/debarred bidder may file the review petition before the Authority in accordance with Rule 19 of the Public Procurement Rules, 2004 to be read with Procedure of filing and disposal of Review Petitions Regulations, 2021.





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: **Purchase Department (Pakistan Institute of Medical Sciences (PIMS))**

The subject of procurement is: **SUPPLY OF BLOOD BAGS, ANTI SERAS, KITS AND DISPOSABLE ITEMS, TO BE USED IN BLOOD BANK AT PIMS ON ANNUAL CONTRACT BASIS, FOR THE FY 2026-2027**

Expected commencement date: **Monday, August 31, 2026**

BDS Clause Number 2

ITB Number 2.1

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

Name and identification number of the Contract: **P45305**

BDS Clause Number 3

ITB Clause Number 3.1

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 Number 8.1

The Bidders may seek clarifications through **EPADS v2.0** : Clarification Date: Wednesday, July 8, 2026

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. Copy of Receipt of Purchase of Tender Documents.
2. FBR online Active Tax Payer List, Valid National Income Tax Number (Enclose copy of NTN Certificate), attach copies. Valid and updated document related to product registration by DRAP are required.
3. Manufacturer, Importer, authorized Distributors / Sole Authorized Distributors may qualify if they meet the criteria, attach copies.
4. Scanned copies of Name & Designation, Specimen Signature of Authorized / Focal Person with Cell No / Whatsapp No and Email address on official firm's letterhead pad.
5. The vendors/firms shall submit an undertaking on stamp paper of at least Rs.100/- by hand to the Purchase Department PIMS and shall also upload the same on the E-PADs, stating therein that the firm is not currently blacklisted / Penalized by any Govt. Deptt, / Hospitals/ International Agencies and NGO's.
6. The vendors/firms shall submit an undertaking on stamp paper of at least Rs.100/- by hand to the Purchase Department PIMS, and shall also upload the same on the E-PADs, stating therein that none of the batch of the product being quoted has been declared spurious by any laboratory anywhere in Pakistan.
7. The vendors/firms shall submit an undertaking on stamp paper of at least Rs.100/- by hand to the Purchase Department PIMS and shall also upload the

same on the E-PADs, stating therein that that the firm has agreed upon all the terms and conditions of the SBDs and upon awarding the contract, the firm will supply item(s) regularly on approved rates for the contract period.

8. The vendors/firms shall submit an undertaking on stamp paper of at least Rs.100/- by hand to the Purchase Department PIMS, and shall also upload the same on the E-PADs, stating therein that the products (imported, Local) for which the bidders intend to bid shall be made freely available for making the stock available in time / for the period as mentioned in TOR of tender.

BDS Clause Number 6

ITB Number 11.1

Items/Lots and threere 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: **180 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**

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

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;

PIMS Hospital, Sector G-8/3., 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: **Thursday, July 23, 2026 11:00 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 : **Thursday**

Date: **Thursday, July 23, 2026**

Time : **11:30 AM**

BDS Clause Number 14

ITB Number 31.1

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.

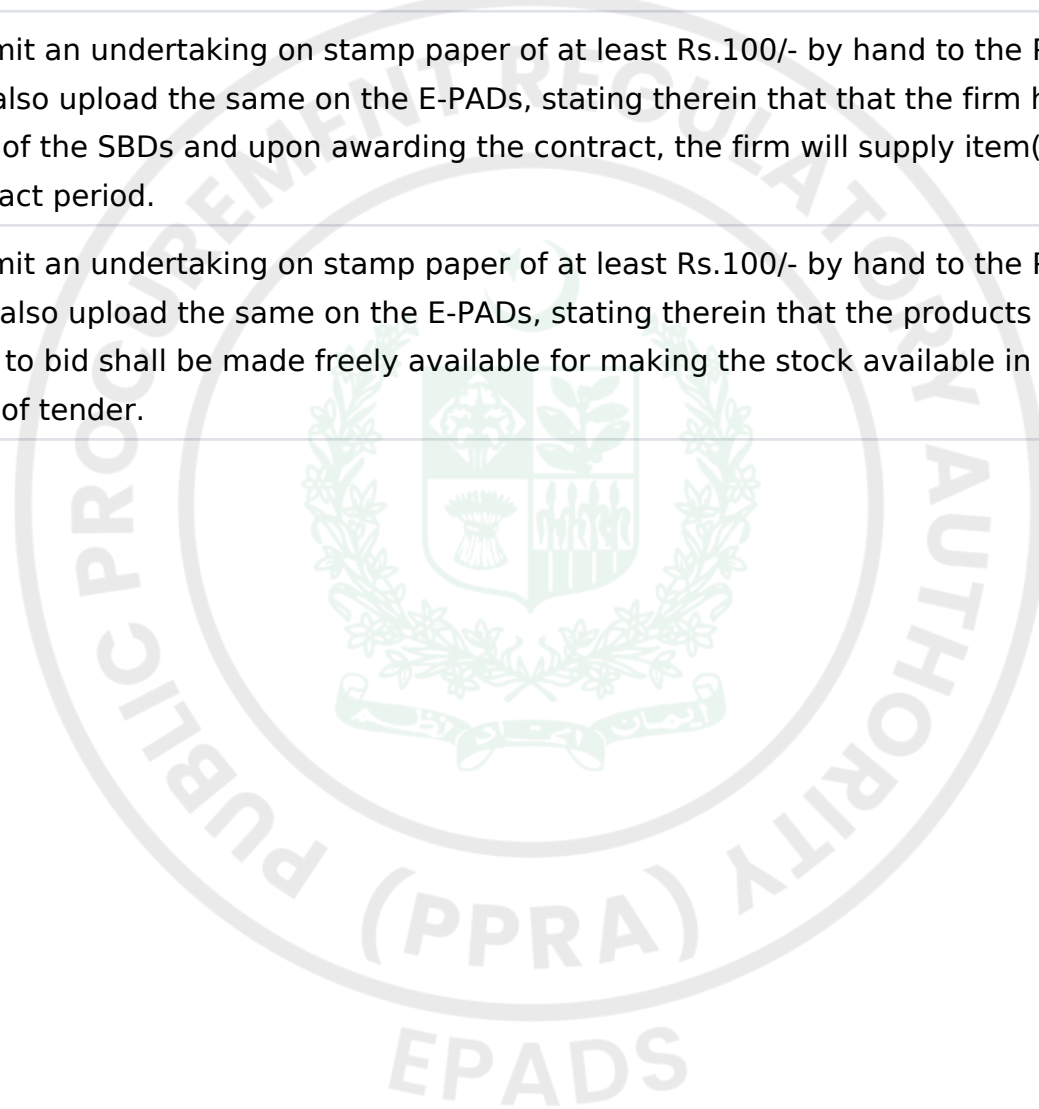
Arbitrator shall be appointed by mutual consent of the both parties.

Eligibility Criteria

Bidder's Type	Required Registration
Any	NADRA CITIZENSHIP (CNIC/NICOP) FBR (NTN) FBR (GSTN)

Eligibility Criteria	Document
Copy of Receipt of Purchase of Tender Documents.	Yes
FBR online Active Tax Payer List, Valid National Income Tax Number (Enclose copy of NTN Certificate), attach copies. Valid and updated document related to product registration by DRAP are required.	Yes
Manufacturer, Importer, authorized Distributors / Sole Authorized Distributors may qualify if they meet the criteria, attach copies.	Yes
Scanned copies of Name & Designation, Specimen Signature of Authorized / Focal Person with Cell No / Whatsapp No and Email address on official firm's letterhead pad.	Yes
The vendors/firms shall submit an undertaking on stamp paper of at least Rs.100/- by hand to the Purchase Department PIMS and shall also upload the same on the E-PADs, stating therein that the firm is not currently blacklisted / Penalized by any Govt. Deptt, / Hospitals/ International Agencies and NGO's.	Yes

<p>The vendors/firms shall submit an undertaking on stamp paper of at least Rs.100/- by hand to the Purchase Department PIMS, and shall also upload the same on the E-PADs, stating therein that none of the batch of the product being quoted has been declared spurious by any laboratory anywhere in Pakistan.</p>	<p>Yes</p>
<p>The vendors/firms shall submit an undertaking on stamp paper of at least Rs.100/- by hand to the Purchase Department PIMS and shall also upload the same on the E-PADs, stating therein that that the firm has agreed upon all the terms and conditions of the SBDs and upon awarding the contract, the firm will supply item(s) regularly on approved rates for the contract period.</p>	<p>Yes</p>
<p>The vendors/firms shall submit an undertaking on stamp paper of at least Rs.100/- by hand to the Purchase Department PIMS, and shall also upload the same on the E-PADs, stating therein that the products (imported, Local) for which the bidders intend to bid shall be made freely available for making the stock available in time / for the period as mentioned in TOR of tender.</p>	<p>Yes</p>



Evaluation Criteria

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

Least Cost Based Selection (LCBS)

Technical Marks	100
Passing Marks	70
Technical Evaluation Criteria	
The firms /Suppliers will provide the Audited Financial Statements for the last 02-years, (Annual Turnover 20 million) (Qualitative)(Doc Required)	50
More than 20 million (50)	
10 million to 20 million (25)	
Less than 10 million (10)	

Proof of experience for supply of Lab Kits, Chemicals, Reagents & Disposable Items ETC, to be used in Pathology Department and Blood Bank. (Attach documentary evidence i.e. experience certificate, copy of award letter etc (Qualitative))(Doc Required)

50

More than 03 years **(50)**

02 years to 03 years **(25)**

01 years to 02 years **(10)**



Items/Lots

Items Without Lots :

Item	UNSPSC	Delivery Schedule	Quantity	Bid Security	Sample Quantity	Manufacturer / Dealer Authorization
Blood Collecting Bags (Single) without Set)	Blood unit collection bags	Address: PIMS Hospital, Sector G-8/3., Islamabad Capital Territory Schedule: As per Supply Order(s) Quantity: 500/Qty	500/Qty	2083 PKR	10	Any
Blood Collecting Bags with diversion pouch triple without set	Blood unit collection bags	Address: PIMS Hospital, Sector G-8/3., Islamabad Capital Territory Schedule: As per Supply Order(s) Quantity: 18000/Qty	18000/Qty	197939 PKR	10	Any

Item	UNSPSC	Delivery Schedule	Quantity	Bid Security	Sample Quantity	Manufacturer / Dealer Authorization
Blood Collecting Bags without diversion pouch (Triple without Set)	Blood unit collection bags	Address: PIMS Hospital, Sector G-8/3., Islamabad Capital Territory Schedule: As per Supply Order(s) Quantity: 18000/Qty	18000/Qty	197939 PKR	10	Any
Blood Collecting Bags with diversion pouch (Quadruple without Set)	Blood unit collection bags	Address: PIMS Hospital, Sector G-8/3., Islamabad Capital Territory Schedule: As per Supply Order(s) Quantity: 600/Qty	600/Qty	8980 PKR	10	Any
Blood Collecting Bags without diversion pouch (Quadruple without Set)	Blood unit collection bags	Address: PIMS Hospital, Sector G-8/3., Islamabad Capital Territory Schedule: As per Supply Order(s) Quantity: 600/Qty	600/Qty	8980 PKR	10	Any

Item	UNSPSC	Delivery Schedule	Quantity	Bid Security	Sample Quantity	Manufacturer / Dealer Authorization
Blood Transfusion Set	Blood transfusion administration kits	Address: PIMS Hospital, Sector G-8/3., Islamabad Capital Territory Schedule: As per Supply Order(s) Quantity: 40000/Qty	40000/Qty	170000 PKR	100	Any
Blood Grouping Sera Anti-A	Blood bank reagents or solutions	Address: PIMS Hospital, Sector G-8/3., Islamabad Capital Territory Schedule: As per Supply Order(s) Quantity: 1600/Qty	1600/Qty	20800 PKR	1	Any
Blood Grouping Sera Anti-B	Blood bank reagents or solutions	Address: PIMS Hospital, Sector G-8/3., Islamabad Capital Territory Schedule: As per Supply Order(s) Quantity: 1600/Qty	1600/Qty	20800 PKR	1	Any

Item	UNSPSC	Delivery Schedule	Quantity	Bid Security	Sample Quantity	Manufacturer / Dealer Authorization
Blood Grouping Sera Anti-D	Blood bank reagents or solutions	Address: PIMS Hospital, Sector G-8/3., Islamabad Capital Territory Schedule: As per Supply Order(s) Quantity: 2000/Qty	2000/Qty	32000 PKR	1	Any
Leukocyte Depletion Filters (RCC Pre-storage)	Blood transfusion administration kits	Address: PIMS Hospital, Sector G-8/3., Islamabad Capital Territory Schedule: As per Supply Order(s) Quantity: 1000/Qty	1000/Qty	56000 PKR	10	Any
Leukocyte Depletion Filters (RCC Bedside)	Blood transfusion administration kits	Address: PIMS Hospital, Sector G-8/3., Islamabad Capital Territory Schedule: As per Supply Order(s) Quantity: 6000/Qty	6000/Qty	269760 PKR	10	Any

Item	UNSPSC	Delivery Schedule	Quantity	Bid Security	Sample Quantity	Manufacturer / Dealer Authorization
Leukocyte Depletion Filters (Platelets Filters)	Blood transfusion administration kits	Address: PIMS Hospital, Sector G-8/3., Islamabad Capital Territory Schedule: As per Supply Order(s) Quantity: 1000/Qty	1000/Qty	63660 PKR	10	Any
Disposable Pasture Pipettes	Serological pipettes	Address: PIMS Hospital, Sector G-8/3., Islamabad Capital Territory Schedule: As per Supply Order(s) Quantity: 280000/Qty	280000/Qty	22008 PKR	100	Any
Test Tube 3" Glass	Centrifuge tubes	Address: PIMS Hospital, Sector G-8/3., Islamabad Capital Territory Schedule: As per Supply Order(s) Quantity: 70000/Qty	70000/Qty	3101 PKR	250	Any

Item	UNSPSC	Delivery Schedule	Quantity	Bid Security	Sample Quantity	Manufacturer / Dealer Authorization
NAT screening kit	Nucleic acid chemiluminescent detection materials	Address: PIMS Hospital, Sector G-8/3., Islamabad Capital Territory Schedule: As per Supply Order(s) Quantity: 60/Qty	60/Qty	576000 PKR	1	Any
Coomb's Sera	Blood bank reagents or solutions	Address: PIMS Hospital, Sector G-8/3., Islamabad Capital Territory Schedule: As per Supply Order(s) Quantity: 12/Qty	12/Qty	117 PKR	1	Any
Bovine Albumin	Blood bank reagents or solutions	Address: PIMS Hospital, Sector G-8/3., Islamabad Capital Territory Schedule: As per Supply Order(s) Quantity: 12/Qty	12/Qty	117 PKR	1	Any

Item	UNSPSC	Delivery Schedule	Quantity	Bid Security	Sample Quantity	Manufacturer / Dealer Authorization
LMB HGB Test Strips, with free placement of Hb Meter	Hemoglobin or hematocrit monitor or meter	Address: PIMS Hospital, Sector G-8/3., Islamabad Capital Territory Schedule: As per Supply Order(s) Quantity: 40000/Qty	40000/Qty	134000 PKR	50	Any
Platelets/ Plasma Mega Unit Kits:- (Apheresis Kit +ACD Solution/ Normal +Saline Fistula Needle)	Blood bank apheresis and donor unit processing analyzer	Address: PIMS Hospital, Sector G-8/3., Islamabad Capital Territory Schedule: As per Supply Order(s) Quantity: 500/Qty	500/Qty	197925 PKR	2	Any
ANTI-H LECTIN (BOMBAY ANTI SERA)	Blood bank reagents or solutions	Address: PIMS Hospital, Sector G-8/3., Islamabad Capital Territory Schedule: As per Supply Order(s) Quantity: 6/Qty	6/Qty	1800 PKR	1	Any

Item	UNSPSC	Delivery Schedule	Quantity	Bid Security	Sample Quantity	Manufacturer / Dealer Authorization
ANTI-A1 LECTIN (DOLICHOSS BIFLORUS) ANTISERA	Blood bank reagents or solutions	Address: PIMS Hospital, Sector G-8/3., Islamabad Capital Territory Schedule: As per Supply Order(s) Quantity: 24/Qty	24/Qty	720 PKR	1	Any
PAEDIATRIC BLOOD BAGS (QUINTUPLE)	Blood unit collection bags	Address: PIMS Hospital, Sector G-8/3., Islamabad Capital Territory Schedule: As per Supply Order(s) Quantity: 500/Qty	500/Qty	500 PKR	5	Any

Related Services of Goods:

No



Items/Lot Specification

Items Without Lots :

Item: Blood Collecting Bags (Single) without Set)

UNSPSC: Blood unit collection bags

Specifications / Requirements:

TS#. 1-5. TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS FOR THE TENDER OF BLOOD BAGS (SINGLE, TRIPLE & QUADRUPLE) FOR PIMS BLOOD BANK ISLAMABAD (FY 2026 – 2027)

S#	TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS	DESCRIPTION
1	SINGLE BLOOD BAG CAPACITY	Single Blood Bag 450 ml
2	TRIPLE BLOOD BAG CAPACITY	Primary bag (450 ml)
		Two satellite bags of upto 300 ml capacity for components
3	QUADRUPLE BLOOD BAG CAPACITY	Primary bag (450 ml)
		Three satellite bags of upto 300 ml capacity for components

TS#. 1-5. TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS FOR THE TENDER OF BLOOD BAGS (SINGLE, TRIPLE & QUADRUPLE) FOR PIMS BLOOD BANK ISLAMABAD (FY 2026 – 2027)

4	MATERIAL USED	Blood collection bag Made up of DEHP (Di-2-ethylhexyl phthalate) plasticized PVC (polyvinyl chloride), collapsible non- vented sterile containers complete with collecting tube for completely closed system to avoid the chances of contamination.
5	DESIGN AND SHAPE	<p>Flexible pre-sterilized</p> <p>Pyrogen free</p> <p>Nontoxic, non-hemolytic, biocompatible material</p> <p>No risk of contamination and air embolism (close system) with leaks proof seals</p> <p>Slit on both sides of the bags should be enough to accommodate 5-10 ml test tubes.</p> <p>The capacity of the bag should be enough to prevent any ballooning/ rupture of the bag from seam when it is filled up with the requisite volume of blood</p>
6	TUBING OF BAG	<p>Flexible non kinking</p> <p>Non sticking</p> <p>Transparent</p> <p>Leak proof</p> <p>The minimum length of tubing from primary bag to the needle should be 80 cm.</p>

TS#. 1-5. TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS FOR THE TENDER OF BLOOD BAGS (SINGLE, TRIPLE & QUADRUPLE) FOR PIMS BLOOD BANK ISLAMABAD (FY 2026 – 2027)

The tube should have multiple printed ID/Segment number. The number should be legible and clear.

A clamp should be provided for closed system

7	NEEDLE	16/ 17 gauge ultra-thin walled, siliconized and straight
		Sharp, regular and smooth margins and beveled tip
		Rust proof
		Tightly fixed with hub covered with sterile guard
		Hermetically sealed
		The needle should not separate from the tube at any point of time, especially while removing it from the vein for the donor safety
8	EXTERNAL PORT	Tamper proof and should not be re-capped
		Easily accessible

TS#. 1-5. TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS FOR THE TENDER OF BLOOD BAGS (SINGLE, TRIPLE & QUADRUPLE) FOR PIMS BLOOD BANK ISLAMABAD (FY 2026 – 2027)

9	PACKAGE	<p>Protective packaging (Aluminum) eliminating microbial contamination on surface maintaining the contents of the bag</p> <p>Easy to handle</p>
10	ANTICOAGULANT	<p>CPDA-1</p> <p>The quantity of anticoagulant (63 ml)</p> <p>Clear & colorless</p> <p>No discoloration on storage at room temperature</p> <p>Manufacturer to supply anticoagulant quality check certificate</p>
11	AIR BUBBLES	<p>Air bubble content in blood bag should be less than 15 ml</p>
12	LABEL	<p>Non peel-off</p> <p>Heat sealed/pressure embossed labels</p> <p>Remain attached between room temperature to - 80 °C with a transparent adhesive</p> <p>Following information must be printed (permanent) on label of blood bag:</p> <p>i. Date of manufacturing</p> <p>ii. Date of expiry</p> <p>iii. Lot number</p> <p>iv. Batch number</p>

**TS#. 1-5. TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS FOR THE TENDER OF BLOOD BAGS (SINGLE, TRIPLE & QUADRUPLE)
FOR PIMS BLOOD BANK ISLAMABAD (FY 2026 – 2027)**

v. Donor number

vi. ABO group

vii. Rh type

viii. Serology
report

ix. Manufacturer
name and
address

x. Anticoagulant
symbol and
chemical
formulation



TS#. 1-5. TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS FOR THE TENDER OF BLOOD BAGS (SINGLE, TRIPLE & QUADRUPLE) FOR PIMS BLOOD BANK ISLAMABAD (FY 2026 – 2027)

The expiry date should be at least 2 years from the date of manufacturing of blood bags and residual shelf life at the time of supply should be at least 3/4th of the total shelf life

13	RESISTANCE TO DISTORTION	Bag shall with stand an acceleration of 5000 g for 30 min at temperature 4°C to 24°C
		Bag should be able to withstand temperature upto-80°C without breakage
14	DIVERSION POUCH	Blood bags with diversion pouch and needle guard will be preferred
15	CERTIFICATION AND APPROVAL	CE marking or any equivalent certification
		Registered with DRAP
		The international marking certificates must be valid; clearly mentioning the brand name of product being referred and name of the company with address given on pack and must be verifiable at company websites
		Products having doubtful or invalid international certificate will be rejected

TS#. 1-5. TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS FOR THE TENDER OF BLOOD BAGS (SINGLE, TRIPLE & QUADRUPLE) FOR PIMS BLOOD BANK ISLAMABAD (FY 2026 – 2027)

16	REFERENCES	Must have references of at least three tertiary care hospitals/ blood banks of Pakistan. Satisfactory reports with address and contact number required.
17	SOLE DISTRIBUTION/ LOCAL OFFICES	The company should have authorization for sole distribution and local sold distribution office and available stock.

Item: Blood Collecting Bags with diversion pouch triple without set

UNSPSC: Blood unit collection bags

Specifications / Requirements:

TS#. 1-5. TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS FOR THE TENDER OF BLOOD BAGS (SINGLE, TRIPLE & QUADRUPLE) FOR PIMS BLOOD BANK ISLAMABAD (FY 2026 – 2027)

S#	TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS	DESCRIPTION
1	SINGLE BLOOD BAG CAPACITY	Single Blood Bag 450 ml
2	TRIPLE BLOOD BAG CAPACITY	Primary bag (450 ml) Two satellite bags of upto 300 ml capacity for components
3	QUADRUPLE BLOOD BAG CAPACITY	Primary bag (450 ml) Three satellite bags of upto 300 ml capacity for components

TS#. 1-5. TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS FOR THE TENDER OF BLOOD BAGS (SINGLE, TRIPLE & QUADRUPLE) FOR PIMS BLOOD BANK ISLAMABAD (FY 2026 – 2027)

4	MATERIAL USED	Blood collection bag Made up of DEHP (Di-2-ethylhexyl phthalate) plasticized PVC (polyvinyl chloride), collapsible non- vented sterile containers complete with collecting tube for completely closed system to avoid the chances of contamination.
5	DESIGN AND SHAPE	<p>Flexible pre-sterilized</p> <p>Pyrogen free</p> <p>Nontoxic, non-hemolytic, biocompatible material</p> <p>No risk of contamination and air embolism (close system) with leaks proof seals</p> <p>Slit on both sides of the bags should be enough to accommodate 5-10 ml test tubes.</p> <p>The capacity of the bag should be enough to prevent any ballooning/ rupture of the bag from seam when it is filled up with the requisite volume of blood</p>
6	TUBING OF BAG	<p>Flexible non kinking</p> <p>Non sticking</p> <p>Transparent</p> <p>Leak proof</p> <p>The minimum length of tubing from primary bag to the needle should be 80 cm.</p>

TS#. 1-5. TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS FOR THE TENDER OF BLOOD BAGS (SINGLE, TRIPLE & QUADRUPLE) FOR PIMS BLOOD BANK ISLAMABAD (FY 2026 – 2027)

The tube should have multiple printed ID/Segment number. The number should be legible and clear.

A clamp should be provided for closed system

7	NEEDLE	16/ 17 gauge ultra-thin walled, siliconized and straight
		Sharp, regular and smooth margins and beveled tip
		Rust proof
		Tightly fixed with hub covered with sterile guard
		Hermetically sealed
		The needle should not separate from the tube at any point of time, especially while removing it from the vein for the donor safety
8	EXTERNAL PORT	Tamper proof and should not be re-capped
		Easily accessible

TS#. 1-5. TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS FOR THE TENDER OF BLOOD BAGS (SINGLE, TRIPLE & QUADRUPLE) FOR PIMS BLOOD BANK ISLAMABAD (FY 2026 – 2027)

9	PACKAGE	<p>Protective packaging (Aluminum) eliminating microbial contamination on surface maintaining the contents of the bag</p> <p>Easy to handle</p>
10	ANTICOAGULANT	<p>CPDA-1</p> <p>The quantity of anticoagulant (63 ml)</p> <p>Clear & colorless</p> <p>No discoloration on storage at room temperature</p> <p>Manufacturer to supply anticoagulant quality check certificate</p>
11	AIR BUBBLES	<p>Air bubble content in blood bag should be less than 15 ml</p>
12	LABEL	<p>Non peel-off</p> <p>Heat sealed/pressure embossed labels</p> <p>Remain attached between room temperature to - 80 °C with a transparent adhesive</p> <p>Following information must be printed (permanent) on label of blood bag:</p> <p>i. Date of manufacturing</p> <p>ii. Date of expiry</p> <p>iii. Lot number</p> <p>iv. Batch number</p>

**TS#. 1-5. TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS FOR THE TENDER OF BLOOD BAGS (SINGLE, TRIPLE & QUADRUPLE)
FOR PIMS BLOOD BANK ISLAMABAD (FY 2026 – 2027)**

v. Donor number

vi. ABO group

vii. Rh type

viii. Serology
report

ix. Manufacturer
name and
address

x. Anticoagulant
symbol and
chemical
formulation



TS#. 1-5. TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS FOR THE TENDER OF BLOOD BAGS (SINGLE, TRIPLE & QUADRUPLE) FOR PIMS BLOOD BANK ISLAMABAD (FY 2026 – 2027)

The expiry date should be at least 2 years from the date of manufacturing of blood bags and residual shelf life at the time of supply should be at least 3/4th of the total shelf life

13	RESISTANCE TO DISTORTION	Bag shall with stand an acceleration of 5000 g for 30 min at temperature 4°C to 24°C
		Bag should be able to withstand temperature upto-80°C without breakage
14	DIVERSION POUCH	Blood bags with diversion pouch and needle guard will be preferred
15	CERTIFICATION AND APPROVAL	CE marking or any equivalent certification
		Registered with DRAP
		The international marking certificates must be valid; clearly mentioning the brand name of product being referred and name of the company with address given on pack and must be verifiable at company websites
		Products having doubtful or invalid international certificate will be rejected

TS#. 1-5. TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS FOR THE TENDER OF BLOOD BAGS (SINGLE, TRIPLE & QUADRUPLE) FOR PIMS BLOOD BANK ISLAMABAD (FY 2026 – 2027)

16	REFERENCES	Must have references of at least three tertiary care hospitals/ blood banks of Pakistan. Satisfactory reports with address and contact number required.
17	SOLE DISTRIBUTION/ LOCAL OFFICES	The company should have authorization for sole distribution and local sold distribution office and available stock.

Item: Blood Collecting Bags without diversion pouch (Triple without Set)

UNSPSC: Blood unit collection bags

Specifications / Requirements:

TS#. 1-5. TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS FOR THE TENDER OF BLOOD BAGS (SINGLE, TRIPLE & QUADRUPLE) FOR PIMS BLOOD BANK ISLAMABAD (FY 2026 – 2027)

S#	TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS	DESCRIPTION
1	SINGLE BLOOD BAG CAPACITY	Single Blood Bag 450 ml
2	TRIPLE BLOOD BAG CAPACITY	Primary bag (450 ml) Two satellite bags of upto 300 ml capacity for components
3	QUADRUPLE BLOOD BAG CAPACITY	Primary bag (450 ml) Three satellite bags of upto 300 ml capacity for components

TS#. 1-5. TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS FOR THE TENDER OF BLOOD BAGS (SINGLE, TRIPLE & QUADRUPLE) FOR PIMS BLOOD BANK ISLAMABAD (FY 2026 – 2027)

4	MATERIAL USED	Blood collection bag Made up of DEHP (Di-2-ethylhexyl phthalate) plasticized PVC (polyvinyl chloride), collapsible non- vented sterile containers complete with collecting tube for completely closed system to avoid the chances of contamination.
5	DESIGN AND SHAPE	<p>Flexible pre-sterilized</p> <p>Pyrogen free</p> <p>Nontoxic, non-hemolytic, biocompatible material</p> <p>No risk of contamination and air embolism (close system) with leaks proof seals</p> <p>Slit on both sides of the bags should be enough to accommodate 5-10 ml test tubes.</p> <p>The capacity of the bag should be enough to prevent any ballooning/ rupture of the bag from seam when it is filled up with the requisite volume of blood</p>
6	TUBING OF BAG	<p>Flexible non kinking</p> <p>Non sticking</p> <p>Transparent</p> <p>Leak proof</p> <p>The minimum length of tubing from primary bag to the needle should be 80 cm.</p>

TS#. 1-5. TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS FOR THE TENDER OF BLOOD BAGS (SINGLE, TRIPLE & QUADRUPLE) FOR PIMS BLOOD BANK ISLAMABAD (FY 2026 – 2027)

The tube should have multiple printed ID/Segment number. The number should be legible and clear.

A clamp should be provided for closed system

7	NEEDLE	16/ 17 gauge ultra-thin walled, siliconized and straight
		Sharp, regular and smooth margins and beveled tip
		Rust proof
		Tightly fixed with hub covered with sterile guard
		Hermetically sealed
		The needle should not separate from the tube at any point of time, especially while removing it from the vein for the donor safety
8	EXTERNAL PORT	Tamper proof and should not be re-capped
		Easily accessible

TS#. 1-5. TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS FOR THE TENDER OF BLOOD BAGS (SINGLE, TRIPLE & QUADRUPLE) FOR PIMS BLOOD BANK ISLAMABAD (FY 2026 – 2027)

9	PACKAGE	Protective packaging (Aluminum) eliminating microbial contamination on surface maintaining the contents of the bag Easy to handle
10	ANTICOAGULANT	CPDA-1 The quantity of anticoagulant (63 ml) Clear & colorless No discoloration on storage at room temperature Manufacturer to supply anticoagulant quality check certificate
11	AIR BUBBLES	Air bubble content in blood bag should be less than 15 ml
12	LABEL	Non peel-off Heat sealed/pressure embossed labels Remain attached between room temperature to - 80 °C with a transparent adhesive Following information must be printed (permanent) on label of blood bag: i. Date of manufacturing ii. Date of expiry iii. Lot number iv. Batch number

**TS#. 1-5. TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS FOR THE TENDER OF BLOOD BAGS (SINGLE, TRIPLE & QUADRUPLE)
FOR PIMS BLOOD BANK ISLAMABAD (FY 2026 – 2027)**

v. Donor number

vi. ABO group

vii. Rh type

viii. Serology
report

ix. Manufacturer
name and
address

x. Anticoagulant
symbol and
chemical
formulation



TS#. 1-5. TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS FOR THE TENDER OF BLOOD BAGS (SINGLE, TRIPLE & QUADRUPLE) FOR PIMS BLOOD BANK ISLAMABAD (FY 2026 – 2027)

The expiry date should be at least 2 years from the date of manufacturing of blood bags and residual shelf life at the time of supply should be at least 3/4th of the total shelf life

13	RESISTANCE TO DISTORTION	Bag shall with stand an acceleration of 5000 g for 30 min at temperature 4°C to 24°C
		Bag should be able to withstand temperature upto-80°C without breakage
14	DIVERSION POUCH	Blood bags with diversion pouch and needle guard will be preferred
15	CERTIFICATION AND APPROVAL	CE marking or any equivalent certification
		Registered with DRAP
		The international marking certificates must be valid; clearly mentioning the brand name of product being referred and name of the company with address given on pack and must be verifiable at company websites
		Products having doubtful or invalid international certificate will be rejected

TS#. 1-5. TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS FOR THE TENDER OF BLOOD BAGS (SINGLE, TRIPLE & QUADRUPLE) FOR PIMS BLOOD BANK ISLAMABAD (FY 2026 – 2027)

16	REFERENCES	Must have references of at least three tertiary care hospitals/ blood banks of Pakistan. Satisfactory reports with address and contact number required.
17	SOLE DISTRIBUTION/ LOCAL OFFICES	The company should have authorization for sole distribution and local sold distribution office and available stock.

Item: Blood Collecting Bags with diversion pouch (Quadruple without Set)

UNSPSC: Blood unit collection bags

Specifications / Requirements:

TS#. 1-5. TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS FOR THE TENDER OF BLOOD BAGS (SINGLE, TRIPLE & QUADRUPLE) FOR PIMS BLOOD BANK ISLAMABAD (FY 2026 – 2027)

S#	TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS	DESCRIPTION
1	SINGLE BLOOD BAG CAPACITY	Single Blood Bag 450 ml
2	TRIPLE BLOOD BAG CAPACITY	Primary bag (450 ml) Two satellite bags of upto 300 ml capacity for components
3	QUADRUPLE BLOOD BAG CAPACITY	Primary bag (450 ml) Three satellite bags of upto 300 ml capacity for components

TS#. 1-5. TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS FOR THE TENDER OF BLOOD BAGS (SINGLE, TRIPLE & QUADRUPLE) FOR PIMS BLOOD BANK ISLAMABAD (FY 2026 – 2027)

4	MATERIAL USED	Blood collection bag Made up of DEHP (Di-2-ethylhexyl phthalate) plasticized PVC (polyvinyl chloride), collapsible non- vented sterile containers complete with collecting tube for completely closed system to avoid the chances of contamination.
5	DESIGN AND SHAPE	<p>Flexible pre-sterilized</p> <p>Pyrogen free</p> <p>Nontoxic, non-hemolytic, biocompatible material</p> <p>No risk of contamination and air embolism (close system) with leaks proof seals</p> <p>Slit on both sides of the bags should be enough to accommodate 5-10 ml test tubes.</p> <p>The capacity of the bag should be enough to prevent any ballooning/ rupture of the bag from seam when it is filled up with the requisite volume of blood</p>
6	TUBING OF BAG	<p>Flexible non kinking</p> <p>Non sticking</p> <p>Transparent</p> <p>Leak proof</p> <p>The minimum length of tubing from primary bag to the needle should be 80 cm.</p>

TS#. 1-5. TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS FOR THE TENDER OF BLOOD BAGS (SINGLE, TRIPLE & QUADRUPLE) FOR PIMS BLOOD BANK ISLAMABAD (FY 2026 – 2027)

The tube should have multiple printed ID/Segment number. The number should be legible and clear.

A clamp should be provided for closed system

7	NEEDLE	16/ 17 gauge ultra-thin walled, siliconized and straight
		Sharp, regular and smooth margins and beveled tip
		Rust proof
		Tightly fixed with hub covered with sterile guard
		Hermetically sealed
		The needle should not separate from the tube at any point of time, especially while removing it from the vein for the donor safety
8	EXTERNAL PORT	Tamper proof and should not be re-capped
		Easily accessible

TS#. 1-5. TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS FOR THE TENDER OF BLOOD BAGS (SINGLE, TRIPLE & QUADRUPLE) FOR PIMS BLOOD BANK ISLAMABAD (FY 2026 – 2027)

9	PACKAGE	<p>Protective packaging (Aluminum) eliminating microbial contamination on surface maintaining the contents of the bag</p> <p>Easy to handle</p>
10	ANTICOAGULANT	<p>CPDA-1</p> <p>The quantity of anticoagulant (63 ml)</p> <p>Clear & colorless</p> <p>No discoloration on storage at room temperature</p> <p>Manufacturer to supply anticoagulant quality check certificate</p>
11	AIR BUBBLES	<p>Air bubble content in blood bag should be less than 15 ml</p>
12	LABEL	<p>Non peel-off</p> <p>Heat sealed/pressure embossed labels</p> <p>Remain attached between room temperature to - 80 °C with a transparent adhesive</p> <p>Following information must be printed (permanent) on label of blood bag:</p> <p>i. Date of manufacturing</p> <p>ii. Date of expiry</p> <p>iii. Lot number</p> <p>iv. Batch number</p>

**TS#. 1-5. TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS FOR THE TENDER OF BLOOD BAGS (SINGLE, TRIPLE & QUADRUPLE)
FOR PIMS BLOOD BANK ISLAMABAD (FY 2026 – 2027)**

v. Donor number

vi. ABO group

vii. Rh type

viii. Serology
report

ix. Manufacturer
name and
address

x. Anticoagulant
symbol and
chemical
formulation



TS#. 1-5. TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS FOR THE TENDER OF BLOOD BAGS (SINGLE, TRIPLE & QUADRUPLE) FOR PIMS BLOOD BANK ISLAMABAD (FY 2026 – 2027)

The expiry date should be at least 2 years from the date of manufacturing of blood bags and residual shelf life at the time of supply should be at least 3/4th of the total shelf life

13	RESISTANCE TO DISTORTION	Bag shall with stand an acceleration of 5000 g for 30 min at temperature 4°C to 24°C
		Bag should be able to withstand temperature upto-80°C without breakage
14	DIVERSION POUCH	Blood bags with diversion pouch and needle guard will be preferred
15	CERTIFICATION AND APPROVAL	CE marking or any equivalent certification
		Registered with DRAP
		The international marking certificates must be valid; clearly mentioning the brand name of product being referred and name of the company with address given on pack and must be verifiable at company websites
		Products having doubtful or invalid international certificate will be rejected

TS#. 1-5. TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS FOR THE TENDER OF BLOOD BAGS (SINGLE, TRIPLE & QUADRUPLE) FOR PIMS BLOOD BANK ISLAMABAD (FY 2026 – 2027)

16	REFERENCES	Must have references of at least three tertiary care hospitals/ blood banks of Pakistan. Satisfactory reports with address and contact number required.
17	SOLE DISTRIBUTION/ LOCAL OFFICES	The company should have authorization for sole distribution and local sold distribution office and available stock.

Item: Blood Collecting Bags without diversion pouch (Quadruple without Set)

UNSPSC: Blood unit collection bags

Specifications / Requirements:

TS#. 1-5. TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS FOR THE TENDER OF BLOOD BAGS (SINGLE, TRIPLE & QUADRUPLE) FOR PIMS BLOOD BANK ISLAMABAD (FY 2026 – 2027)

S#	TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS	DESCRIPTION
1	SINGLE BLOOD BAG CAPACITY	Single Blood Bag 450 ml
2	TRIPLE BLOOD BAG CAPACITY	Primary bag (450 ml) Two satellite bags of upto 300 ml capacity for components
3	QUADRUPLE BLOOD BAG CAPACITY	Primary bag (450 ml) Three satellite bags of upto 300 ml capacity for components

TS#. 1-5. TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS FOR THE TENDER OF BLOOD BAGS (SINGLE, TRIPLE & QUADRUPLE) FOR PIMS BLOOD BANK ISLAMABAD (FY 2026 – 2027)

4	MATERIAL USED	Blood collection bag Made up of DEHP (Di-2-ethylhexyl phthalate) plasticized PVC (polyvinyl chloride), collapsible non- vented sterile containers complete with collecting tube for completely closed system to avoid the chances of contamination.
5	DESIGN AND SHAPE	<p>Flexible pre-sterilized</p> <p>Pyrogen free</p> <p>Nontoxic, non-hemolytic, biocompatible material</p> <p>No risk of contamination and air embolism (close system) with leaks proof seals</p> <p>Slit on both sides of the bags should be enough to accommodate 5-10 ml test tubes.</p> <p>The capacity of the bag should be enough to prevent any ballooning/ rupture of the bag from seam when it is filled up with the requisite volume of blood</p>
6	TUBING OF BAG	<p>Flexible non kinking</p> <p>Non sticking</p> <p>Transparent</p> <p>Leak proof</p> <p>The minimum length of tubing from primary bag to the needle should be 80 cm.</p>

TS#. 1-5. TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS FOR THE TENDER OF BLOOD BAGS (SINGLE, TRIPLE & QUADRUPLE) FOR PIMS BLOOD BANK ISLAMABAD (FY 2026 – 2027)

The tube should have multiple printed ID/Segment number. The number should be legible and clear.

A clamp should be provided for closed system

7	NEEDLE	16/ 17 gauge ultra-thin walled, siliconized and straight
		Sharp, regular and smooth margins and beveled tip
		Rust proof
		Tightly fixed with hub covered with sterile guard
		Hermetically sealed
		The needle should not separate from the tube at any point of time, especially while removing it from the vein for the donor safety
8	EXTERNAL PORT	Tamper proof and should not be re-capped
		Easily accessible

TS#. 1-5. TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS FOR THE TENDER OF BLOOD BAGS (SINGLE, TRIPLE & QUADRUPLE) FOR PIMS BLOOD BANK ISLAMABAD (FY 2026 – 2027)

9	PACKAGE	<p>Protective packaging (Aluminum) eliminating microbial contamination on surface maintaining the contents of the bag</p> <p>Easy to handle</p>
10	ANTICOAGULANT	<p>CPDA-1</p> <p>The quantity of anticoagulant (63 ml)</p> <p>Clear & colorless</p> <p>No discoloration on storage at room temperature</p> <p>Manufacturer to supply anticoagulant quality check certificate</p>
11	AIR BUBBLES	<p>Air bubble content in blood bag should be less than 15 ml</p>
12	LABEL	<p>Non peel-off</p> <p>Heat sealed/pressure embossed labels</p> <p>Remain attached between room temperature to - 80 °C with a transparent adhesive</p> <p>Following information must be printed (permanent) on label of blood bag:</p> <p>i. Date of manufacturing</p> <p>ii. Date of expiry</p> <p>iii. Lot number</p> <p>iv. Batch number</p>

**TS#. 1-5. TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS FOR THE TENDER OF BLOOD BAGS (SINGLE, TRIPLE & QUADRUPLE)
FOR PIMS BLOOD BANK ISLAMABAD (FY 2026 – 2027)**

v. Donor number

vi. ABO group

vii. Rh type

viii. Serology
report

ix. Manufacturer
name and
address

x. Anticoagulant
symbol and
chemical
formulation



TS#. 1-5. TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS FOR THE TENDER OF BLOOD BAGS (SINGLE, TRIPLE & QUADRUPLE) FOR PIMS BLOOD BANK ISLAMABAD (FY 2026 – 2027)

The expiry date should be at least 2 years from the date of manufacturing of blood bags and residual shelf life at the time of supply should be at least 3/4th of the total shelf life

13	RESISTANCE TO DISTORTION	Bag shall with stand an acceleration of 5000 g for 30 min at temperature 4°C to 24°C
		Bag should be able to withstand temperature upto-80°C without breakage
14	DIVERSION POUCH	Blood bags with diversion pouch and needle guard will be preferred
15	CERTIFICATION AND APPROVAL	CE marking or any equivalent certification
		Registered with DRAP
		The international marking certificates must be valid; clearly mentioning the brand name of product being referred and name of the company with address given on pack and must be verifiable at company websites
		Products having doubtful or invalid international certificate will be rejected

TS#. 1-5. TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS FOR THE TENDER OF BLOOD BAGS (SINGLE, TRIPLE & QUADRUPLE) FOR PIMS BLOOD BANK ISLAMABAD (FY 2026 – 2027)

16	REFERENCES	Must have references of at least three tertiary care hospitals/ blood banks of Pakistan. Satisfactory reports with address and contact number required.
17	SOLE DISTRIBUTION/ LOCAL OFFICES	The company should have authorization for sole distribution and local sold distribution office and available stock.

Item: Blood Transfusion Set

UNSPSC: Blood transfusion administration kits

Specifications / Requirements:

Sr.	TECHNICAL SPECIFICATIONS	DESCRIPTION
1	Purpose of Equipment	Used for transfusion of RCC
2	Type of Blood Transfusion Set	Double chamber with filter
3	Material	Natural rubber latex
4	Shelf Life	3 Years from the date of manufacturer
5	Spike	ABS Material
6	Drip Chamber	PVC Material
7	Connector	Luer Slip
8	Injection Port	Latex Injection Port

Sr.	TECHNICAL SPECIFICATIONS	DESCRIPTION
9	Air Vent	Without built-in air vent
10	Roller thumb controller	Specially designed roller (thumb) controller after accurate regulation of infusion rate

Item: Blood Grouping Sera Anti-A

UNSPSC: Blood bank reagents or solutions

Specifications / Requirements:

S#	TECHNICAL SPECIFICATIONS	DESCRIPTION
1	Description	Blood grouping sera (A, B & D) are used for detecting the blood group by slide and tube test
2	Type of antibodies	Monoclonal human or animal antibodies
3	Volume	10 ml/ Bottle
4	Color of reagents	<ul style="list-style-type: none"> · Anti A Blue · Anti B Yellow · Anti D Colorless
5	Sealed Lid Packing	<ul style="list-style-type: none"> · Preferably sealed lid packing
6	Preservative	0.1 % sodium azide
7	Agglutination Titer	i. Titer result of monoclonal Anti A and Anti B antisera must be 1:256 or above

S#	TECHNICAL SPECIFICATIONS	DESCRIPTION
	ii. Titer result of monoclonal anti D antisera must be 1:128 or above	
8	References	Must have references of at least three tertiary care hospitals/ blood banks of Pakistan (Preferably local). Satisfactory reports with address and contact number required.
9	Certification	It should be FDA approved, CE Marked or any other equivalent certification
10	DRAP registration	Preferably registered with DRAP
11	Anti sera A, B & D must be of one brand	All anti sera A, B & D must be of one brand
Note:		
i. Only monoclonal high titer approved reagents should be used National Guidelines "Quality Control in Transfusion Medicine".		
ii. Specifications are according to the National Guidelines "Quality Control in Transfusion Medicine".		

S#	TECHNICAL SPECIFICATIONS	DESCRIPTION
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iii. Clause# 8 is mandatory

Item: Blood Grouping Sera Anti-B

UNSPSC: Blood bank reagents or solutions

Specifications / Requirements:

S#	TECHNICAL SPECIFICATIONS	DESCRIPTION
1	Description	Blood grouping sera (A, B & D) are used for detecting the blood group by slide and tube test
2	Type of antibodies	Monoclonal human or animal antibodies
3	Volume	10 ml/ Bottle
4	Color of reagents	<ul style="list-style-type: none"> · Anti A Blue · Anti B Yellow · Anti D Colorless
5	Sealed Lid Packing	· Preferably sealed lid packing
6	Preservative	0.1 % sodium azide
7	Agglutination Titer	<ul style="list-style-type: none"> i. Titer result of monoclonal Anti A and Anti B antisera must be 1:256 or above ii. Titer result of monoclonal anti D antisera must be 1:128 or above

S#	TECHNICAL SPECIFICATIONS	DESCRIPTION
8	References	Must have references of at least three tertiary care hospitals/ blood banks of Pakistan (Preferably local). Satisfactory reports with address and contact number required.
9	Certification	It should be FDA approved, CE Marked or any other equivalent certification
10	DRAP registration	Preferably registered with DRAP
11	Anti sera A, B & D must be of one brand	All anti sera A, B & D must be of one brand
Note:		
		i. Only monoclonal high titer approved reagents should be used National Guidelines "Quality Control in Transfusion Medicine".
		ii. Specifications are according to the National Guidelines "Quality Control in Transfusion Medicine".
		iii. Clause# 8 is mandatory

Item: Blood Grouping Sera Anti-D

UNSPSC: Blood bank reagents or solutions

Specifications / Requirements:

S#	TECHNICAL SPECIFICATIONS	DESCRIPTION
1	Description	Blood grouping sera (A, B & D) are used for detecting the blood group by slide and tube test

S#	TECHNICAL SPECIFICATIONS	DESCRIPTION
2	Type of antibodies	Monoclonal human or animal antibodies
3	Volume	10 ml/ Bottle
4	Color of reagents	<ul style="list-style-type: none"> · Anti A Blue · Anti B Yellow · Anti D Colorless
5	Sealed Lid Packing	· Preferably sealed lid packing
6	Preservative	0.1 % sodium azide
7	Agglutination Titer	<ul style="list-style-type: none"> i. Titer result of monoclonal Anti A and Anti B antisera must be 1:256 or above ii. Titer result of monoclonal anti D antisera must be 1:128 or above
8	References	Must have references of at least three tertiary care hospitals/ blood banks of Pakistan (Preferably local). Satisfactory reports with address and contact number required.
9	Certification	It should be FDA approved, CE Marked or any other equivalent certification
10	DRAP registration	Preferably registered with DRAP
11	Anti sera A, B & D must be of one brand	All anti sera A, B & D must be of one brand
Note:		
i. Only monoclonal high titer approved reagents should be used National Guidelines "Quality Control in Transfusion Medicine".		
ii. Specifications are according to the National Guidelines "Quality Control in Transfusion Medicine".		

S#	TECHNICAL SPECIFICATIONS	DESCRIPTION
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iii. Clause# 8 is mandatory

Item: Leukocyte Depletion Filters (RCC Pre-storage)

UNSPSC: Blood transfusion administration kits

Specifications / Requirements:

S#	TECHNICAL SPECIFICATIONS	DESCRIPTION
1	Description	The Leukocyte Reduction Filter for Red Blood Cells is indicated for the depletion of leukocytes in a single unit of red blood cells collected from a healthy adult
2	Filtration Technique	Filtration by gravity, no saline filter prime or flush is required
3	Types of Filters	1. RCC (Pre-Storage) Filter
		2. RCC (Bed Side) Filter
		3. Platelets Filter
4	Satellite blood bag with RCC Filter (Pre Storage)	RCC Filter (Pre Storage) must have a satellite blood bag for leukodepleted RCC
5	References	Local references will be preferred
6	Certification	FDA approved or CE Marked or any equivalent certification (Preferably FDA approved)
7	Surface coating technology	Preferably ZISC Technology

S#	TECHNICAL SPECIFICATIONS	DESCRIPTION
8	Ambient Temperature	20 to 24 °C
9	Filtration Time of	Minimum time for RCC (Bed-side) and Platelets filters
	· RCC filter (Pre-storage)	
	· RCC filter (Bed-side)	
	· Platelets filter	
10	Residual Volume	· Upto 10 Minutes for RCC (Pre-storage)
	· RCC filter (Pre-storage)	
	· RCC filter (Bed-side)	· 25 ml
11	Leukocyte reduction rate	> 99.0 %
12	RCC/ Platelets Recovery	> 80 %
13	Leukocyte depletion efficiency	High leukocyte depletion efficiency 1x 10(e)6 WBC/unit
14	Provision of technical services	Successful vendor will be responsible to depute one technical assistant for doing filtration procedures successfully and to avoid wastage of precious products

Item: Leukocyte Depletion Filters (RCC Bedside)

UNSPSC: Blood transfusion administration kits

Specifications / Requirements:

S#	TECHNICAL SPECIFICATIONS	DESCRIPTION
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1	Description	The Leukocyte Reduction Filter for Red Blood Cells is indicated for the depletion of leukocytes in a single unit of red blood cells collected from a healthy adult
2	Filtration Technique	Filtration by gravity, no saline filter prime or flush is required
3	Types of Filters	1. RCC (Pre-Storage) Filter 2. RCC (Bed Side) Filter 3. Platelets Filter
4	Satellite blood bag with RCC Filter (Pre Storage)	RCC Filter (Pre Storage) must have a satellite blood bag for leukodepleted RCC
5	References	Local references will be preferred
6	Certification	FDA approved or CE Marked or any equivalent certification (Preferably FDA approved)
7	Surface coating technology	Preferably ZISC Technology
8	Ambient Temperature	20 to 24 °C
9	Filtration Time of	Minimum time for RCC (Bed-side) and Platelets filters
	· RCC filter (Pre-storage)	
	· RCC filter (Bed-side)	
	· Platelets filter	
10	Residual Volume	· Upto 10 Minutes for RCC (Pre-storage)
	· RCC filter (Pre-storage)	
	· RCC filter (Bed-side)	· 25 ml
11	Leukocyte reduction rate	> 99.0 %

12	RCC/ Platelets Recovery	> 80 %
13	Leukocyte depletion efficiency	High leukocyte depletion efficiency 1x 10(e)6 WBC/unit
14	Provision of technical services	Successful vendor will be responsible to depute one technical assistant for doing filtration procedures successfully and to avoid wastage of precious products

Item: Leukocyte Depletion Filters (Platelets Filters)

UNSPSC: Blood transfusion administration kits

Specifications / Requirements:

S#	TECHNICAL SPECIFICATIONS	DESCRIPTION
1	Description	The Leukocyte Reduction Filter for Red Blood Cells is indicated for the depletion of leukocytes in a single unit of red blood cells collected from a healthy adult
2	Filtration Technique	Filtration by gravity, no saline filter prime or flush is required
3	Types of Filters	1. RCC (Pre-Storage) Filter 2. RCC (Bed Side) Filter 3. Platelets Filter
4	Satellite blood bag with RCC Filter (Pre Storage)	RCC Filter (Pre Storage) must have a satellite blood bag for leukodepleted RCC
5	References	Local references will be preferred
6	Certification	FDA approved or CE Marked or any equivalent certification (Preferably FDA approved)

S#	TECHNICAL SPECIFICATIONS	DESCRIPTION
7	Surface coating technology	Preferably ZISC Technology
8	Ambient Temperature	20 to 24 °C
9	Filtration Time of	Minimum time for RCC (Bed-side) and Platelets filters
	· RCC filter (Pre-storage)	
	· RCC filter (Bed-side)	
	· Platelets filter	
10	Residual Volume	· Upto 10 Minutes for RCC (Pre-storage)
	· RCC filter (Pre-storage)	
	· RCC filter (Bed-side)	· 25 ml
11	Leukocyte reduction rate	> 99.0 %
12	RCC/ Platelets Recovery	> 80 %
13	Leukocyte depletion efficiency	High leukocyte depletion efficiency 1×10^6 WBC/unit
14	Provision of technical services	Successful vendor will be responsible to depute one technical assistant for doing filtration procedures successfully and to avoid wastage of precious products

Item: Disposable Pasture Pipettes

UNSPSC: Serological pipettes

Specifications / Requirements:

S#	TECHNICAL SPECIFICATIONS	DESCRIPTION
1	Description	Known as dropper used for transferring of blood samples or liquids
2	Material	Plastic
3	Size	Upto 3.0 ml
4	Graduation	Must be graduated upto 3.0 ml
5	Type	Disposable

Item: Test Tube 3" Glass

UNSPSC: Centrifuge tubes

Specifications / Requirements:

S#	TECHNICAL SPECIFICATIONS	DESCRIPTION
1	Description	Plain transparent glass tube to be used for blood grouping and cross matching
2	Material	Glass
3	Size	3"
4	Transparency	Must be transparent to check visible agglutination of blood grouping

Item: NAT screening kit

UNSPSC: Nucleic acid chemiluminescent detection materials

Specifications / Requirements:

S#	TECHNICAL SPECIFICATIONS	DESCRIPTION
1	Description	NAT screening kit for HBV, HCV & HIV 1 & 2
2	Compatibility	Must be compatible with Cobas 5800/ 6800 by Roche
3	Manufacturer	Roche
4	Consumables & Accessories	All accessories & consumables including calibrators, controls etc. will be responsibility of vendor on FOC basis
5	Certification	FDA approved or CE Marked or any equivalent certification

Item: Coomb's Sera

UNSPSC: Blood bank reagents or solutions

Specifications / Requirements:

S#	TECHNICAL SPECIFICATIONS	DESCRIPTION
1	Description	Coomb's sera/ Anti-Human Globulins (Anti-IgG) are used in the direct antiglobulin test to detect the in vivo coating on human red blood cells with IgG, and for indirect antiglobulin test for antibody screening and identification, and Cross match.
2	Type of antibodies	Monoclonal/ Polyclonal human or animal antibodies
3	Volume	10 ml/ Bottle

S#	TECHNICAL SPECIFICATIONS	DESCRIPTION
4	Sealed Lid Packing	Preferably sealed lid packing
5	Agglutination Titer	Sera with greater titer will be preferred
6	References	Local references will be preferred.
7	Certification	FDA approved or CE Marked or any equivalent certification (Preferably FDA approved)
8	DRAP registration	Preferably registered with DRAP

Item: Bovine Albumin

UNSPSC: Blood bank reagents or solutions

Specifications / Requirements:

S#	TECHNICAL SPECIFICATIONS	DESCRIPTION
1	Description	Bovine Albumin Sera is used for compatibility testing, for unexpected antibody testing and antigen-antibody interactions enhancement
2	Type of antibodies	Monoclonal human or animal antibodies
3	Volume	10 ml/ Bottle
4	Sealed Lid Packing	Preferably sealed lid packing
5	Agglutination Titer	Sera with greater titer will be preferred
6	References	Local references will be preferred.

S#	TECHNICAL SPECIFICATIONS	DESCRIPTION
7	Certification	FDA approved or CE Marked or any equivalent certification (Preferably FDA approved)
8	DRAP registration	Preferably registered with DRAP

Item: LMB HGB Test Strips, with free placement of Hb Meter

UNSPSC: Hemoglobin or hematocrit monitor or meter

Specifications / Requirements:

Sr#	TECHNICAL SPECIFICATIONS / TERMS & CONDITIONS	DESCRIPTION
1	HGB Meter Description	Testing of blood Hemoglobin (HGB) level (Blood donors)
2	Principle	Photometric
3	Detection System	Relectance Photometry/ Met-hemoglobin
4	Measurement range	3 - 25 g/dl
5	Sample Volume	Upto 10 ul
6	Sample Type	Whole blood (Venous) and capillary blood
7	Certification	FDA approved or CE Marked or any equivalent certification (Preferably FDA approved)
8	Number of equipments	Three HGB Meters must be provided
9	Repair/ Maintenance of equipments	Repair and maintenance of Hb meters will be responsibility of vendor

Sr#	TECHNICAL SPECIFICATIONS / TERMS & CONDITIONS	DESCRIPTION
10	References	1 - 2 Local references with satisfactory reports should be provided
11	Wavelength	Multiple wavelength ranges from 450 nm to 750 nm
12	Measuring Time	Upto 5 seconds
13	Training of staff	Training of staff will be responsibility of vendor
14	Battery	Preferably have a rechargeable battery
15	Memory	More than 100 test results

Item: Platelets/ Plasma Mega Unit Kits:- (Apheresis Kit +ACD Solution/ Normal +Saline Fistula Needle)

UNSPSC: Blood bank apheresis and donor unit processing analyzer

Specifications / Requirements:

S#	TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS	DESCRIPTION
1	Apheresis System Description	· Automated cell separator
		· Continuous Flow System
		· Floor Top model with lockable wheels (casters)
2	Component Types	· Platelets (Mega Units)
		· Plasma Apheresis

S#	TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS	DESCRIPTION
3	Principle	· Incorporation of polyvinyl tubing that draws blood from the patient and moves it through centrifuges and/or filters to separate blood products.
		· The blood is then returned to the patient via tubing or is collected in bags, often suspended from a pole, for donation or disposal.
4	Detection System	· Must have patented optical detection system that continuously manages separated layers allowing the platelet and white blood cell layer to accumulate, preferably AIM System
		· Targeted components should be removed efficiently by system.
5	Hemolysis/ Red Cells sensor	There should be hemolysis/ red cells sensor for plasma line monitoring and platelets sensor on cell line
6	Centrifugation Speed	· 0 - 2200 RPM
7	Corporeal Volume	· Upto 185 ml
8	Data Management	· Prints reports or exports the data
		· Captures and stores procedure data
		· Tracks and records alarms during each of your procedures
		· Have capacity to store data upto 100 procedures
9	Patient comfort and safety	· Pump precision contributes to optimized fluid balance
		· Low volume tubing set that can accommodate smaller patients
		· Low operating noise

S#	TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS	DESCRIPTION
10	Incorporated seal safe system for sealing tubes	· Ready to use tubing sets designed to handle multiple procedure types.
		· Snap- in-place cassette should be loaded in few steps.
		· System should verify the correct tubing set.
11	Highly maneuverable system	· Must have large, durable wheels on pivoting casters with advanced wheel pedal
		· Telescoping IV pole that can be easily moved up or down and folding screen for moving and storage
12	Display	· Must have high resolution, colored touch screen
		· Must be user friendly
		· Should provide clear alarm messages that can be seen at a glance
13	References	There must be at least 2 analyzers with same offered model across the country in the tertiary care hospital or well reputed Blood Banks/ Institutes with highly satisfactory results (provide certificate) with reference address and telephone numbers.
14	Certification	· The company should be ISO certified.
		· Apheresis analyzer and kits must be FDA approved or CE marked or any equivalent certification. Preferably FDA approved. (Certificate must be provided).
15	Physical dimensions (HxWxD)	· Weight upto 130 Kg
		· Height: (lowered IV pole) upto 180 cm & (extended IV pole) upto 180 cm
		· Width upto 60 cm
		· Depth upto 90 cm

S#	TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS	DESCRIPTION
.	Floor space required upto 0.5 m square	
16	Temperature & Humidity	<ul style="list-style-type: none"> · Room temperature (18 - 32 OC). · Humidity 40 - 80 % RH.
17	Power supply & Battery back up	<ul style="list-style-type: none"> · 220 / 230 / 240 V 50 Hz · Built- in
18	Number of equipments	One system is required which will be installed in blood bank
19	Consumables & Accessories	All consumables and accessories will be the responsibility of vendor
20	Printout Feature	Analyzers should have built-in printout feature or attached external laser printer
21	UPS Backup	Analyzer must be supplied with preferably 15 minutes UPS (uninterrupted power supply) backup and other electrical connections. In case of any malfunctioning/ problem in UPS, it must be replaced immediately but not repaired. UPS batteries must be changed by the vendor as and when required.

S#	TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS	DESCRIPTION
22	Maintenance of equipments	Maintenance of equipment will be according to manufacturer booklet i.e. daily, weekly and monthly. Maintenance sticker to be applied on the analyzers and regularly updated. This will be documented in log book signed by the vendor engineer and checked by "Blood Bank Manager" and verified by "Incharge Blood Bank".
23	Consumables/ parts/ accessories will be provided by the vendor free of cost	Consumables (i.e. QC & cleaning solution)/ parts/ accessories and printer papers will be provided by the vendor free of cost. PIMS will not be liable for additional payments in respect of consumables/ parts/ accessories.
24	Repair, training & documentation of analyzer	All repairs of analyzer, training and documentation will be vendor's responsibility. This must be verified by consultant pathologist.
25	Performing tests from outside lab	In case of any breakdown/ failure of analyzers/ kits etc., company will be responsible to perform all pending tests from any well reputed outside lab. It's all expenses will be paid by company.
26	Up-gradation of analyzers	Analyzer will be upgraded to higher version as and when available by manufacturer if specifications and test cost remains same.
27	Transportation of Reagents	Transportation of the reagents to the hospital/ center according to standard protocol will be responsibility of vendor.
28	Service Workshop	Must have service workshop at Islamabad/ Rawalpindi with adequate engineering services and back up facilities with adequate spare parts, reagents and consumables.
29	Interfacing & Installation of computers with Analyzer	Installation of new latest computer & interfacing of analyzer with BTIS will be responsibility of vendor at the time of installation.

S#	TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS	DESCRIPTION
30	Maintenance of room temperature	Maintenance of room temperature will be the responsibility of vendor
31	Provision of technical services	Successful vendor will be responsible to depute one technical assistant in blood bank for performing apheresis procedures successfully and to avoid wastage of precious products

Item: ANTI-H LECTIN (BOMBAY ANTI SERA)

UNSPSC: Blood bank reagents or solutions

Specifications / Requirements:

S#	TECHNICAL SPECIFICATIONS	DESCRIPTION
1	Description	Blood grouping sera (Anti-H lectin) is used for detecting the Bombay phenotype by slide and tube test. It differentiates the Bombay phenotype (Oh) from ordinary group O blood.
2	Reagent Type	Anti-H Lectin (Plant lectin)
3	Source	Purified extract of Ulex europaeus seeds
4	Principle	Agglutinates RBCs carrying H antigen
5	Test Methods	Slide and tube agglutination techniques
6	Presentation	Ready-to-use liquid reagent

S#	TECHNICAL SPECIFICATIONS	DESCRIPTION
7	Storage	2-8°C
8	Shelf Life	Typically 24 months when stored properly
9	Pack Size	2 mL, 5 mL, 10 mL (manufacturer dependent)
10	Interpretation	Agglutination = H antigen present
		No agglutination = H antigen absent (suggestive of Bombay phenotype)
11	Sealed Lid Packing	Preferably sealed lid packing
12	References	Preferably have references of at least two tertiary care hospitals/ blood banks of Pakistan (Preferably local). Satisfactory reports with address and contact number required.
13	Certification	Preferably FDA approved or CE Marked or any other equivalent certification
14	DRAP registration	Preferably registered with DRAP

Item: ANTI-A1 LECTIN (DOLICHOSS BIFLORUS) ANTISERA

UNSPSC: Blood bank reagents or solutions

Specifications / Requirements:

S#	TECHNICAL SPECIFICATIONS	DESCRIPTION
1	Description	Anti-A1 Lectin (Dolichos biflorus) Antisera

S#	TECHNICAL SPECIFICATIONS	DESCRIPTION
3	Application	Detection of A1 antigen on human red blood cells for differentiation of A1 and A2 subgroups and investigation of ABO discrepancies
4	Source	Standardized extract of Dolichos biflorus seeds
5	Reactivity	Agglutinates A1 red blood cells; non-reactive with A2 and Group O red blood cells
6	Pack Size	10 mL or equivalent manufacturer standard pack
7	Storage Temperature	2-8°C (refrigerated)
8	Shelf Life	Minimum 12 months at the time of delivery
9	Quality Control	Each lot must be quality controlled and provide clear, specific agglutination with A1 cells
10	Sealed Lid Packing	Preferably sealed lid packing
12	Certification	Preferably FDA approved or CE Marked or any other equivalent certification
13	DRAP registration	Preferably registered with DRAP

Item: PAEDIATRIC BLOOD BAGS (QUINTUPLE)

UNSPSC: Blood unit collection bags

Specifications / Requirements:

S#	TECHNICAL SPECIFICATIONS/ TERMS & CONDITIONS	DESCRIPTION
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1	BLOOD BAG CAPACITY	· Quintuple Blood Bag 450 ml
2	PILOT BLOOD BAG CAPACITY	· Having 4/5 pilot bags of 100 - 200 ml each
3	MATERIAL USED	· Blood collection bag Made up of DEHP (Di-2-ethylhexyl phthalate) plasticized PVC (polyvinyl chloride), collapsible non-vented sterile containers complete with collecting tube for completely closed system to avoid the chances of contamination.
4	DESIGN AND SHAPE	<ul style="list-style-type: none"> · Flexible pre-sterilized · Pyrogen free · Nontoxic, non-hemolytic, biocompatible material · No risk of contamination and air embolism (close system) with leaks proof seals · Slit on both sides of the bags should be enough to accommodate 5-10 ml test tubes. · The capacity of the bag should be enough to prevent any ballooning/ rupture of the bag from seam when it is filled up with the requisite volume of blood
5	TUBING OF BAG	<ul style="list-style-type: none"> · Flexible non kinking · Non sticking · Transparent · Leak proof · The minimum length of tubing from primary bag to the needle should be 80 cm. · The tube should have multiple printed ID/Segment number. The number should be legible and clear. · A clamp should be provided for closed system

6	NEEDLE	· 16/ 17 gauge ultra-thin walled, siliconized and straight
		· Sharp, regular and smooth margins and beveled tip
		· Rust proof
		· Tightly fixed with hub covered with sterile guard
		· Hermetically sealed
		· The needle should not separate from the tube at any point of time, especially while removing it from the vein for the donor safety
7	EXTERNAL PORT	· Tamper proof and should not be re-capped
		· Easily accessible
8	PACKAGE	· Protective packaging (Aluminium) eliminating microbial contamination on surface maintaining the contents of the bag
		· Easy to handle
9	ANTICOAGULANT	· CPDA-1
		· The quantity of anticoagulant (63 ml)
		· Clear & colorless
		· No discoloration on storage at room temperature
		· Manufacturer to supply anticoagulant quality check certificate
10	AIR BUBBLES	· Air bubble content in blood bag should be less than 15 ml
11	LABEL	· Non peel-off
		· Heat sealed/pressure embossed labels

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°C
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transparent
adhesive

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Following
information
must
be
printed
(permanent)
on
label



of
blood
bag

i.
Date
of
manufacturing

ii.
Date
of
expiry

iii.
Lot
number

iv.
Batch
number

v.
Donor
number

vi.
ABO
group



vii.
Rh
type

viii.
Serology
report

ix.
Manufacturer
name
and
address

x.
Anticoagulant
symbol
and
chemical
formulation



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The
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12	RESISTANCE TO DISTORTION	· Bag shall with stand an acceleration of 5000 g for 30 min at temperature 4°C to 24°C
		· Bag should be able to withstand temperature upto-80°C without breakage
13	DIVERSION POUCH	· Blood bags with diversion pouch and needle guard will be preferred
14	CERTIFICATION AND APPROVAL	· CE marking or any equivalent certification
		· Registered with DRAP
		· The international marking certificates must be valid; clearly mentioning the brand name of product being referred and name of the company with address given on pack and must be verifiable at company websites
		· Products having doubtful or invalid international certificate will be rejected
15	REFERENCES	· Must have references of at least three tertiary care hospitals/ blood banks of Pakistan. Satisfactory reports with address and contact number required.
16	SOLE DISTRIBUTION/ LOCAL OFFICES	· The company should have authorization for sole distribution and local sold distribution office and available stock.

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. "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;
- c. "The Contract" means an agreement enforceable by law;
- d. "The Contract Price" means the price payable to the Bidder under the Contract for the full and proper performance of its contractual obligations;
- e. "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;
- f. "GCC" means the General Conditions of Contract contained in this section;
- g. "SCC" means the Special Conditions of Contract by which the GCC may be amended or supplemented;
- h. "Day" means calendar day unless indicated otherwise.
- i. "Effective Date" means the date on which this Contract comes into force and effect.
- j. "The Bidder" means the individual or corporate body whose Bids to provide the Goods has been accepted by the Procuring Agency;
- k. "The Project Site," where applicable, means the place or places named in Bids Data Sheet and technical Specifications;
- l. "Government" means the Government of Pakistan;
- m. "Subcontractor" means any entity to which the Bidder subcontracts any part of the Goods.
- n. "Service" means any object of procurement other than goods or works;
- o. "Party" means the Procuring Agency or the Bidder, as the case may be, and "Parties" means both of them;
- p. "Foreign Currency" means any currency other than the currency of the country of the Procuring Agency;

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

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

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

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. Delivery/Location

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

7. Authorized Representatives / Authority of Member in charge

7.1 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 Services

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 Services, 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 15** 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 of the Services, 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.

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 15**.

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 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

23.2 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.

23.3 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.

23.4 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**.

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.

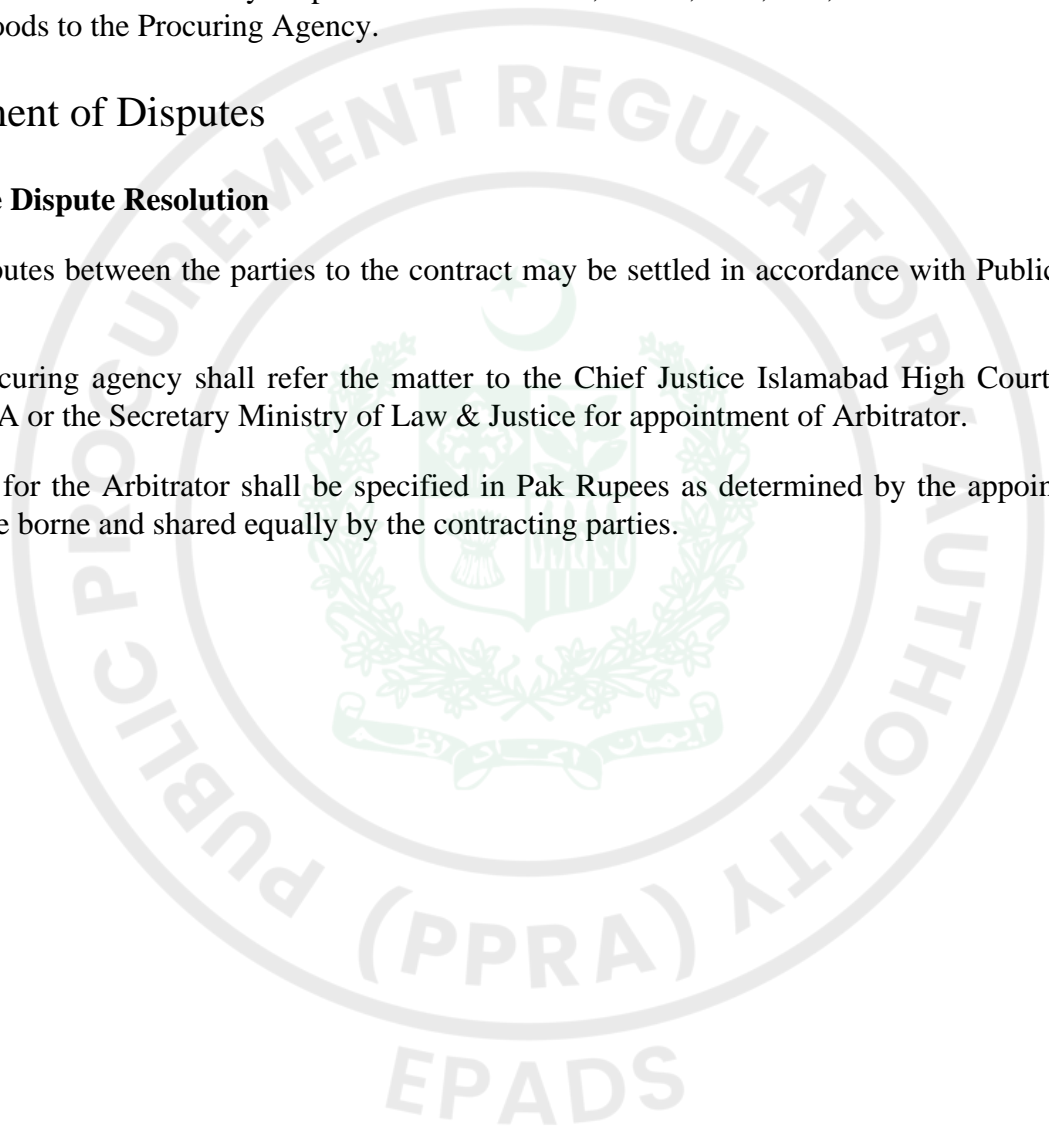
H. Settlement of Disputes

37. Alternate Dispute Resolution

37.1 The disputes between the parties to the contract may be settled in accordance with Public Procurement Rules, 2004.

37.2 The procuring agency shall 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.





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: Purchase Department (Pakistan Institute of Medical Sciences (PIMS)), Assistant Director PIMS Hospital, Sector G-8/3., Islamabad Capital Territory

The Supplier is:

The title of the subject procurement is: SUPPLY OF BLOOD BAGS, ANTI SERAS, KITS AND DISPOSABLE ITEMS, TO BE USED IN BLOOD BANK AT PIMS ON ANNUAL CONTRACT BASIS, FOR THE FY 2026-2027

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:

Procuring Agency:

Purchase Department (Pakistan Institute of Medical Sciences (PIMS)), Assistant Director
PIMS Hospital, Sector G-8/3., Islamabad Capital Territory
+92-312-345-6789
adpurchase@pims.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:

Purchase Department (Pakistan Institute of Medical Sciences (PIMS)), Assistant Director
PIMS Hospital, Sector G-8/3., Islamabad Capital Territory
+92-312-345-6789
adpurchase@pims.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 **0.50%** to **10.00%** 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:

The amount of performance guarantee shall be **2.00%** of the contract price in acceptable form of **Pay Order, Call at Deposit, Bank Guarantee**

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;

Insurance Certificate;

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

- 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 by sole arbitrator in accordance with Arbitration Act 1940. The arbitrator shall be appointed by mutual consent of the both parties. The Arbitration shall take place in Islamabad, Pakistan and proceedings will be conducted in English language.
- 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.: **P45305**

To: **Purchase Department (Pakistan Institute of Medical Sciences (PIMS)), Assistant Director PIMS Hospital, Sector G-8/3., 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.



Contract Form

SECTION IX: CONTRACT FORMS

THIS AGREEMENT made the ____ day of _____ 20____ between **Purchase Department (Pakistan Institute of Medical Sciences (PIMS)), Assistant Director PIMS Hospital, Sector G-8/3., 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., **SUPPLY OF BLOOD BAGS, ANTI SERAS, KITS AND DISPOSABLE ITEMS, TO BE USED IN BLOOD BANK AT PIMS ON ANNUAL CONTRACT BASIS, FOR THE FY 2026-2027 (P45305)** 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.

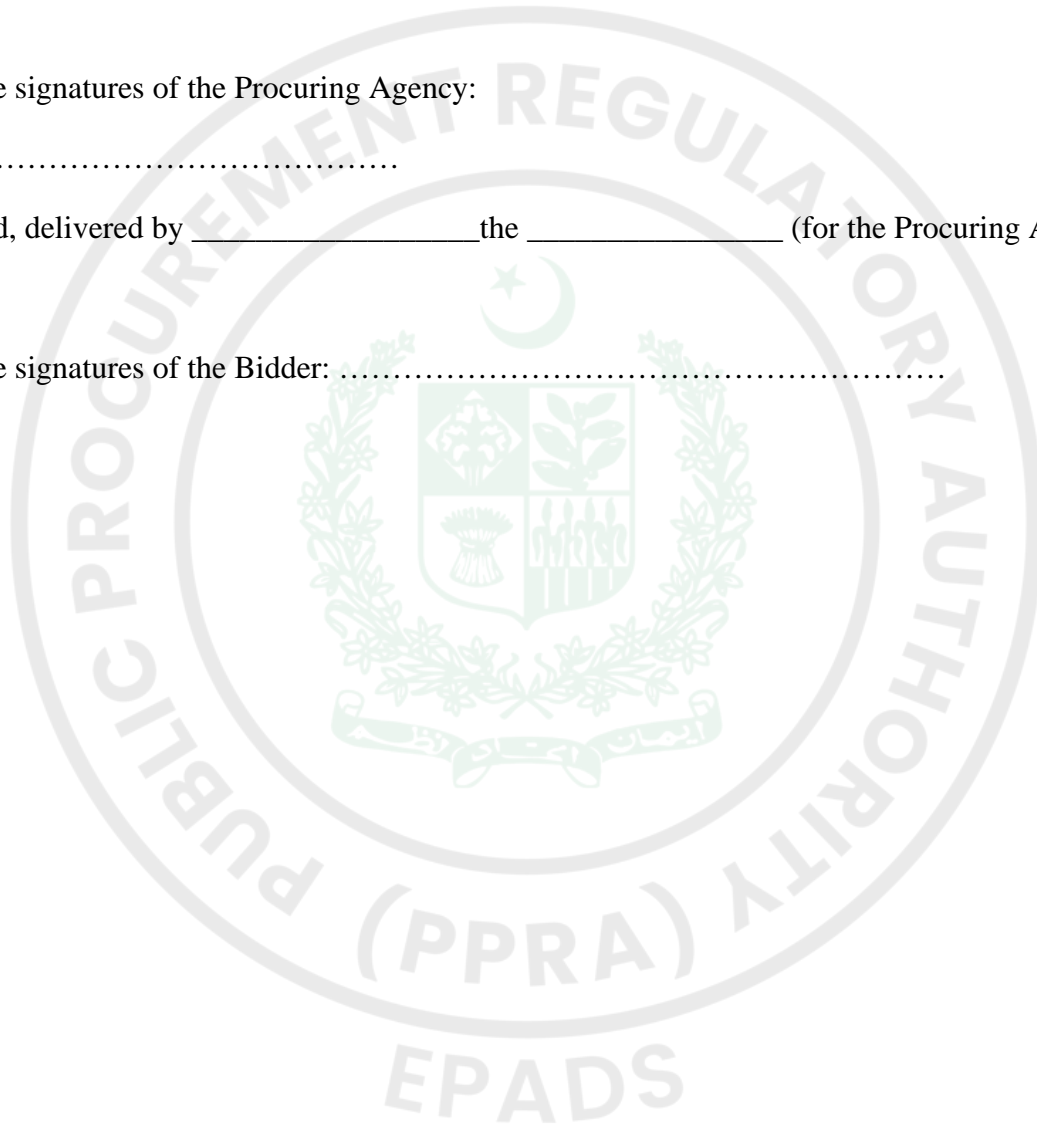
Signed, sealed, delivered by _____ the _____ (for the Procuring Agency)

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: **Purchase Department (Pakistan Institute of Medical Sciences (PIMS)), Assistant Director PIMS Hospital, Sector G-8/3., 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



Required Technical Specifications

Technical Table Inputs (Vendor)

Name of Items	Brand Name	Manufacturer Name	DRAP Registration No
Blood Collecting Bags (Single) without Set) (Item/Job/Position)			
Blood Collecting Bags with diversion pouch triple without set (Item/Job/Position)			
Blood Collecting Bags without diversion pouch (Triple without Set) (Item/Job/Position)			
Blood Collecting Bags with diversion pouch (Quadruple without Set) (Item/Job/Position)			
Blood Collecting Bags without diversion pouch (Quadruple without Set) (Item/Job/Position)			
Blood Transfusion Set (Item/Job/Position)			
Blood Grouping Sera Anti-A (Item/Job/Position)			
Blood Grouping Sera Anti-B (Item/Job/Position)			

Name of Items	Brand Name	Manufacturer Name	DRAP Registration No
Blood Grouping Sera Anti-D (Item/Job/Position)			
Leukocyte Depletion Filters (RCC Pre-storage) (Item/Job/Position)			
Leukocyte Depletion Filters (RCC Bedside) (Item/Job/Position)			
Leukocyte Depletion Filters (Platelets Filters) (Item/Job/Position)			
Disposable Pasture Pipettes (Item/Job/Position)			
Test Tube 3" Glass (Item/Job/Position)			
NAT screening kit (Item/Job/Position)			
Coomb's Sera (Item/Job/Position)			
Bovine Albumin (Item/Job/Position)			
LMB HGB Test Strips, with free placement of Hb Meter (Item/Job/Position)			
Platelets/ Plasma Mega Unit Kits:- (Apheresis Kit +ACD Solution/ Normal +Saline Fistula Needle) (Item/Job/Position)			

Technical Specification Parameters for T.S # 1-5

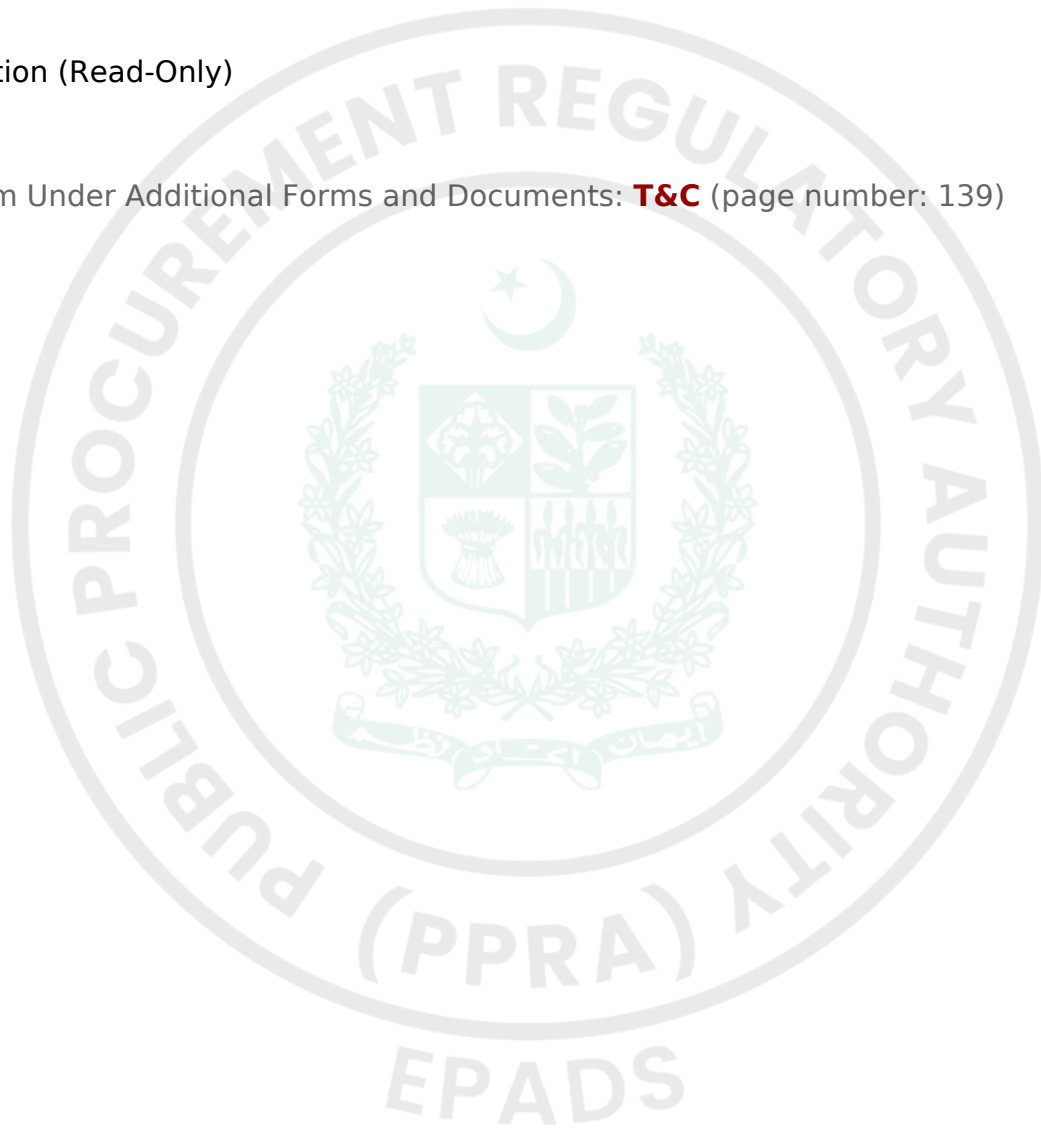
Information (Read-Only)

See Form Under Additional Forms and Documents: **Technical Specification Parameters for T.S # 1-5** (page number: 138)

T&C

Information (Read-Only)

See Form Under Additional Forms and Documents: **T&C** (page number: 139)





Procurement Forms







Additional Forms and Documents

TS#. 1-5. PARAMETERS FOR TECHNICAL EVALUATION FOR THE TENDER OF BLOOD BAGS FOR PIMS BLOOD BANK (FY 2026 – 2027)

S#	GENERAL SPECIFICATIONS PARAMETERS	DESCRIPTION				
1	Brand Name					
2	Country of origin					
3	Sample Lot Number					
4	Sample Expiry date					
5	Total Capacity					
6	CE marked or any equivalent certification					
7	Registered with DRAP					
8	References (At least 03)					
9	Sole Distributor/ Local Office					
10	None of the batch of the product being quoted has been declared spurious by any laboratory anywhere in Pakistan					
11	Firm/ company has not been blacklisted / penalized by any government department/ hospitals/ international agencies and NGOs					
	TECHNICAL SPECIFICATIONS PARAMETERS	Poor (Marks = 1)	Below average (Marks = 2)	Average (Marks = 3)	Good (Marks = 4)	Excellent (Marks = 5)
12	Material used					
13	Design and Shape					
14	Tubing of bag					
15	Needle					
16	External port					
17	Package					
18	Anticoagulant					
19	Air Bubbles					
20	Label					
21	Resistance to distortion					
22	Hemolysis					
23	HCT					
24	Platelets Yield					
TOTAL MARKS (S# 12 to 24)						65

NOTE:

1. Clause# 6, 7, 8, 9, 10 & 11 are mandatory
2. Only those bidders will be technically qualified who will score ≥ 70 % marks.

S. #	☛ Special Instructions for Vendors (Please read carefully)
1	Technical brochures /Literature brand country of origin of quoted item/material must also be uploaded with the bid where required.
2	All vendors will drop the original Pay Order/ CDR on account of bid security along with original Affidavit / undertaking along with copy of Tender Receipt in envelop duly sign and seal by the firms authority/focal person in the Drop Box placed at Purchase Department before 11:00 hrs on the specified date as mentioned in the advertisement. (Cross Cheque / Open Cheque / Bank Guarantee not acceptable).
3	The Vendors / Firms quoting two or more brands of item in one line / column shall be liable to rejection

A. GENERAL CONDITIONS:

1.	The contract will be valid for <i>Financial Year 2026-2027</i>; however, the contract can be extended after the mutual consent of both parties as PPRA Rules.
2.	Bid validity is 180-days from the date of opening the tender.
3.	Earnest money will be forfeited if a bidder withdraws his bid during the period of bid validity.
4.	The supply will be made within 35-days after the issuance of award letter.
5.	Initially the supplier/bidder will submit call deposit/bid money in the name of Executive Director PIMS. After the issuance of LOI/ Award letter the supplier will replace the deposit at call / bid money with 2% performance guarantee of total amount of the awarded items. It will be released on completion of successful contract on producing of NOC from Store Keepers Medical, Pharmacist Incharge stores and Chief Pharmacist.
6.	The rates quoted in the financial bid Performa i.e. items List will be final, and no change therein will be accepted after the tenders are opened
7.	The approved rates will be valid for the whole contract period and there will be no increase in rates and there is no escalation clause
8.	The successful Vendors / Firms will never stop the daily supply even their bills /payment are delayed due to some unavoidable circumstances
9.	Income tax will be deducted according to the Government rules at source Supplier has to submit any other liable tax receipts f applicable
10.	All items will be received on FOR basis, PIMS Islamabad.
11.	All the regulations / rules framed / enforced by the Federal Government / PIMS from time to time will be binding upon the firms.
12.	In case of any dispute, Executive Director PIMS the Competent Authority of PIMS will be the final Authority.
13.	The Competent Authority of PIMS reserves the right to accept or reject tender as per PPRA rules.
14.	Letter of Intent will be issued to successful /most advantageous bidder in terms of lowest bidder. The approved bidder will submit the following documents/performance guarantee within 07-days positively. <ul style="list-style-type: none"> i) Integrity Pact on stamp paper worth of Rs.100/- if the contract cost is more than 10.00million. ii) 2% Performance Guarantee of total amount of awarded items in the name of Executive Director PIMS Islamabad from any scheduled bank of Pakistan.

	After the receiving the above mentioned documents and performance guarantee, <u>the bid money of firm will be released and proper award letter will issued accordingly.</u>
--	---

B. SPECIFIC TERMS & CONDITIONS:

1	The kits/Chemicals shall be accompanied by the necessary warranty where applicable in accordance with the provision of the Drugs Act 1976 and rules framed therein. (However, the warranty will be supplied at the time of delivery).
2	Thermoliable items e.g. Lab kits & Chemicals. Vaccines, Sera insulin, Ophthalmic preparations and infusion will be supplied under specific storage conditions and the supplier shall be responsible to maintain the cold chain. Contrary to this the delivery will not be accepted and the Hospital will not be responsible for any inconvenience.
3	Bills of the supplies will be submitted with the supply/delivery challan. For late submission of bills, Institute shall not be responsible for delay in payment.
4	The samples of product/kits/chemicals reagents shall be provided for the purpose of quality evaluation of specifications:- a) The committee is authorized to approve a product without sample presentation if not required, by virtue of its well-known brand name. b) The committee reserves the right to reject a product if bidder is unable to present the sample if desired so. Such an offer will be ignored/ rejected and the lowest offer will be decided among the rest of the bidder. c) Failure to provide sample(s) action will be taken against firm as penalty clause # 6 of the terms & conditions of the tender.
5	The preference shall be given to the manufacturers directly participating in the tender, however in case of authorized distributor the recent and original distribution letter shall be required. In case of Importer, documents of Sole Agent for Pakistan, duly verified from the Country of Origin shall be required.

C. RESPONSIBILITIES

1	In case of any discrepancy, less weight, short supply etc the supplier will be held responsible.
2	All items supplied will be in accordance with the Drugs Act 1976 "Labeling and packaging rules".
3	The following words in shape of insignia (Large size) should be printed with undeletable ink in bold letter on each bottle/strip/label outer packing, inner packing and ampoules etc (NOT FOR SALE, FOR PIMS USE ONLY).
4	Packing should be worthy of transportation by Road, Sea, or mail as to ensure their contents being free from loss of damage due to faulty packing.
5	The supplier once awarded a product will be responsible to keep a contract with the organization for day to day supply orders, within due time. Non-compliance to a registered posted letter on given address, Telephone number or E.mail will be considered non-compliance of the supply order as per clause.
6	The bidder will be responsible to supply the quality products.
7	Port of shipment certificate etc, if required.
8	Verification of documents by the related embassy wherever required.
9	All reagents and chemicals should have expiry on all individual bottles where ever applicable.

11	The reference list Institutions where the equipment and / or reagents are being used will be provided by the company.
12	The company should also provide ISO Certification whenever required.

D. PENALITIES

1	Test / analysis of drugs from Government Drugs Testing Laboratory will be done according to the prescribed procedure as per Drug Act, 1976. If a drug is found substandard, adulterated etc, the same will not be returned to the supplier and will be seized OR destroyed by the Inspector of Drugs as the case may be. The payment of defective drugs will not be made to the supplier. The supplier will be responsible to provide the fresh stock of standard quality against the confiscated stock within 20-days against the quantity or amount equivalent to defective goods will be deducted from the bills of the firm. Further action against the firm will be taken as per Drugs Act 1976 and rules framed thereunder.
2	The manufacturer/importer. Sole distributors of sub-standard, adulterated or contaminated medicine can be blacklisted by the competent authority, or as per judgment of the Drugs Court or any other Authority whose decision will be final and in accordance with the offence and hence their earnest money will not be released till the case is decided by the Drugs Court or any other relevant authority.
3	No items will be accepted having expiry less than 75% of its shelf life in case of locally manufactured and 60% for the imported items at the time of delivery. In case of expiry of drugs, the supplier shall have to replace the quantity thus expired and under taking regarding this shall be given at the time of supply along with delivery challan. Warranty of drugs will have to be provided at the time of delivery of drugs / items. In case of expiry of drugs, the supplier shall have to replace the quantity thus expired, with long expiry batch, three month before the date of its expiry.
4	<ul style="list-style-type: none"> • At any instance, if prices quoted in the bills are found higher, than Trade Price approved by the Ministry of Health, the Institute has the right to impose penalty as per decision of the Executive Director PIMS. • Repletion of incomplete supplies are delay in services will render the contract liable to penalty as per penalties clause No.5.
5	<p><u>Non-compliance of supply order: -</u></p> <p>If the supply order is not completed within 35-days one or more of the following penalties can be applied against the firm according to the gravity of situations: -</p> <ol style="list-style-type: none"> a) A penalty @ 0.5% per day of the amount of pending item(s) of the supply order shall be imposed. b) Risk purchase will be made at the cost of supplier and amount will be deducted from the bills of supplier. c) 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. d) Total or partial earnest money (CDR)/ Performance Guarantee will be forfeited. e) The firm will be debarred as per the PPRA rules. f) 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.
6	<u>Non provide of samples:-</u>

	<p>Failure to provide samples of the quoted item(s) for evaluation will be considered a non-serious commitment on the part of bidder, it will be treated as under:-</p> <p>a) Rs.5,000/- penalty for failure to provide each sample in the quoted bid, up to five sample.</p> <p>b) If failed to provide samples of more than five items Rs.10,000/- penalty for each sample.</p> <p>c) If failed to provide samples of more than ten items complete CDR will be forfeited and entire bid will be considered for rejection.</p>
7	<p>After submission of a bid any request by the bidder for change in the price or any content of the bid will be considered as an act of non-compliance of supply order and will be dealt as per penalty clause # 5.</p>

I/We have read and agree with the above-mentioned terms and conditions.

Name of Firm & Owner _____

Signature with Stamp _____

Correct Postal Address _____

Landline Telephone/Mobile Number _____

Valid email address _____