**BIDDING DOCUMENTS**

PROCUREMENT OF

**DISPOSABLE ITEMS**

**FOR CATH LAB, STROKE & ELECTROPHYSIOLOGY DEPARTMENT**

FOR THE FINANCIAL YEAR 2023-24

**THROUGH FRAMEWORK CONTRACT**



**RAWALPINDI INSTITUTE OF CARDIOLOGY**

**RAWAL ROAD, RAWALPINDI**

**Phone No: 051-9281111-20**

**Fax No: 051-9281357**

**E-Mail:** **purchaseric272@gmail.com**

**Table of Contents**

Contents

[PAGE MARKING / INDEX CERTIFICATE 4](#_Toc130206950)

[PAPERS (CHECK LIST) 6](#_Toc130206951)

[BID DATA SHEET 7](#_Toc130206952)

[SECTION I 8](file:///E%3A%5C2023-24%20TENDER%5CBIDDING%20DOCUMENTS%202023-24%5C01%29%2008-04-23%20-%20%20Bidding%20Documents%20for%20Medicine%202023-24.docx#_Toc130206953)

[INVITATION TO BID 8](file:///E%3A%5C2023-24%20TENDER%5CBIDDING%20DOCUMENTS%202023-24%5C01%29%2008-04-23%20-%20%20Bidding%20Documents%20for%20Medicine%202023-24.docx#_Toc130206954)

[LETTER OF INVITATION 9](#_Toc130206955)

[SECTION II 10](file:///E%3A%5C2023-24%20TENDER%5CBIDDING%20DOCUMENTS%202023-24%5C01%29%2008-04-23%20-%20%20Bidding%20Documents%20for%20Medicine%202023-24.docx#_Toc130206956)

[INSTRUCTINO TO BIDDERS (ITB) 10](file:///E%3A%5C2023-24%20TENDER%5CBIDDING%20DOCUMENTS%202023-24%5C01%29%2008-04-23%20-%20%20Bidding%20Documents%20for%20Medicine%202023-24.docx#_Toc130206957)

[1. Scope of Bid 11](#_Toc130206958)

[2. Source of Funds 11](#_Toc130206959)

[3. Eligible Bidders 11](#_Toc130206960)

[4. Corrupt or Fraudulent Practices and Mechanism to Debar/Blacklist the Defaulted Bidder 11](#_Toc130206961)

[5. Eligible Goods and Services 12](#_Toc130206962)

[6. Cost of Bidding 12](#_Toc130206963)

[7. Bidding for Selective Items 13](#_Toc130206964)

[THE BIDDING PROCEDURE 13](#_Toc130206965)

[8. The Governing Rules 13](#_Toc130206966)

[9. Applicable Bidding Procedure 13](#_Toc130206967)

[THE BIDDING DOCUMENTS 13](#_Toc130206968)

[10. Contents of the Bidding Documents 13](#_Toc130206969)

[11. Clarification(s) on Bidding Documents 14](#_Toc130206970)

[12. Amendment(s) to the Bidding Documents 15](#_Toc130206971)

[PREPARATION OF BIDS 15](#_Toc130206972)

[13. Language of Bids 15](#_Toc130206973)

[14. Documents comprising the Bids 15](#_Toc130206974)

[15. Bid Price. 15](#_Toc130206975)

[16. Bid Currencies. 16](#_Toc130206976)

[17. Samples 16](#_Toc130206977)

[18. Documentation on Eligibility of Bidders 16](#_Toc130206978)

[19. Documentation on Eligibility of Goods 16](#_Toc130206979)

[20. Bid Security. 16](#_Toc130206980)

[21. Bid Validity 17](#_Toc130206981)

[22. Format and Signing of Bids. 17](#_Toc130206982)

[SUBMISSION OF BIDS 17](#_Toc130206983)

[23. Sealing and Marking of Bids 17](#_Toc130206984)

[24. Deadline for Submission of Bids 18](#_Toc130206985)

[25. Late Bids 18](#_Toc130206986)

[26. Withdrawal of Bids 18](#_Toc130206987)

[OPENING AND EVALUATION OF BIDS 18](#_Toc130206988)

[27. Opening of Bids by the Procuring Agency 18](#_Toc130206989)

[28. Clarification of Bids 19](#_Toc130206990)

[29. Preliminary Examination 19](#_Toc130206991)

[30. Evaluation of Bids 19](#_Toc130206992)

[31. Qualification of Bidder 19](#_Toc130206993)

[32. Rejection of Bids 20](#_Toc130206994)

[33. Re-Bidding 20](#_Toc130206995)

[34. Announcement of Evaluation Report 20](#_Toc130206996)

[35. Contacting the Procuring Agency 20](#_Toc130206997)

[36. Grievance Redressal 21](#_Toc130206998)

[37. Acceptance of Bid and Award Criteria 21](#_Toc130206999)

[38. Procuring Agency’s Right to vary quantities at the time of Award 21](#_Toc130207000)

[39. Notification of Award 21](#_Toc130207001)

[40. Limitation on Negotiations 22](#_Toc130207002)

[41. Signing of Contract 22](#_Toc130207003)

[SECTION III 23](file:///E%3A%5C2023-24%20TENDER%5CBIDDING%20DOCUMENTS%202023-24%5C01%29%2008-04-23%20-%20%20Bidding%20Documents%20for%20Medicine%202023-24.docx#_Toc130207004)

[SCHEDULE OF REQUIREMENT & TECHNICAL SPECIFICATION 23](file:///E%3A%5C2023-24%20TENDER%5CBIDDING%20DOCUMENTS%202023-24%5C01%29%2008-04-23%20-%20%20Bidding%20Documents%20for%20Medicine%202023-24.docx#_Toc130207005)

[Schedule of Requirements: 24](#_Toc130207006)

[ANNUAL DEMAND OF MEDICINE, SRUGICAL DISPOSABLE FOR THE FINANCIAL YEAR 2023-24 25](#_Toc130207007)

[(THROUGH FRAMEWORK CONTRACT) 25](#_Toc130207008)

[SECTION IV 39](file:///E%3A%5C2023-24%20TENDER%5CBIDDING%20DOCUMENTS%202023-24%5C01%29%2008-04-23%20-%20%20Bidding%20Documents%20for%20Medicine%202023-24.docx#_Toc130207009)

[EVALUATION CRITERIA 39](file:///E%3A%5C2023-24%20TENDER%5CBIDDING%20DOCUMENTS%202023-24%5C01%29%2008-04-23%20-%20%20Bidding%20Documents%20for%20Medicine%202023-24.docx#_Toc130207010)

[EVALUATION CRITERIA 40](#_Toc130207011)

[1. COMPULSORY PARAMETERS: 40](#_Toc130207012)

[2. ORDINARY PARAMETERS: 41](#_Toc130207013)

[SECTION V 43](file:///E%3A%5C2023-24%20TENDER%5CBIDDING%20DOCUMENTS%202023-24%5C01%29%2008-04-23%20-%20%20Bidding%20Documents%20for%20Medicine%202023-24.docx#_Toc130207014)

[BID FORM 43](file:///E%3A%5C2023-24%20TENDER%5CBIDDING%20DOCUMENTS%202023-24%5C01%29%2008-04-23%20-%20%20Bidding%20Documents%20for%20Medicine%202023-24.docx#_Toc130207015)

[BID COVER SHEET 44](#_Toc130207016)

[BID FORM 1 45](#_Toc130207017)

[BID FORM 2 46](#_Toc130207018)

[BID FORM 3(A) 47](#_Toc130207019)

[BID FORM 3(B) 48](#_Toc130207020)

[BID FORM 4 49](#_Toc130207021)

[BID FORM 5(A) 50](#_Toc130207022)

[BID FORM 5(B) 51](#_Toc130207023)

[BID FORM 6 52](#_Toc130207024)

[SECTION VI 53](file:///E%3A%5C2023-24%20TENDER%5CBIDDING%20DOCUMENTS%202023-24%5C01%29%2008-04-23%20-%20%20Bidding%20Documents%20for%20Medicine%202023-24.docx#_Toc130207025)

[DRAFT STANDARD CONTRACT 53](file:///E%3A%5C2023-24%20TENDER%5CBIDDING%20DOCUMENTS%202023-24%5C01%29%2008-04-23%20-%20%20Bidding%20Documents%20for%20Medicine%202023-24.docx#_Toc130207026)

[Contract Form 54](#_Toc130207027)

[ANNEX: A 58](#_Toc130207028)

[SCHEDULE OF REQUIREMENTS: 58](#_Toc130207029)

[ANNEX. B 59](#_Toc130207030)

[Special Conditions of the Contract & Technical Specifications 59](#_Toc130207031)

[ANNEX. C 61](#_Toc130207032)

[PRICE SCHEDULE SUBMITTED BY THE BIDDER 61](#_Toc130207033)

[ANNEX. D 62](#_Toc130207034)

[PURCHASER’S NOTIFICATION OF AWARD 62](#_Toc130207035)

[ANNEX. E 63](#_Toc130207036)

[PURCHASE ORDER 63](#_Toc130207037)

[ANNEX. F 64](#_Toc130207038)

[PAYMENT SCHEDULE 64](#_Toc130207039)

[ANNEX. G 65](#_Toc130207040)

[GENERAL CONDITIONS OF CONTRACT (GCC) 65](#_Toc130207041)

# PAGE MARKING / INDEX CERTIFICATE

# PAGE MARKING / INDEX CERTIFICATE

I Mr. / Miss / Mrs. \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ do hereby certify on the behalf of M/S (firm name)\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ that the bidding documents submitted for tender of \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ contain total pages \_\_\_\_\_\_\_\_\_\_\_.

Moreover, the page marking is done and index has been prepared which is marked as page no \_\_\_\_\_\_\_\_\_\_.

Name of authorized person \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

Designation \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

CNIC No. \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

Mailing Address \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

Contact No. (Land Line) \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

Contact No. (Mobile) \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

E-mail Address \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

**NOTE:**

***Technical Bid should be properly tagged / binding / page numbering, otherwise the procuring agency has right to reject the bid and its decision will be final which cannot be challenged in any court of law.***

**Signature: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**

**Stamp (Firm) \_\_\_\_\_\_\_\_\_\_\_\_**

**Dated \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**

I Mr. / Miss / Mrs. \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ do hereby certify on the behalf of M/S (firm name)\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ that the bidding documents submitted for tender of \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ contain total pages \_\_\_\_\_\_\_\_\_\_\_.

Moreover, the page marking is done and index has been prepared which is marked as page no \_\_\_\_\_\_\_\_\_\_.

Name of authorized person \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

Designation \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

CNIC No. \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

Mailing Address \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

Contact No. (Land Line) \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

Contact No. (Mobile) \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

E-mail Address \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

**NOTE:**

***Technical Bid should be properly tagged / binding / page numbering, otherwise the procuring agency has right to reject the bid and its decision will be final which cannot be challenged in any court of law.***

**Signature: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**

**Stamp (Firm) \_\_\_\_\_\_\_\_\_\_\_\_**

**Dated \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**

**COPY OF CNIC (ATTACH HERE)**

**Name :\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**

**Father Name: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**

**CNIC No. \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**

**Address. \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**

**(Mandatory to attach copy of CNIC)**

**ORIGINAL TENDER PURCHASE RECEIPT (ATTACH HERE)**

**Tender Fee Receipt No. \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**

**Date: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**

**Amount Rs. \_\_\_\_\_\_\_\_\_\_\_\_\_\_**

**(Mandatory to attach Original Purchase Receipt)**

**COPY OF BID SECURITY (ATTACH HERE)**

**Bank Name: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**

**Call Deposit Receipt / Bank Guarantee No: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**

**Date \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**

**Amount of Bid Security: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**

**(Mandatory to attach copy of bid security)**

# PAPERS (CHECK LIST)

|  |  |  |  |
| --- | --- | --- | --- |
| **S.#** | **Detail** | **Yes / No** | **Page No** |
|  | Index & page marking certificate by the bidder |  |  |
|  | Attested copy of CNIC |  |  |
|  | Original purchased receipt of bid documents |  |  |
|  | Copy of Bid Security 2% of estimated price |  |  |
|  | Technical Proposal / Offer of quoted items (on firm letter head) |  |  |
|  | Bidding Documents signed & stamped by bidder (all pages) |  |  |
|  | NTN Certificate |  |  |
|  | GST Certificate |  |  |
|  | Professional Tax Certificate |  |  |
|  | Drug Manufacturing / Sale License |  |  |
|  | Drug Registration Certificate (DRC) |  |  |
|  | Good Manufacturing Practices (GMP) Certificate(if applicable) |  |  |
|  | Embassy Attested Free sale certificate (if applicable) |  |  |
|  | Batch Capacity of manufacturer for the quoted item / product |  |  |
|  | Documentary evidence of product experience in market (attested copy of relevant documents) |  |  |
|  | Attested copy of list of client (where product is available in open market) |  |  |
|  | Audit Balance Sheet (Last Financial Year) |  |  |
|  | Fresh Samples of quoted Items (3 packs of each quoted item)  |  |  |
|  | **Affidavit (on judicial paper)** |  |  |
|  | Affidavit regarding Good / services supplied are to be provided by bidder in accordance with Government instruction |  |  |
|  | Affidavit regarding firm is not legally penalized |  |  |
|  | Firm has not provided these stocks below the price if so, he I legally bound to pay rate difference if the bid is accepted  |  |  |
|  | Affidavit / Undertaking Regarding Offered prices are not more than trade price. |  |  |
|  | Affidavit from the sole agent(s) that their product(s) are freely available with same brand name in the country of manufacturer & is safe for human consumption |  |  |
|  | Affidavit / Undertaking Regarding Non Cancellation / Suspension of Drug Registration of quoted product of the bidder by Drug Regulatory Authority of Pakistan within last two years |  |  |
|  | Affidavit / Undertaking Regarding Non Declaration of Spurious / Adulterated batch by DTLs of Punjab / any Competent Lab of quoted item within last two years |  |  |
|  | Affidavit / Undertaking Regarding Firm is not blacklisted |  |  |

**Signature: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**

**Stamp (Firm) \_\_\_\_\_\_\_\_\_\_\_\_**

**Dated \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**

# BID DATA SHEET

|  |  |  |
| --- | --- | --- |
| **ITB Reference** | **Description** | **Detail** |
| N/A | Bid reference number | **RIC/PO/3054/23, Dated 27-03-2023** |
| N/A | Commencement of sale ofBidding Documents | From the date of advertisement, onall working days during officehours |
| N/A | Last date & time of sale ofBidding Documents | **10-04-2023, 01:00 PM** |
| N/A | Pre-Bid Meeting | **03-04-2023, 11:00 AM** |
| ITB Clause 24 | Last date and time for thereceipt of bids | **11-04-2023, , 11:00 AM** |
| ITB Clause 27 | Date, time and venue ofopening of technical bids | **11-04-2023, 11:30 AM****At MS Office in RIC, Rawalpindi** |
| ITB Clause 16 | Bid currency | PKR on free delivery to Consignee’s end basis including all Ex-work, Transportation, Storage Charges till the destination. |
| ITB Clause 13 | Language of bid | English or Urdu |
| ITB Clause 20 | Amount of bid security | 2% of the Estimated Price (Estimated price Mention against each item of the list) |
| ITB Clause 21 | Bid validity period | 120 Days |
| ITB Clause 09 | Bidding procedure | Single Stage – Two Envelopbidding procedure |
| ITB Clause 27 | Address for Communication  | **Medical Superintendent****Rawalpindi Institute of Cardiology****Rawal Road, Rawalpindi****051-9281111-20****Purchaseric272@gmail.com** |

# SECTION I

# INVITATION TO BID

## LETTER OF INVITATION

**PROCUREMENT OF DISPOSABLE ITEMS FOR CATH LAB, STROKE & EP DEPARTMENT THROUGH FRAMEWORK CONTRACT**

Dated; \_\_\_\_\_\_\_\_\_\_\_\_

Dear Sir/ Madam.

1. **Rawalpindi Institute of Cardiology, Rawal Road, Rawalpindi,** invites sealed bids from the eligible bidders **(original manufacturers/their authorized sole agents/ and in case of imported goods their authorized agents/importers/suppliers in Pakistan for supply of Goods)** for supply of Medicine / Drugs in quantities and specifications more specifically described in Section III of the Bidding Documents.
2. Bidding shall be conducted as per the procedure specified in the Bidding Documents.
3. A complete set of original Bidding Documents shall be purchased from the Account office of Rawalpindi Institute of Cardiology, Rawal Road, Rawalpindi with submission of written application on letter head and a copy of CNIC along with payment of non-refundable fee of **Rs.1,000/-** (One thousand only) in all working days during office hours till **10-04-2023,** and the same can be examined online at the PPRA website [**www.ppra.punjab.gov.pk**](http://www.ppra.punjab.gov.pk/) and [**www.ric.gop.pk**](http://www.ric.gop.pk) until the closing date for the submission of bids.

1. Sealed bids are required to be submitted by the interested bidders on **11-04-2023, at 11:00 AM** positively in the Purchase office of Rawalpindi Institute of Cardiology, Rawal Road Rawalpindi. The bids received till the stipulated date & time shall be opened on the same day at **11:30 AM** in the presence of the bidders or their authorized representatives (who choose to attend) by the purchase committee. Bid Security of 2% of the estimated price in the shape of Pay Order/Bank Draft/Deposit at Call/Irrevocable Bank Guarantee from any scheduled bank is required to be furnished with the Financial Bid and copy of bid security attached with Technical Bid otherwise bid will be rejected. **Late bids shall not be entertained.**
2. Single Stage – Two Envelopes bidding procedure shall be applied. The envelopes shall be marked as **“TECHNICAL PROPOSAL**” AND **FINANCIAL PROPOSAL**” in bold and legible letters.
3. All bids (Financial) must be accompanied with a bid security which is **2%** of the Estimated Price in the form of Call Deposit Receipt / Irrevocable Bank Guarantee from any scheduled bank in the name of Executive Director, Rawalpindi Institute of Cardiology, Rawalpindi.
4. In case the date of opening or last date of sale of tender documents is declared as a public holiday by the government or non-working day due to any reason, the next official working day shall be deemed to be the date of sale, submission and opening of tenders accordingly. The time and venue shall remain the same.

**Medical Superintendent**

**Rawalpindi Institute of Cardiology**

**Rawal Road, Rawalpindi**

**051-9281111-20**

# SECTION II

# INSTRUCTION TO BIDDERS (ITB)

## Scope of Bid

Rawalpindi Institute of Cardiology, Rawal Road, Rawalpindi, invites bids for supply of Medicine / Drugs specified in the Section III, Schedule of Requirements & Technical Specifications.

## Source of Funds

The Government of Punjab allocated funds in the specific head of account **(A03927 Purchase of Drugs & Medicine etc)**.

## Eligible Bidders

* + 1. This Invitation for Bids is open to all **original manufacturers/their authorized sole agents/ and in case of imported goods their authorized agents/importers/suppliers in Pakistan for supply of Goods** more specifically described in the Section III, Schedule of Requirements & Technical Specifications.
		2. Government-owned enterprises in Pakistan may participate only if they are legally and financially autonomous and authorized to participate in bidding.
		3. The Agent/Supplier/Importer must possess valid authorization from the Manufacturer and shall have to submit a copy of Memorandum of Association/Partnership deed registered with the Registrar of Companies. However, in case of Manufacturer, they should have a documentary proof as prescribed in the Section V, Bid Form, to the effect that they are the original Manufacturer of the required specifications of Goods.
		4. Bidders under a declaration of ineligibility for corrupt and fraudulent practices issued by any Government (Federal, Provincial or Local) or a public sector organization are NOT ELIGIBLE.

## Corrupt or Fraudulent Practices and Mechanism to Debar/Blacklist the Defaulted Bidder

* + 1. The Punjab Procurement Regulatory Authority, Government of Punjab, defines Corrupt and Fraudulent Practices as “the offering, giving, receiving, or soliciting of anything of value to influence the action of a public official or the contractor in the procurement process or in contract execution to the detriment of the procuring agency; or misrepresentation of facts in order to influence a procurement process or the execution of a contract, collusive practices among bidders (prior to or after bid submission) designed to establish bid prices at artificial, non-competitive levels and to deprive the procuring agency of the benefits of free and open competition and any request for, or solicitation of anything of value by any public official in the course of the exercise of his duty; it may include any of the following practices:
			1. coercive practice by impairing or harming, or threatening to impair or harm, directly or indirectly, any party or the property of the party to influence the actions of a party to achieve a wrongful gain or to cause a wrongful loss to another party;
			2. collusive practice by arrangement between two or more parties to the procurement process or contract execution, designed to achieve with or without the knowledge of the procuring agency to establish prices at artificial, noncompetitive levels for any wrongful gain;
			3. corrupt practice by offering, giving, receiving or soliciting, directly or indirectly, of anything of value to influence the acts of another party for wrongful gain;
			4. fraudulent practice by any act or omission, including a misrepresentation, that knowingly or recklessly misleads, or attempts to mislead, a party to obtain a financial or other benefit or to avoid an obligation;
			5. obstructive practice by harming or threatening to harm, directly or indirectly, persons or their property to influence their participation in a procurement process, or affect the execution of a contract or deliberately destroying, falsifying, altering or concealing of evidence material to the investigation or making false statements before investigators in order to materially impede an investigation into allegations of a corrupt, fraudulent, coercive or collusive practice; or threatening, harassing or intimidating any party to prevent it from disclosing its knowledge of matters relevant to the investigation or from pursuing the investigation, or acts intended to materially impede the exercise of inspection and audit rights;
1. Indulgence in corruption and fraudulent practices is liable to result in rejection of Bids, cancellation of contracts, debarring and blacklisting of the Bidder, for a stated or indefinite period of time.
2. The following are the events which would lead to initiate under Rule 21 of PPRA Rules 2014 Blacklisting / Debarment process;
	1. Submission of false fabricated / forged documents for procurement in tender.
	2. Not attaining required quality of work.
	3. Inordinate tardiness in accomplishment of assigned/agreed responsibilities / contractual obligations resulting loss to procuring agency / Government.
	4. Non execution of work as per terms & condition of contract.
	5. Any unethical or unlawful professional or business behavior detrimental to good conduct and integrity of the public procurement process.
	6. Involvement in any sort of tender fixing.
	7. Persistent and intentional violation of important conditions of contract
	8. Non-adherence to quality specification despite being importunately pointed out.
	9. Security consideration of the State i.e., any action that jeopardizes the security of the State or good repute of the procuring agency.

**PROCEDURE:** A notice will be issued to the agency / individual seeking it/his explanation for the lapses committed by it/him. The explanation will be required within 07 days from the date of issue, (time will be fixed depending upon the intensity of lapses). In case its/his explanation is found unsatisfactory, a show cause notice shall be issued providing an opportunity of being heard followed by decision for blacklistment for a maximum period of three years depending upon the intensity of lapses. The letter for debarring the agency/individual will be published on PPRA website. Once the blacklisting order is issued it shall not be revoked ordinarily unless as provided under Rule-21 of the procurement Rules 2014.

## Eligible Goods and Services

* 1. All goods and related services to be supplied under the contract shall have their origin in eligible source coun­tries, defined in the *Bid Data Sheet (BDS)*, and all expenditures made under the contract will be limited to such goods and related services.
	2. For purposes of this clause, “origin” means the place where the goods are mined, grown, or produced, or the place from which the related services are supplied. Goods are produced when, through manufacturing, processing, or substantial and major assembly of components, a commercially-recognized product is obtained that is substantially different in basic characteristics or in purpose or utility from its components.
	3. The origin of goods and services is distinct from the nationality of the Bidder. In any case, the requirements of rules 10 & 26 of PPR-14, shall be followed.

## Cost of Bidding

* 1. The Bidder shall bear all costs associated with the preparation and submission of its Bid, and the Procuring Agency named in the Bid Data Sheet, hereinafter referred to as “the Procuring Agency,” will in no case be responsible or liable for those costs, regardless of the conduct or outcome of the Bidding process.

## Bidding for Selective Items

* 1. A Bidder, if he so chooses, can bid for selective items from the list of goods provided in the Section III i.e., Schedule of Requirements & Technical Specifications. A Bidder is also at a liberty to bid for all the goods mentioned in the Section III i.e., Schedule of Requirements & Technical Specifications.

**However, Bidders cannot bid for partial quantities of an item mentioned in Section III** i.e., Schedule of Requirements & Technical Specifications. THE BID MUST BE FOR THE WHOLE QUANTITY OF AN ITEM REQUIRED IN THE SECTION III i.e., SCHEDULE OF REQUIREMENTS & TECHNICAL SPECIFICATIONS.

## THE BIDDING PROCEDURE

## The Governing Rules

* 1. The Bidding procedure shall be governed by the Punjab Procurement Rules, 2014, of the Government of the Punjab.

## Applicable Bidding Procedure

* 1. The bidding procedure is governed by Rule 38 “Procedures for Selection of Contractors” sub-rule (2)(a) “Single stage – Two Envelops bidding procedure”. Bidders are advised also to refer to the Bid Data Sheet above to confirm the Bidding procedure applicable in the present bidding process

9.2 The bidding procedure prescribed in the Bid Data Sheet above is explained in the table below.

**Single Stage: Two Envelope Bidding Procedure**

Single stage two envelopes bidding procedure shall be used for procurement of such goods where the bids are to be evaluated on technical and financial grounds and the procedure for single stage two envelopes shall be:

1. the bid shall be a single package consisting of two separate envelopes, containing separately the financial and the technical proposals;
2. the envelopes shall be marked as “Financial Proposal” and “Technical Proposal”;
3. in the first instance, the “Technical Proposal” shall be opened and the envelope marked as “Financial Proposal” shall be retained unopened in the custody of the procuring agency;
4. the procuring agency shall evaluate the technical proposal in the manner prescribed in advance, without reference to the price and shall reject any proposal which does not conform to the specified requirements;
5. during the technical evaluation no amendments in the technical proposal shall be permitted;
6. after the evaluation and approval of the technical proposals, the procuring agency shall open the financial proposals of the technically accepted bids, publically at a time, date and venue announced and communicated to the bidders in advance, within the bid validity period;
7. the financial bids found technically nonresponsive shall be returned un-opened to the respective bidders; and
8. the lowest evaluated bidder shall be awarded the contract;

## THE BIDDING DOCUMENTS

## Contents of the Bidding Documents

* 1. The goods required, applicable bidding procedures, and Contract terms are prescribed in the Bidding Documents. The bidding documents includes;
1. Invitation to bids
2. Instructions to Bidders (ITB) (Section-II)
3. Schedule of Requirements & Technical Specifications (Section-III)
4. Evaluation Criteria (Section-IV)
5. Bid Forms (Section-V)
	1. *Letter of Intention*
	2. *Affidavit*
	3. *Technical Forms*
	4. *Financial Forms*
6. Draft Standard Contract (Section-VI)
	1. *Contract Form*
	2. *General Conditions of the Contract*
	3. *Special Conditions of Contract,*

10.2 The “Invitation for Bids” is not a formal part of the Bidding Documents and is included as a reference only. In case of discrepancies between the Invitation for Bid and the Bidding Documents listed in 10.1 above, the Bidding Documents shall take precedence.

* 1. The Bidder is expected to examine all instructions, forms, terms and specifications in the Bidding Documents. Failure to furnish all information required by the Bidding Documents or to submit a bid not substantially responsive to the Bidding Documents in every respect shall be at the Bidder’s risk and may result in the rejection of its bid.

## Clarification(s) on Bidding Documents

1. A prospective Bidder requiring any clarification of the Bidding documents may notify the Procuring Agency in writing or by email at the Procuring Agency’s address indicated in Invitation to Bid/ Tender Notice/ Advertisement. The Procuring Agency will respond in writing to any request for clarification of the Bidding documents which it receives no later than seven (7) days prior to the deadline for the submission of Bids prescribed in the Bid Data Sheet. Written copies of the Procuring Agency’s response (including an explanation of the query but without identifying) will be sent to all prospective Bidders that have received the Bidding documents.
2. A prospective Bidder requiring any clarification of the Bidding Documents may notify the Procuring Agency in writing or in electronic form that provides record of the content of communication at the Procuring Agency's address indicated in the **BDS.**
3. The Procuring Agency will within three (3) working days after receiving the request for clarification, respond in writing or in electronic form to any request for clarification provided that such request is received not later than seven (7) days prior to the deadline for the submission of Bids**.** However, this clause shall not apply in case of alternate methods of Procurement.
4. Copies of the Procuring Agency's response will be forwarded to all identified Prospective Bidders through an expeditious identified source of communication, e.g. e-mail etc., including a description of the inquiry, but without identifying its source.
5. Should the Procuring Agency deem it necessary to amend the Bidding Documents as a result of a clarification
6. 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 Documents**.**
7. 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 transmitted promptly to all prospective Bidders who have obtained the Bidding Documents. 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. Non-attendance at the pre-Bid meeting will not be a cause for disqualification of a Bidder.

## Amendment(s) to the Bidding Documents

* 1. At any time prior to the deadline for submission of bids, the Procuring Agency, for any reason, whether at its own initiative or in response to a clarification(s) requested by a prospective Bidder, may modify the Bidding Documents by amendment(s).
	2. All prospective Bidders that have received the Bidding Documents shall be notified of the amendment(s) in writing through Post, E-mail or Fax, and shall be binding on them.
	3. In order to allow prospective Bidders reasonable time for taking the amendment(s) into account in preparing their bids, the Procuring Agency, at its discretion, may extend the deadline for the submission of bids.

## PREPARATION OF BIDS

## Language of Bids

* 1. All correspondences, communications, associated with preparation of Bids, clarifications, amendments, submissions shall be written either in English or Urdu or both languages. 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 passages in English or Urdu, in which case, for purposes of interpretation of the Bid, the said translation shall take precedence.

## Documents comprising the Bids

* 1. The Bid shall comprise of the Bid Forms of this Bidding Documents and all those ancillary documentation that are prescribed for the eligibility of the bidders and goods and ancillary services that are found necessary and highlighted in the Bid Forms in Section V.
	2. The Bidder shall complete the Bid Forms and an appropriate Price Schedule (Financial Bid) furnished in the bidding documents, indicating the goods to be supplied, a brief description of the goods, their general and specific characteristics, ancillary services that the bidder is willing or required to provide along with the proposed price.

## Bid Price.

* 1. The Bidder shall indicate on the appropriate form, prescribed in this Bidding Documents, the unit prices and total bid price of the goods, it proposes to supply under the Contract.
	2. Form prescribed for quoting of prices is to be filled in very carefully, preferably typed. Any alteration/correction must be initialed. Every page is to be signed and stamped at the bottom. Tender Enquiry Number of the quoted item may be marked with red/yellow marker.
	3. The Bidder should quote the prices of goods according to the technical specifications as provided in Section III of this document. The technical specifications of goods, different from the required specifications, shall straightway be rejected.
	4. The Bidder is required to offer a competitive price. All prices must **include the taxes and duties**, **where applicable** and all Ex-work & inland transportation & storage charges till the destination (on free delivery to Consignee’s end basis). If there is no mention of taxes, the offered/quoted price shall be considered as inclusive of all prevailing taxes/duties.
	5. The benefit of exemption from or reduction in the taxes and duties shall be passed on to the Procuring Agency.
	6. Prices offered should be for the entire quantity of an item demanded in the Section III i.e., Schedule of Requirement & Technical Specifications; partial quantity offers shall straightaway be rejected. Conditional offer shall also be considered as non-responsive bid.
	7. No request for increase in price due to market fluctuation in the cost of goods and services shall be entertained.

## Bid Currencies.

* 1. Prices shall be quoted in Pak Rupees.

## Samples

* 1. The Bidder shall provide samples of quoted goods along with the bid at his own cost and in a quantity prescribed by the Procuring Agency in Section III.

## Documentation on Eligibility of Bidders

* 1. Bidder shall furnish, as part of its bid (Bid Form) as specified in Section V, documents establishing the Bidder’s eligibility to bid and its qualifications to perform the Contract if its bid is accepted.
	2. The documentary evidence of the Bidder’s eligibility to bid shall establish to the Procuring Agency’s satisfaction that the Bidder, at the time of submission of its bid, is an eligible as defined under ITB Clause 3 above.

## Documentation on Eligibility of Goods

* 1. The Bidder shall furnish, as part of its bid (Bid Form) as specified in Section V, documents establishing the eligibility and conformity to the bidding documents of all goods, which the Bidder proposes to supply under the Contract.

## Bid Security.

* 1. The Bidder shall furnish separately against each quoted item/ Tender Enquiry, as part of its financial bid, a Bid Security of 2% of the estimated price (denominated in Pak Rupees) in the shape of Pay Order/Bank Draft/Deposit at Call/Irrevocable Bank Guarantee from any scheduled bank (as per the format provided in the Bidding Documents) in the name of the Purchaser. Failure to furnish the prescribed Bid Security shall result in the rejection of bid. Bid Security must have a minimum validity period of **One Hundred &Twenty (120) Days** from the last date for submission of the Bids or until furnishing of the Performance Security, whichever is later.
	2. The Bid Security shall be forfeited by the Purchaser, on the occurrence of any/all of the following conditions
		+ 1. If the Bidder withdraws its bid during the period of bid validity specified in the bidding documents; or
			2. If the bidder does not accept the corrections of his Total Bid Price; or
			3. If the Bidder, having been notified for the acceptance of the bid by the Purchaser during the period of the bid validity, fails or refuses to furnish the Performance Security, in accordance with the Bidding Documents.
	3. Unsuccessful bidder’s bid security shall be discharged or returned soon after announcement of the successful bids. The successful Bidder’s bid security shall be discharged upon signing of contract and furnishing the performance guarantee.

## Bid Validity

* 1. Bids shall remain valid for the period identified in the Bid Data Sheet after the date of opening of technical bid prescribed by the Procuring Agency. A bid valid for a shorter period shall be rejected by the Procuring Agency as non-responsive.
	2. A procuring agency shall ordinarily be under an obligation to process and evaluate the bids within the stipulated bid validity period but, under exceptional circumstances and for reasons to be recorded in writing, if an extension is considered necessary, all the bidders shall be requested to extend their respective bid validity period but such extension shall not be for more than the original period of bid validity.
	3. A Bidder who,-
		1. agrees to the extension of the bid validity period shall also extend the validity of the bid bond or security for the extended period of the bid validity;
		2. agrees to the procuring agency’s request for extension of bid validity period shall not be permitted to change the substance of the bid; and
		3. does not agree to an extension of the bid validity period shall be allowed to withdraw the bid without forfeiture of the bid bond or security.

## Format and Signing of Bids.

* 1. The Bidder shall prepare and submit its bid and provide original documents, as appropriate. Copies of any documents must be signed and stamped by the bidder.
	2. The Bid shall be accompanied by the original receipt for payment made for the purchase of the bidding documents. In an event where the Bidder has downloaded the bidding documents from the web, he will require to get the original payment receipt of the prescribed fee from the Procuring Agency well before the date of submission of bid.
	3. The original bid shall be typed or written in indelible ink. All documents should contain proper page marking, attached in sequence as indicated for evaluation in the bidding document and signatures of authorized person. Moreover, signing and stamping of each page of bidding document/form is mandatory.
	4. Any interlineations, erasures, or overwriting shall be valid only if they are initialed by the person or persons signing the bid.

## SUBMISSION OF BIDS

## Sealing and Marking of Bids

* 1. The envelopes shall be marked as “**FINANCIAL PROPOSAL**” and “**TECHNICAL PROPOSAL**” in bold and legible letters to avoid confusion.

Similarly, the Bidder shall seal the proposals/bids in separate envelopes. The envelopes shall then be sealed in an outer envelope.

* 1. The inner and outer envelopes shall:
		1. be addressed to the Procuring Agency at the address given in the Invitation for Bids; and
		2. Bid Reference No. indicated in the Bid Data Sheet, Tender Enquiry No. indicated in Section III, Schedule of Requirements & Technical Specifications and a statement: “DO NOT OPEN BEFORE,” the time and the date specified in the Bid Data Sheet for opening of Bids.
	2. The inner envelopes shall also indicate the name and address of the Bidder to enable the bid to be returned unopened in case it is declared as “non-responsive” or “late”.
	3. If the outer as well as inner envelope is not sealed and marked as required by 23.1 to 23.4 above the Procuring Agency shall assume no responsibility for the bid’s misplacement or premature opening.

## Deadline for Submission of Bids

* 1. **All bids should be submitted in tape binding.** All documents should contain proper page marking. Bids must be submitted by the Bidder and received by the Procuring Agency at the address on the time and date specified in the Bid Data Sheet. Bids received later than **the time and date specified in the Bid Data Sheet will stand summarily rejected.**

24.2 The Procuring Agency may, in its discretion, extend the prescribed deadline for the submission of bids by amending the bidding documents in accordance with ITB Clause 12 above, in which case all rights and obligations of the Procuring Agency and Bidders previously subject to the deadline shall thereafter be subject to the deadline as extended.

## Late Bids

* 1. Any bid received by the Procuring Agency after the deadline for submission of bids prescribed by the Procuring Agency pursuant to ITB Clause 24 shall be rejected and returned unopened to the Bidder.

## Withdrawal of Bids

* 1. The Bidder may withdraw its bid after the bid’s submission and prior to the deadline prescribed for submission of bids.
	2. No bid may be withdrawn in the period between deadline for submission of bids and the expiration of the period of bid validity specified in Bid Data Sheet. Withdrawal of a bid during this period may result in forfeiture of the Bid Security submitted by the Bidder, pursuant to the ITB Clause 20 above.

## OPENING AND EVALUATION OF BIDS

## Opening of Bids by the Procuring Agency

* 1. All bids received, shall be opened by the Procuring Agency publically in the presence of the Bidders or their authorized representatives, who choose to attend the bid opening, on the date, time and venue prescribed in the Bid Data Sheet.
	2. The opening of Bids shall be subject to the Bidding Procedure prescribed in the Bid Data Sheet and elaborated in ITB Clause 9 above.
	3. All Bidders in attendance shall sign an attendance sheet.
	4. The Procuring Agency shall open one Bid at a time and read out aloud its contents which may include name of the Bidder, items bided/quoted for and unit prices and total amount of the Bid (if applicable). The Procuring Agency may choose to announce any other details which it deems appropriate if not in conflict with the Punjab Procurement Rules-2014, specifically Rule 30 (Opening of Bids)
	5. The Procuring Agency shall have the minutes of the Bid opening (technical and when applicable financial) recorded.
	6. No bid shall be rejected at technical proposal/bid opening, except for late bids, which shall be returned unopened to the Bidder
	7. The financial bids found having without Bid Security shall also be returned unannounced to the Bidders. However, prior to return to the Bidder, the Chairman of the Purchase/Procurement Committee shall record a statement giving reasons for return of such bid(s).

## Clarification of Bids

* 1. During evaluation of the bids, the Procuring Agency may, at its discretion, ask the Bidder for a clarification of its bid. The request for clarification and the response shall be in writing, and no change in the prices or substance of the bid shall be sought, offered, or permitted.

## Preliminary Examination

* 1. The Procuring Agency shall examine the bids to determine whether they are complete, whether any computational errors have been made, whether required sureties have been furnished, whether the documents have been properly signed, and whether the bids are generally in order.
	2. In the financial bids the arithmetical errors shall be rectified on the Following basis
1. If there is a discrepancy between the unit price and the total price that is obtained by multiplying the unit price and quantity, the unit price shall prevail, and the total price shall be corrected.
2. If the Bidder does not accept the correction of the errors, its bid shall be rejected, and its Bid Security may be forfeited.
3. If there is a discrepancy between words and figures, the amount in words will be prevail.
	1. Prior to the detailed evaluation, the Procuring Agency shall determine the substantial responsiveness of each bid to the bidding documents. For purposes of this clause, a substantially responsive bid is one, which conforms to all the terms and conditions of the bidding documents without material deviations. Deviations from, or objections or reservations to critical provisions, such as those concerning Applicable Laws, Taxes & Duties and internationally recognized best practices shall be deemed to be a material deviation for technical proposals and Bid Security for financial proposals. The Procuring Agency’s determination of a bid’s responsiveness is to be based on the contents of the bid itself without recourse to extrinsic evidence.
	2. If a bid is not substantially responsive, it shall be rejected by the Procuring Agency and may not subsequently be made responsive by the Bidder by correction of the nonconformity.

## Evaluation of Bids

* 1. The Procuring Agency shall evaluate and compare the bids, which have been determined to be substantially responsive in accordance with ITB Clause 29 above.
	2. All bids shall be evaluated in accordance with the Evaluation Criteria and other terms and conditions set forth in these bidding documents i.e., Rule 32 of PPR 2014.
	3. For the purposes of comparison of bids quoted in different currencies, the price shall be converted into Pak Rupees. The rate of exchange shall be the selling rate, prevailing on the date of opening of bids specified in the bidding documents, as notified by the State Bank of Pakistan/National Bank of Pakistan on that day.
	4. A bid once opened in accordance with the prescribed procedure shall be subject to only those rules, regulations and policies that are in force at the time of issue of notice for invitation of bids.

## Qualification of Bidder

* 1. A procuring agency, at any stage of the procurement proceedings, having credible reasons for, or prima facie evidence of, any defect in the capacity or otherwise of a contractor, whether or not prequalified, may require the contractor to provide such further information concerning the professional, technical, financial, legal or managerial competence as the procuring agency may decide.
	2. Such qualification shall only be laid down after recording reasons thereof in writing. They shall form part of the records of that procurement proceeding.
	3. The Procuring Agency shall determine to its satisfaction whether a Bidder, technically and financially qualified and even having the lowest evaluated responsive bid is qualified to perform the Contract satisfactorily.
	4. The determination can take into account the Bidder’s financial, technical, and production capabilities. It shall 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. Further, during the process of technical evaluation of Bidder, the Procuring Agency may inspect the manufacturing plant/production capacity/warehousing system/practices by a team of experts for assessment, if it deems necessary.
	5. An affirmative determination shall be a prerequisite for award of the Contract to the Bidder. A negative determination shall result in rejection of the Bidder’s bid, in which event the Procuring Agency shall proceed to the next lowest evaluated bid to make a similar determination of that Bidder’s capabilities to perform satisfactorily.
	6. The procuring agency shall disqualify a contractor on the ground that he had provided false, fabricated or materially incorrect information.

## Rejection of Bids

(1) The procuring agency may reject all bids or proposals at any time prior to the acceptance of a bid or proposal.

(2) The procuring agency shall upon request communicate to any bidder, the grounds for its rejection of all bids or proposals, but shall not be required to justify those grounds.

(3) The procuring agency shall incur no liability, solely by virtue of its invoking sub-rule (1) towards the bidders.

(4) The bidders shall be promptly informed about the rejection of the bids, if any.

(5) A procuring agency may, for reasons to be recorded in writing, restart bidding process from any prior stage if it is possible without violating any principle of procurement contained in rule 4 and shall immediately communicate the decision to the bidders.

## Re-Bidding

* 1. If the Procuring Agency rejected all bids in pursuant to ITB Clause 32, it may proceed with the process of fresh bidding but before doing that it shall assess the reasons for rejection and may, if necessary, revise specifications, evaluation criteria or any other condition for bidders.

## Announcement of Evaluation Report

* 1. The Procuring Agency shall announce the results of the bid evaluation in form of a report, not inconsistent with Rule 37 of the Punjab Procurement Rules, 2014, giving justification for acceptance or rejection of bids at least ten days prior to the award of procurement Contract

## Contacting the Procuring Agency

* 1. Subject to ITB Clause 28 above, no Bidder shall contact the Procuring Agency on any matter relating to its bid, from the time of the bid opening to the time of announcement of Evaluation Repot. If a Bidder wishes to bring additional information to the notice of the Procuring Agency, it should do so in writing.
	2. Any effort by a Bidder to influence the Procuring Agency in its decisions on bid evaluation, bid comparison, or Contract award may result in the rejection of the Bidder’s bid. Canvassing by any Bidder at any stage of the bid evaluation is strictly prohibited. Any infringement shall lead to disqualification**.**

## Grievance Redressal

* 1. As per Rule-67 of PPR-14, Procuring Agency shall constitute a Grievance Redressed Committee (GRC) comprising of odd number of persons with proper powers and authorization to address the complaints. The GRC shall not have any of the members of the Procurement Evaluation Committee. The Committee may preferably have one subject specialist depending upon the nature of the procurement in addition to one person with legal background as per their availability to the Procuring Agency.
	2. Any Bidder feeling aggrieved can file its written complaint against the eligibility parameters or any other terms and conditions prescribed in the Bidding documents found contrary to provision of Rule 33 and the same shall be addressed by the GRC well before the proposal submission deadline.
	3. Any party can file its written complaint against the eligibility parameters or any other terms and conditions prescribed in the bidding documents found contrary to provision of Rule 34 and the same shall be addressed by the GRC well before the proposal submission deadline.
	4. Any Bidder feeling aggrieved by any act of the Procuring Agency after the submission of his Bid may lodge a written complaint concerning his grievances not later than ten days after the announcement of the Final evaluation reports. In case of single stage - two envelope bidding procedure any bidder feeling aggrieved from technical evaluation may file a grievance within 5 days of announcement of the technical evaluation report. After completion of the technical evaluation process, the procuring agency shall immediately upload the technical evaluation report on the website of PPRA for obtaining/ receiving grievance petitions from the prospective bidders (if any).
	5. In case, the complaint is filed after the issuance of the final evaluation report, the complainant cannot raise any objection on technical evaluation of the report. Provided that the complainant may raise the objection on any part of the final evaluation report in case where single stage one envelop bidding procedure is adopted.
	6. The GRC shall investigate and decide upon the complaint within fifteen days of the receipt of the complaint. Mere fact of lodging of a complaint shall not warrant suspension of the procurement process.

**AWARD OF CONTRACT**

## Acceptance of Bid and Award Criteria

* 1. The Bidders whose bid is found to be most closely conforming to the Evaluation Criteria prescribed in Section IV and having the lowest evaluated bid, if not in conflict with any other law, rules, regulations or policy of the Punjab Government, shall be awarded the Contract, within the original or extended period of bid validity.

## Procuring Agency’s Right to vary quantities at the time of Award

* 1. The Procuring Agency reserves the right at the time of award of Contract to increase or decrease, the quantity of goods originally specified in Section III i.e., Schedule of Requirements & Technical Specifications without any change in unit price and other terms & conditions.

## Notification of Award

* 1. Prior to the expiration of the period of bid validity, the Procuring Agency shall notify to the successful Bidder in writing that its bid has been accepted.
	2. The notification of award shall constitute the formation of the Contract between the Procuring Agency and the successful Bidder.
	3. The enforcement of the Contract shall be governed by Rule 63 of Punjab Procurement Rules-2014.

## Limitation on Negotiations

* 1. Save and otherwise provided in PPR-2014, Procuring Agency shall not negotiate with any bidder.

## Signing of Contract

* 1. After the completion of the Contract **Negotiations** the Procuring Agency shall send the Bidder the Contract Form provided in the bidding documents, incorporating all agreements between the Parties.
	2. Within **ONE week** of receipt of the Contract Form, the successful Bidder and the Procuring Agency shall sign the Contract in accordance with the legal requirements in vogue.
	3. If the successful Bidder, after completion of all codal formalities shows an inability to sign the Contract then its Bid Security shall stand forfeited and the firm may be blacklisted and de-barred from future participation, whether temporarily or permanently.
	4. The Contract shall become effective upon affixation of signature of the Procuring Agency and the selected Bidder on the Contract document, and shall be governed by the terms and conditions mutually agreed in the contract, bidding documents & relevant laws/rules.
	5. The contract is to be made on stamp paper worth of Rs. @ 25 paisa per every one hundred rupees of the total value of the contract, under section 22(A)(B) of schedule 1 of Stamp Duty Act 1899 read with Finance Act 1995 (Act-VI of 1995) Notification No.JAW/HD/8-21/77 (PG) dated 1st January, 2014.
1. **Performance Guarantee.**
	1. On the date of signing of Contract, the successful Bidder shall furnish a Performance Guarantee, on the Form and in the mannered prescribed by the Procuring Agency.
	2. The Bid Security submitted by the bidder at the time of submitting its bid shall be returned to the Bidder upon submission of Performance Guarantee.
	3. Failure to provide a Performance Guarantee by the Bidder is a sufficient ground for annulment of the award and forfeiture of Bid Security. In such event the Procuring Agency may award the contract to the next lowest evaluated bidder or call for new bid.
2. **Price Reasonability Certificate.**
	1. The supplier shall Certifies on judicial stamp paper that the prices quoted to the Rawalpindi Institute of Cardiology, Rawal Road Rawalpindi, against the items mentioned at Tender Enquiry. No. \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ are not more than the trad prices as per MRP (Maximum Retail Price) Fixed by the Federal Government under Drugs Act, 1976/DRAP Act 2012

All supplies will comply with the provision of Drugs Act, 1976/DRAP Act, 2012 and rules framed there under.

# SECTION III

# SCHEDULE OF REQUIREMENT & TECHNICAL SPECIFICATION

## Schedule of Requirements:

The supplies shall be delivered in accordance with the Purchase Orders as per following schedule of requirements: -

**Respective Consignee’s End: Rawalpindi Institute of Cardiology, Rawal Road, Rawalpindi**

Free Delivery to Consignee’s end (DDP) Basis:

|  |  |
| --- | --- |
| **MODE OF PENALTY** | **DELIVERY OF 100% QUANTITY AS PER PURCHASE ORDER** |
| Without Recovery of Late Delivery Charges | 45 days or earlier(to be determined by the Procuring Agency) |
| With Recovery of Late Delivery Charges @0.067% per day | After 45 (Forty Five) days or earlier (to be determined by the Procuring Agency) and decided by concerned consignee on the formal request of supplier with proper justification. |
| Maximum Rate of Late Delivery Charges | Maximum limit of Late Delivery Charges is 10% after which contract will be cancelled with all legal and codal formalities |
| Risk Purchase | After expiry of prescribed delivery period the Procuring Agency may proceed for risk purchase (at the risk and cost of defaulter) to ensure the un-interrupted healthcare service to the patients |

## ANNUAL DEMAND OF DISPOSABLE ITEMS FOR CATH LAB, STROKE & ELECTROPHYSIOLOGY DEPARTMENT FOR THE FY 2023-24

## (THROUGH FRAMEWORK CONTRACT)

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| **S #** | **DETAIL** | **Unit** | **Qty Req 2023-24** | **Estimated Unit Rate** |
|   | **Coronary**  |   |  |  |
| 1 | Coronary Semi Compliant / compliant balloon with tip entry profile 0.017, 2 mm in diameter and 15mm long (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **2500** |  **10,000**  |
| 2 | Coronary Semi Compliant / compliant balloon with tip entry profile 0.017, 2 mm in diameter and 20mm long (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **500** |  **10,000**  |
| 3 | Coronary Semi Compliant / compliant balloon with tip entry profile 0.017, 2.5 mm in diameter and 15mm long (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **2500** |  **10,000**  |
| 4 | Coronary Semi Compliant / compliant balloon with tip entry profile 0.017, 2.5 mm in diameter and 20mm long (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **500** |  **10,000**  |
| 5 | Coronary Semi complaint / complaint Balloons with tip entry profile 0.017, 1.2mm or less in diameter and 15mm long (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **200** |  **10,000**  |
| 6 | Coronary Semi Compliant / compliant balloon with tip entry profile 0.016, 2 mm in diameter and 15mm long (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **2500** |  **10,000**  |
| 7 | Coronary Semi Compliant / compliant balloon with tip entry profile 0.016, 2 mm in diameter and 20mm long (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **500** |  **10,000**  |
| 8 | Coronary Semi Compliant / compliant balloon with tip entry profile 0.016, 2.5 mm in diameter and 15mm long (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **2500** |  **10,000**  |
| 9 | Coronary Semi Compliant / compliant balloon with tip entry profile 0.016, 2.5 mm in diameter and 20mm long (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **500** |  **10,000**  |
| 10 | Coronary Semi complaint / complaint Balloons with tip entry profile 0.016, 1.2mm or less in diameter and 15mm long (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **200** |  **10,000**  |
| 11 | Coronary Non-Compliant balloons with tip entry profile 0.017 or more, 5-8mm long in diameters from 2.5-4.5mm (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **500** |  **10,000**  |
| 12 | Coronary Non-Compliant balloons with tip entry profile 0.017 or more, 12mm long in diameters from 2.5-4.5mm (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **2000** |  **10,000**  |
| 13 | Coronary Non-Compliant balloons with tip entry profile 0.017 or more, 20mm long in diameters from 2.5-4.5mm (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **500** |  **10,000**  |
| 14 | Coronary Non-Compliant balloons with tip entry profile 0.016 or less, 5-8mm long in diameters from 2.5-4.5mm (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **500** |  **10,000**  |
| 15 | Coronary Non-Compliant balloons with tip entry profile 0.016 or less, 12mm long in diameters from 2.5-4.5mm (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **2000** |  **10,000**  |
| 16 | Coronary Non-Compliant balloons with tip entry profile 0.016 or less, 20mm long in diameters from 2.5-4.5mm (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **500** |  **10,000**  |
| 17 | Coronary Non-Compliant balloons 5mm diameter, 5-8mm long (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **75** |  **10,000**  |
| 18 | Coronary Non-Compliant balloons 5mm diameter, 20mm long (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **150** |  **10,000**  |
| 19 | Coronary Non-Compliant balloons 5mm diameter, 25mm long (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **50** |  **10,000**  |
| 20 | Coronary Non-Compliant balloons 5.5mm diameter, 5-8 mm long (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **30** |  **10,000**  |
| 21 | Coronary Non-Compliant balloons 5.5mm diameter, 20 mm long (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **15** |  **10,000**  |
| 22 | Coronary Non-Compliant balloons 6mm diameter, 5-8 mm long (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **20** |  **10,000**  |
| 23 | Coronary Non-Compliant balloons 6mm diameter, 20 mm long (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **15** |  **10,000**  |
| 24 | Coronary over the wire balloon, 1.2mm diameter and 12 or 15mm long (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **300** |  **13,200**  |
| 25 | Coronary over the wire balloon, 2mm diameter and 12mm long (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **50** |  **13,200**  |
| 26 | Coronary Drug Coated balloon 20mm long in 2, 2.5, 3, 3.5 and 4 mm diameters with randomized control trials and evidence of use in Europe, Japan or USA | No | **500** |  **85,000**  |
| 27 | Coronary Drug Coated balloon 25mm long in 2, 2.5, 3, 3.5 and 4 mm diameters with randomized control trials and evidence of use in Europe, Japan or USA | No | **500** |  **85,000**  |
| 28 | Coronary Drug Coated balloon 35mm long in 2, 2.5, 3, 3.5 and 4 mm diameters with randomized control trials and evidence of use in Europe, Japan or USA | No | **100** |  **85,000**  |
| 29 | Coronary Cutting balloon | No | **75** |  **132,000**  |
| 30 | Coronary covered stent (single stent design) 15 or 16mm long in 2.5, 3, 3.5, 4, 4.5 and 5mm diameters | No | **10** |  **238,000**  |
| 31 | Dilatation Catheter O.P.N.C with high pressure inflation device or equivalent  | No | **20** |  **54,000**  |
| 32 | Inflation Device with accessories including Y-connector with click system  | No | **5000** |  **6,900**  |
| 33 | Mani fold 3 port | No | **10000** |  **800**  |
| 34 | Y-Hemostasis valve set (click system) 6F | No | **1500** |  **2,200**  |
| 35 | Y-Hemostasis valve set (click system) 8F | No | **100** |  **2,200**  |
| 36 | Pressure tube 150cm (for manifold) | No | **10000** |  **180**  |
| 37 | Radial Sheath 4 to 6F | No | **10000** |  **2,000**  |
| 38 | Cylinder radial sheath 6 in 7 | No | **25** |  **2,000**  |
| 39 | M coat Radial sheath 6F | No | **50** |  **6,000**  |
| 40 | Femoral Sheaths 4 to 6F | No | **3000** |  **2,000**  |
| 41 | Femoral Sheaths 7F to 11F | No | **500** |  **2,300**  |
| 42 | Femoral Access Long sheaths 6 and 8F with cannula, usable length 23cm | No | **50** |  **11,000**  |
| 43 | Coronary Aspiration Catheter (evidence of use in USA, Japan or Europe) | No | **3000** |  **33,000**  |
| 44 | FFR Wire | No | **100** |  **15,000**  |
| 45 | IVUS Catheter | No | **100** |  **162,000**  |
| 46 | OFDI Catheter | No | **10** |  **455,000**  |
| 47 | PTCA GUIDE Catheter 6F JR 3.5 with inner diameter 0.071 inch,100cm long (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **300** |  **7,200**  |
| 48 | PTCA GUIDE Catheter 6F JR4 with inner diameter 0.071 inch,100cm long (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **2000** |  **7,200**  |
| 49 | PTCA GUIDE Catheter 6F JL 3.5 with inner diameter 0.071 inch,100cm long (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **100** |  **7,200**  |
| 50 | PTCA GUIDE Catheter 6F JL4 with inner diameter 0.071 inch,100cm long (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **50** |  **7,200**  |
| 51 | PTCA GUIDE Catheter 6F XB/EBU 3 with inner diameter 0.071 inch,100cm long (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **2000** |  **7,200**  |
| 52 | PTCA GUIDE Catheter 6F XB/EBU 3.5 with inner diameter 0.071 inch,100cm long (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **300** |  **7,200**  |
| 53 | PTCA GUIDE Catheter 6F with inner diameter 0.071 inch,100cm long, in shapes other then mentioned above (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **200** |  **7,200**  |
| 54 | PTCA GUIDE Catheter 7F AL 0.75 with inner diameter 0.081 inch or more, 100cm long (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **150** |  **7,200**  |
| 55 | PTCA GUIDE Catheter 7F AL 1 with inner diameter 0.081 inch or more,100cm long (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **50** |  **7,200**  |
| 56 | PTCA GUIDE Catheter 7F XB/EBU 3.5 with inner diameter 0.081 inch or more,100cm long (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **150** |  **7,200**  |
| 57 | PTCA GUIDE Catheter 7F JR4 with inner diameter 0.081 inch or more,100cm long (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **75** |  **7,200**  |
| 58 | PTCA GUIDE Catheter 7F AL 0.75 with inner diameter 0.081 inch or more, 90cm long (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **25** |  **7,200**  |
| 59 | PTCA GUIDE Catheter 7F AL 1 with inner diameter 0.081 inch or more, 90cm long (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **15** |  **7,200**  |
| 60 | PTCA GUIDE Catheter 7F XB/EBU 3.5 with inner diameter 0.081 inch or more, 90cm long (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **25** |  **7,200**  |
| 61 | PTCA GUIDE Catheter 7F JR4 with inner diameter 0.081 inch or more, 90cm long (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **15** |  **7,200**  |
| 62 | PTCA GUIDE Catheter 7 F XBRCA with inner diameter 0.081 inch or more,100cm long (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **20** |  **7,200**  |
| 63 | PTCA GUIDE Catheter 8F AL 0.75 with inner diameter 0.090 inch or more,100cm long (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **50** |  **7,200**  |
| 64 | PTCA GUIDE Catheter 8F AL1 with inner diameter 0.090 inch or more,100cm long (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **20** |  **7,200**  |
| 65 | PTCA GUIDE Catheter 8F XB/EBU 3.5 with inner diameter 0.090 inch or more,100cm long (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **50** |  **7,200**  |
| 66 | PTCA GUIDE Catheter 8F JR4 with inner diameter 0.090 inch or more,100cm long (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **20** |  **7,200**  |
| 67 | PTCA GUIDE Catheter 8F XBRCA with inner diameter 0.090 inch or more,100cm long (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **15** |  **7,200**  |
| 68 | PTCA GUIDE Catheter 8F MP with inner diameter 0.090 inch or more,100cm long (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **50** |  **7,200**  |
| 69 | PTCA GUIDE Catheter 7 & 8 F all shapes other than mentioned above,100cm long (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **50** |  **7,200**  |
| 70 | PTCA GUIDE Catheter 6F JR 3.5 with inner diameter 0.070 inch,100cm long (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **200** |  **7,200**  |
| 71 | PTCA GUIDE Catheter 6F JR4 with inner diameter 0.070 inch,100cm long (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **1000** |  **7,200**  |
| 72 | PTCA GUIDE Catheter 6F XB/EBU 3 with inner diameter 0.070 inch,100cm long (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **1000** |  **7,200**  |
| 73 | PTCA GUIDE Catheter 6F XB/EBU 3.5 with inner diameter 0.070 inch,100cm long (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **200** |  **7,200**  |
| 74 | Diagnostic Catheter 6F JL 3.5 (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **5000** |  **2,400**  |
| 75 | Diagnostic Catheter 6F JL4 (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **100** |  **2,400**  |
| 76 | Diagnostic Catheter 6F JR 3.5 (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **1000** |  **2,400**  |
| 77 | Diagnostic Catheter 6F JR4 (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **5000** |  **2,400**  |
| 78 | Diagnostic Catheter 5F JR 3.5 (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **300** |  **2,400**  |
| 79 | Diagnostic Catheter 5F JR4 (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **100** |  **2,400**  |
| 80 | Diagnostic Catheter 5F JL 3.5 (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **300** |  **2,400**  |
| 81 | Diagnostic Catheter 5F JL4 (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **100** |  **2,400**  |
| 82 | 5F SIM 2 Diagnostic Catheter with soft tip (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **500** |  **5,000**  |
| 83 | Diagnostic Catheter VTK 5F (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **50** |  **5,000**  |
| 84 | Diagnostic Catheter Cobra 2 (C2) 4F (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **25** |  **5,000**  |
| 85 | Diagnostic catheter Radial Tig 5 F | No | **3000** |  **3,300**  |
| 86 | Diagnostic Catheter Vert 125cm (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **100** |  **5,000**  |
| 87 | Diagnostic Catheter All shapes & Sizes other than mentioned above (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **500** |  **2,400**  |
| 88 | Rota Wire | No | **100** |  **30,000**  |
| 89 | Rota Link Plus or equivalent latest rota burr system (All Sizes) with compatibility to RIC console | No | **100** |  **132,000**  |
| 90 | Glide wire angled hydrophilic coating 0.035 x 260cm (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **1500** |  **8,000**  |
| 91 | Glide wire angled hydrophilic coating 0.035 x 150cm (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **1500** |  **4,500**  |
| 92 | Exchange wire normal 260cm (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **8000** |  **1,800**  |
| 93 | 0.014 Coronary Workhorse Wire with floppy tip and tip load 0.7 gram or less (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **5000** |  **9,500**  |
| 94 | 0.014 Coronary Workhorse Wire with inner coil technology or hybrid core technology or hybrid coating technology (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **2000** |  **17,000**  |
| 95 | Polymer jacketed Guide wire pilot 50 or equivalent  | No | **500** |  **12,000**  |
| 96 | Polymer jacketed Guide wire pilot 200 or equivalent  | No | **500** |  **12,000**  |
| 97 | Polymer jacketed Guide wire Fielder XT or equivalent  | No | **500** |  **25,000**  |
| 98 | PTCA Guide wire Sion or equivalent  | No | **25** |  **14,500**  |
| 99 | PTCA Guide wire RG3 or equivalent  | No | **15** |  **27,500**  |
| 100 | Penetrating coronary Guide wire Confianza Pro 12 or progress 200-T or equivalent with tip load 12gm or more | No | **100** |  **23,000**  |
| 101 | PTCA Guide wire Gaia 3 or equivalent  | No | **200** |  **26,500**  |
| 102 | PTCA Guide wire Grand Slam or BHW or equivalent  | No | **25** |  **12,000**  |
| 103 | PTCA Guide wire Sion Black or equivalent  | No | **15** |  **27,500**  |
| 104 | PCI Guide Wire Suoh 3 or equivalent | No | **10** |  **27,500**  |
| 105 | PTCA Guide wire Fielder XTA or equivalent  | No | **10** |  **27,500**  |
| 106 | Tr Band | No | **50** |  **2,000**  |
| 107 | Femoral closure Device Angiosel or equivalent with bioabsorbable anchor and collagen system | No | **100** |  **41,000**  |
| 108 | Femoral percutaneous suture system Proglide or equivalent  | No | **75** |  **58,000**  |
| 109 | Concerto Microcoils for Septal Perforation 0.014 or Equivalent  | No | **20** |  **85,000**  |
| 110 | PTMC Balloon + mullen sheath +Transeptal needle + 0.032 J TIP 150cm guide wire  | No | **250** |  **250,000**  |
| 111 | Angioplasty Drape Set  | No | **10000** |  **3,300**  |
| 112 | EVAR/TEVAR Grafts with accessories (Main body grafts upto 36mm and Limb grafts upto 28mm size) | Case | **5** |  **1,700,000**  |
| 113 | TAVI Valve with loading and delivery system | No | **25** |  **2,700,000**  |
| 114 | Introducer Sheath 14, 16, 18, 20F | No | **25** |  **30,000**  |
| 115 | Lunderquist / Confida / Safari Pre Looped Super/Extra stiff guide wire or equivalent for TAVI procedure | No | **35** |  **38,500**  |
| 116 | Aortic Valvuplasty Balloon (semi-compliant) (evidence of use in Europe, Japan or USA) | No | **15** |  **125,000**  |
| 117 | Aortic Valvuplasty Balloon (non-compliant) (US FDA approved) | No | **30** |  **125,000**  |
| 118 | Vascular Dilators 10, 12, 14F | No | **25** |  **15,000**  |
| 119 | Renal denervation catheter with accessories | No | **10** |  **300,000**  |
|   | **CENTRAL PURCHASE** |  |  |  |
| 120 | Everolimus eluting, US/Japenese FDA approved stent, strut thickness 81 micron or less, in all manufactured sizes of the latest brand being supplied in Pakistan | No | **2000** |  **75,000**  |
| 121 | Sirolimus eluting, US/Japenese FDA approved stent, strut thickness 81 micron or less, in all manufactured sizes of the latest brand being supplied in Pakistan | No | **2000** |  **75,000**  |
| 122 | US/Japenese FDA approved drug eluting stents (DES), other than everolimus or sirolimus drugs, strut thickness 81 micron or less, in all manufactured sizes of the latest brand being supplied in Pakistan | No | **2000** |  **75,000**  |
| 123 | Coronary micro catheter non tapered (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **250** |  **130,000**  |
| 124 | Dual lumen microcatheter (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **20** |  **130,000**  |
| 125 | Coronary micro catheter torqueable tapered septal dilator longest length (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **200** |  **130,000**  |
| 126 | Angled coronary microcatheter (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **15** |  **130,000**  |
| 127 | Guide catheter extention (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **100** |  **170,000**  |
|   | **Peripheral**  |  |  |  |
| 128 | Radifocus Guide Wire M Stiff tip Angled 0.035 x 260 or equivalent (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **100** |  **8,500**  |
| 129 | Peripheral work horse Wire 0.014” (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **200** |  **35,000**  |
| 130 | Peripheral PTCA Balloon 0.035 System OTW, 5mm diameter in 20,40, 80 and 120mm length (longest shaft length) (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **60** |  **25,000**  |
| 131 | Peripheral PTCA Balloon 0.035 System OTW, 6mm diameter in 20, 40 and 80mm length (longest shaft length) (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **40** |  **25,000**  |
| 132 | Peripheral PTCA high pressure Balloon 0.035 System OTW, 8mm diameter in 20, 40 and 80mm length with RBP of 24atm or more (longest shaft length) (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **40** |  **45,000**  |
| 133 | Peripheral PTCA high pressure Balloon 0.035 System OTW, 10mm diameter in 20, 40 and 80mm length with RBP of 24atm or more (longest shaft length) (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **30** |  **45,000**  |
| 134 | Peripheral PTCA high pressure Balloon 0.035 System OTW, 12mm diameter in 40 and 80mm length with RBP of 24atm or more (longest shaft length) (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **20** |  **45,000**  |
| 135 | Peripheral PTCA high pressure Balloon 0.035 System OTW, 14mm diameter in 40 and 80mm length with RBP of 24atm or more (longest shaft length) (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **15** |  **45,000**  |
| 136 | Peripheral PTCA Balloon 0.014 System OTW, 3mm diameter in 60 and 150mm length (longest shaft length) (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **20** |  **35,000**  |
| 137 | Peripheral PTCA Balloon 0.014 System OTW, 2.5mm diameter in 60 and 150 mm length (longest shaft length) (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **40** |  **35,000**  |
| 138 | Peripheral PTCA Balloon 0.014 System OTW, 2mm diameter in 80 and 220mm length (longest shaft length) (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **40** |  **35,000**  |
| 139 | Peripheral PTCA Balloon 0.014 System OTW, 1.5mm diameter in 40mm length (longest shaft length) (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **20** |  **35,000**  |
| 140 | Peripheral Balloon premounted Stents OTW 0.035, 6mm diameter in 27, 37 and 57mm lengths (longest shaft length) (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **10** |  **100,000**  |
| 141 | Peripheral Balloon premounted Stents OTW 0.035, 8mm diameter in 27, 37 and 57mm lengths (longest shaft length) (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **30** |  **100,000**  |
| 142 | Peripheral Balloon premounted Stents OTW 0.035, 10mm diameter in 25 or 27, 37 and 57mm lengths (longest shaft length) (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **15** |  **100,000**  |
| 143 | Renal Stent 0.014, 6mm diameter in 12mm length (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **4** |  **100,000**  |
| 144 | Renal Stent 0.014, 6mm diameter in 14 or 15mm and 18 or 19mm lengths (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **4** |  **100,000**  |
| 145 | Renal Stent 0.014, 7mm diameter in 12mm length (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **4** |  **100,000**  |
| 146 | Renal Stent 0.014, 7mm diameter in 15mm and 18 or 19mm lengths (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **4** |  **100,000**  |
| 147 | Self Expending Stent for Limbs longest shaft length OTW 0.035, 6mm diameter in 40, 80 and 120mm lengths (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **30** |  **100,000**  |
| 148 | Self Expending Stent for Limbs longest shaft length OTW 0.035, 8mm diameter in 40 and 60mm lengths (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **30** |  **100,000**  |
| 149 | Peripheral Graft Stent 6mm diameter in 27 and 37mm lengths (longest shaft length) (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **6** |  **300,000**  |
| 150 | Peripheral Graft Stent 8mm diameter in 27 and 37mm lengths (longest shaft length) (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA)  | No | **6** |  **300,000**  |
| 151 | Peripheral Graft Stent 10mm diameter in 37mm lengths (longest shaft length) (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **4** |  **300,000**  |
| 152 | Peripheral infusion catheter longest Length or equivalent (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **10** |  **75,000**  |
| 153 | Peripheral Suppport Catheter 0.014 (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **50** |  **30,000**  |
| 154 | Peripheral Suppport Catheter 0.035 (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **50** |  **30,000**  |
| 155 | Peripheral Micro Catheter inner lumen 0.027" (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **10** |  **110,000**  |
| 156 | Uterine fibrord embolization particles (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **20** |  **25,000**  |
| 157 | Pushable coils 0.018" (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **20** |  **30,000**  |
| 158 | Torque device 0.035" | No | **150** |  **2,500**  |
| 159 | 4fr Micro Puncture kit (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **50** |  **30,000**  |
| 160 | Guiding Sheaths straight tip, 6F in 90cm length with option of connecting Y connector (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **150** |  **60,000**  |
| 161 | Guiding Sheaths straight tip, 6F in 55 or 65cm length with option of connecting Y connector (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **50** |  **60,000**  |
| 162 | Angiojet Catheter (Coronary/Arterial/Venous/Pulmonary) | No | **8** |  **450,000**  |
| 163 | Jet stream Catheter | No | **2** |  **450,000**  |
| 164 | Reterivable IVC Filter with legs (non Cage design) (Europeon CE Mark/Japenese/US FDA Approved, with evidence of use in Europe, Japan or USA) | No | **5** |  **150,000**  |
| 165 | IVC Filter Retriver Kit (Europeon CE Mark/Japenese/US FDA Approved, with evidence of use in Europe, Japan or USA) | No | **5** |  **150,000**  |
| 166 | Femoral Sheath 16F 45CM | No | **3** |  **45,000**  |
| 167 | Femoral Sheath 9F 55 or 65CM | No | **3** |  **45,000**  |
| 168 | Peripheral Vascular Plugs | No | **10** |  **180,000**  |
| 169 | Peripheral Guide Catheter MP1, 6F in 55or 65 cm length (Europeon CE Mark/Japanese/US FDA with evidence of use in Europe, Japan or USA) | No | **50** |  **9,500**  |
|   | **Stroke / Neuro Intervention** |  |  |  |
| 170 | Sofia Plus Neuro Aspiration Catheter or equivalent with 0.070" inner lumen, longest length (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **300** |  **250,000**  |
| 171 | React Neuro Aspiration Catheter or equivalent with 0.071" inner lumen, longest length (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **100** |  **230,000**  |
| 172 | Sofia Distal access / Support Catheter 5F or equivalent, maximum outer diameter 0.068, Longest length (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **50** |  **330,000**  |
| 173 | Stent Retriever Parametric Desgin for proximal and distal treatment with compatible microcatheter for delivery (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **300** |  **350,000**  |
| 174 | Stent Retriever with drop zone technology with compatible microcatheter for delivery | No | **50** |  **350,000**  |
| 175 | Stent Retriever with interlinked cage technology with compatible microcatheter for delivery | No | **50** |  **350,000**  |
| 176 | Neuro Micro Catheter, Longest Length, with 0.021 inch inner diameter compatible with stent retrievers (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **200** |  **50,000**  |
| 177 | Neuro intervention workhorse wire 0.014 (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **500** |  **50,000**  |
| 178 | Self-Expanding Carotid Stent with Open Cell Design, tapered stents, 6x8x40mm and 7x10x40mm (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **40** |  **100,000**  |
| 179 | Self-Expanding Carotid Stent with closed cell design (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **60** |  **125,000**  |
| 180 | Carotid Filter covering vessel diameters including 7mm (wire mounted) (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **40** |  **175,000**  |
| 181 | Carotid Filter covering vessel diameters including 7mm (any wire filter) (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **60** |  **125,000**  |
| 182 | Carotid proximal protection device Moma or Equivalent (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **25** |  **130,000**  |
| 183 | Dedicated Neurovascular Long Sheath 6F, straight tip, 90cm long (Japenese/US FDA Approved, with evidence of use in Japan/USA or valid US FDA CFG) | No | **100** |  **70,000**  |
| 184 | Balloon Tip Guide Catheter  | No | **100** |  **130,000**  |
|   | **Paeds Cardiology** |   |  |  |
| 185 | ASD Device with delivery system With clamp | No | **40** |  **324,000**  |
| 186 | ASD Device with delivery system with Screw | No | **40** |  **250,000**  |
| 187 | PDA Device with delivery system Long Shank | No | **15** |  **230,000**  |
| 188 | PDA Device with delivery system reverse Shank | No | **40** |  **220,000**  |
| 189 | PDA Device with delivery system Normal Shank | No | **40** |  **220,000**  |
| 190 | Diagnostic Catheter for Paeds  All sizes | No | **500** |  **2,500**  |
| 191 | Giude Wire - 014x180 | No | **10** |  **1,500**  |
| 192 | Giude Wire - 018x260 | No | **10** |  **2,500**  |
| 193 | Giude Wire - 035x150 | No | **50** |  **1,500**  |
| 194 | Giude Wire - 035x260 | No | **100** |  **2,500**  |
| 195 | Gliude Wire - 018x260 | No | **15** |  **2,500**  |
| 196 | Gliude Wire - 021x260 | No | **10** |  **2,500**  |
| 197 | Gliude Wire - 035x150 | No | **30** |  **1,500**  |
| 198 | Gliude Wire - 035x260 | No | **50** |  **2,500**  |
| 199 | Super Stiff or Extra Stiff | No | **300** |  **8,500**  |
| 200 | Velvoplasty Balloon low profile (All Size) | No | **50** |  **80,000**  |
| 201 | VSD Device Muscular Diferent Size  | No | **10** |  **250,000**  |
| 202 | Bib Balloon | No | **20** |  **222,000**  |
| 203 | Septostomy Balloon | No | **20** |  **95,000**  |
| 204 | Balloon Mounted Covered Stent Diferent Size  | No | **20** |  **510,000**  |
| 205 | CP Stent Covered With Delivery system Diferent Size  | No | **20** |  **650,000**  |
| 206 | Aneurysmal VSD Device or Equivalent With Delivery system | No | **10** |  **290,000**  |
| 207 | Infantile VSD Device or Equivalent With Delivery system | No | **40** |  **290,000**  |
| 208 | Wedge Pressure Catheter or Equivalent | No | **5** |  **55,000**  |
| 209 | Vascular Plugs | No | **10** |  **85,000**  |
| 210 | Single loop snare all sizes | No | **25** |  **95,000**  |
| 211 | Snare Catheter (3loop) all sizes | No | **25** |  **100,000**  |
| 212 | Pulmonary Valve With Delivery system | No | **10** |  **5,000,000**  |
|   | **Electrophysiology** |   |  |  |
| 213 | T.P.M Lead 5 & 6F |   | No | **1000** |  **9,000**  |
| 214 | Single Chamber Permanent Pacemaker  | Full Body MRI Safe, 3T Latest (or equivalent) ESSENTIO VR EL MRI Pacing System | No | **300** |  **419,000**  |
| 215 | Dual Chamber Permanent Pacemaker | MRI Conditional, 1.5T | No | **50** |  **585,000**  |
| 216 | Dual Chamber Permanent Pacemaker | Full body MRI safe, 3T – Latest model, not older than 5 years since FDA approval | No | **150** |  **585,000**  |
| 217 | ICD Single Chamber or VIGILANT EL ICD (or equivalent) minimum longevity >12 years | Complete set | No | **50** |  **1,800,000**  |
| 218 | ICD Double or VIGILANT EL ICD (or equivalent) | Complete set | No | **6** |  **2,200,000**  |
| 219 | CRT-P VISIONIST X4 CRT-P (or equivalent) complete set minimum longevity > 10 years | Complete set | No | **80** |  **1,850,000**  |
| 220 | CRT-D VIGILANT X4 CRT-D (or equivalent) complete set | Complete set | No | **2** |  **3,800,000**  |
| 221 | Subcutaneous ICD  | Complete set | No | **4** |  **1,450,000**  |
| 222 | His bundle pacing with dual IPG complete set | Complete set | No | **50** |  **450,000**  |
| 223 | His bundle pacing ventricular (his) lead  | with delivery system | No | **40** |  **600,000**  |
| 224 | Quadripolar LV lead for CRT  | with delivery system | No | **10** |  **450,000**  |
| 225 | Coronary sinus venogram balloon  |   | No | **20** |  **100,000**  |
| 226 | Implantable Loop Recorders  | Linq or DX or equivalent | No | **20** |  **450,000**  |
| 227 | Cryoballoon set or Equivalent  | compatible with cryoconsole | No | **50** |  **900,000**  |
| 228 | Cryoballoon only  | compatible with cryoconsole | No | **40** |  **500,000**  |
| 229 | Flexcath Steerable long sheath for Cryoballoon delivery (or equivalent) | compatible with cryoconsole | No | **40** |  **120,000**  |
| 230 | Achieve (multipolar) mapping catheter (or equivalent)  | compatible with cryoconsole | No | **20** |  **250,000**  |
| 231 | Cryofreeze Focal Ablation Catheter  | compatible with cryoconsole | No | **10** |  **450,000**  |
| 232 | Disposable Drape set for Pacing  | with instrument as per hospital requirement | No | **300** |  **18,000**  |
| 233 | Blazer Prime Temperature Ablation Catheter or equivalent |   | No | **50** |  **306,526**  |
| 234 | Blazer II HTD Temperature Ablation Catheter, 4mm tip Standard curve (or equivalent) |   | No | **40** |  **268,210**  |
| 235 | Blazer II Temperature Ablation Catheter  | Standard curve, standard distal (or equivalent) | No | **20** |  **268,210**  |
| 236 | Blazer II Temperature Ablation Catheter  | Standard, Asymmetric Curve (or equivalent) | No | **10** |  **268,210**  |
| 237 | Blazer II Temperature Ablation Catheter  | Large Curve (or equivalent) | No | **40** |  **268,210**  |
| 238 | Blazer II Temperature Ablation Catheter  | Large Curve, Medium Distal (or equivalent) | No | **30** |  **268,210**  |
| 239 | Blazer II Temperature Ablation Catheter )  | Standard Curve, Medium Distal (or equivalent) | No | **40** |  **268,210**  |
| 240 | Blazer II Temperature Ablation Catheter  | Standard Curve, Extended Distal (or equivalent) | No | **40** |  **268,210**  |
| 241 | Cable, Thermistor For Blazer Ablation Catheters |   | No | **8** |  **229,895**  |
| 242 | Cable, Recorder-APM (POD to Junction Box IEGM cable) |   | No | **8** |  **84,295**  |
| 243 | Blazer Open-Irrigated Ablation Catheter | Large curve (or equivalent) | No | **50** |  **919,578**  |
| 244 | Cable, Blazer OI to Maestro POD |   | No | **5** |  **383,157**  |
| 245 | Blazer DX-20 7F Bidirectional | Duodecapolar Diagnostic Catheter (or equivalent) | No | **10** |  **651,368**  |
| 246 | Cable, Blazer DX  | Duodecapolar Catheter (or equivalent) | No | **8** |  **265,705**  |
| 247 | Orbiter ST 7F | Steerable Diagnostic Catheter, Duodecapolar (or equivalent) | No | **10** |  **175,000**  |
| 248 | Protected 24-pin Orbiter STTM cable 120cm |   | No | **5** |  **190,000**  |
| 249 | Polaris X™ 2.5,5,2.5mm (or equivalent) | Steerable Decapolar Catheter | No | **20** |  **306,526**  |
| 250 | Polaris X™ 2,10,2mm (or equivalent) | Steerable Decapolar Catheters | No | **10** |  **306,526**  |
| 251 | 10-pole male Quick Connect to 10 pins 2mm 5 ft. (or equivalent) |   | No | **5** |  **160,926**  |
| 252 | Dyn XT, 6F, 110cm,10E - 2,5,2mm (or equivalent) | Steerable Catheter, Large Curve | No | **80** |  **199,242**  |
| 253 | Dyn XT 6F | Steerable Catheter (Quadripolar, Hexapolar, Octapolar)Standard Curve  | No | **40** |  **199,242**  |
| 254 | Quadripolar steerable diagnostic EP catheter (or equivalent) | Dynamic Tip  | No | **20** |  **252,884**  |
| 255 | Woven Fixed Curve Catheter, 6F 5E,10,10,267mm,120cm 120cm (or equivalent) |  Cournand Curve  | No | **50** |  **199,242**  |
| 256 | Woven Fixed Curve Diagnostic Catheter 6F 4E,2,5,2mm,125cm (or equivalent) | Josephson Curve  | No | **100** |  **199,242**  |
| 257 | Viking Fixed Curve 6F Diagnostic Catheter (or equivalent) | Hisser Curve  | No | **30** |  **91,957**  |
| 258 | Viking Fixed Curve Diagnostic Catheter6F 4E,2,5,2mm,115cm (or equivalent) | Josephson Curve  | No | **50** |  **91,957**  |
| 259 | Viking Fixed Curve Diagnostic Catheter 6F 4E,2,5,2mm,115cm (or equivalent) | Cournand Curve  | No | **50** |  **91,957**  |
| 260 | Viking Fixed Curve Diagnostic Catheter 6F , 4E, 10,10,10mm, 115cm (or equivalent) | Cournand Curve  | No | **80** |  **91,957**  |
| 261 | Viking Fixed Curve Diagnostic Catheter5F 4E,2,5,2mm,115cm (or equivalent) | Josephson Curve  | No | **15** |  **91,663**  |
| 262 | Viking Fixed Curve Diagnostic Catheter5F 4E,2,5,2mm, 115cm (or equivalent) |  Hisser Curve  | No | **15** |  **91,957**  |
| 263 | Viking Fixed Curve Diagnostic Catheter5F 10E,2,5,2mm, 115cm (or equivalent) | CS | No | **15** |  **153,263**  |
| 264 | Cable 125cms (or equivalent) | Protected 4- Pin, | No | **6** |  **84,294**  |
| 265 | Cable125cm (or equivalent) | Easy-Mate 4- Pin, | No | **6** |  **84,294**  |
| 266 | Cable-232125cm (or equivalent) | Protected 6-Pin Easy-Mate, | No | **6** |  **99,842**  |
| 267 | Cable 120cm (or equivalent) | 10-Pin Surelink, | No | **5** |  **114,947**  |
| 268 | Cable 120cm (or equivalent) |  Protected 4-Pin Surelink, | No | **10** |  **68,968**  |
| 269 | Cable 120cm Red, Blue,Yellow (or equivalent) | Protected 4-Pin Surelink, | No | **10** |  **68,968**  |
| 270 | Zurpaz/Direx Sheath (Steerable) (or equivalent) | Steerable Sheath, SymmetricMedium Curve  | No | **50** |  **421,473**  |
| 271 | TSXFS Fixed Sheath 8.5F, 60cm, All Curves (or equivalent) |   | No | **140** |  **168,589**  |
| 272 | TSX Transseptal Needle 18GA, 71cm, Lrg Crv (or equivalent)  |   | No | **140** |  **162,509**  |
| 273 | TSXFS Fixed Sheath 8.5F, 79.4cm, 55Crv (or equivalent) |   | No | **3** |  **210,736**  |
| 274 | TSX Transseptal Needle 18GA, 89cm, Lrg Crv (or equivalent) |   | No | **3** |  **200,000**  |
| 275 | Safesept needle for transeptal puncture (or equivalent) |   | No | **10** |  **95,000**  |
| 276 | Valleylab Ground Pad Pack 1x50 (or equivalent) |   | No | **15** |  **90,000**  |
| 277 | Needle eye snare (for percutaneous lead extraction) |   | No | **10** |  **250,000**  |
| 278 | Open-Irrigated Tubing Kit MetriQ |   | No | **50** |  **54,473**  |
| 279 | mechanical lead extraction sheath  | Tightrail, siterail (or equivalent) | No | **10** |  **350,000**  |
| 280 | 3D Navigational, Thermocool Smart touch Unidirectional catheter (or equivalent)  | F curve | No | **80** |  **841,000**  |
| 281 | 3D Navigational, Thermocool Smart touch bi-directional catheter (or equivalent)  | D/F, F curve | No | **30** |  **915,700**  |
| 282 | 3D Navigational Multipolar catheter Pentaray (or equivalent) |   | No | **70** |  **600,000**  |
| 283 | Smartable System irrigation tubing set (or equivalent) |   | No | **80** |  **29,700**  |
| 284 | 3D Navigational, Thermocool Navstar Catheter (or equivalent) |   | No | **25** |  **820,000**  |
| 285 | Cable for thermocool  | Smart touch catheter | No | **30** |  **200,000**  |
| 286 | Cable for thermocool  | Navstar catheter | No | **10** |  **200,000**  |
| 287 | Cable for multipolar  | Pentaray catheter | No | **40** |  **200,000**  |
| 288 | 3D mapping external reference patch |   | No | **50** |  **110,000**  |
| 289 | Ground patch  | Compatible with 3D mapping ablation PIU | No | **50** |  **25,000**  |
| 290 | Defibrillator patches  | Compatible with hospital defibrillator | No | **300** |  **45,000**  |
| 291 | Intracardiac echo (ICE) catheter (e.g. accunav) or equivalent  | compatible with ICE console (GE) | No | **10** |  **600,000**  |
| 292 | Percutaneous Left Atrial Appendage Occlusion Device (e.g. watchman) or equivalent |   | No | **10** |  **1,500,000**  |
| 293 | Bioptome for Myocardial Biopsy |   | No | **10** |  **400,000**  |
| 294 | Vizigo steerable long sheath (or equivalent) | Medium curve, Compatible with 3D mapping system | No | **30** |  **493,000**  |
| 295 | Decanav Steerable Catheter (or equivalent) | Compatible with 3D mapping system | No | **40** |  **445,000**  |

**Note:-**

1. The bidder shall provide samples of the quoted items (sample containing 3 packs of each quoted item along with its bid).
2. Certificate regarding fulfillments of requirements under Bio safety Act. 2005 and the rules framed there under must be attached for Vaccines/Sera, Biotechnical products etc.
3. For thermolabile drugs for which storage temperature is 2-8 degree centigrade. The firm shall be bound to produce batch wise cold chain data from the source of origin & thermo log data from factory to Consignee’s end.
4. Any further information can be obtained from the Purchase Department of Rawalpindi Institute of Cardiology, Rawalpindi
5. ***All taxes are applicable as per government rules.***
6. ***Mentioned GST (where applicable) item wise separately in financial bid otherwise it will be consider that rates are inclusive of GST.***
7. Hospital can reduce the quantity according to the budget.
8. All supply will be packed as per Government packing rules.
9. Company will pay the DTL fee (where applicable).
10. Quoted rates will be applicable on Model Pharmacy, Zakat, Pakistan Bait ul Mal, CM Grant, PM Health Program, Panel Patients, etc.
11. The supplier firm (s) shall be bound by the clause of the contract which state that the near expiry / unused disposables should be timely replace by supplier firm upon requested of the procuring agency.

# SECTION IV

# EVALUATION CRITERIA

## EVALUATION CRITERIA

## COMPULSORY PARAMETERS:

Failure to comply with any compulsory parameter will result in disqualification of bidder.

|  |  |  |
| --- | --- | --- |
| **PARAMETERS** | **DOCUMENTS REQUIRED** | **COMPLIANCE STATUS****(Yes/No)** |
| Tender Purchase Receipt, CNIC, Bid Security | * Original Tender Purchase Receipt, Copy of CNIC, Copy of bid security.
 |  |
| Drug Manufacturing / Sale License | * Valid Drug Manufacturing License issued by DRAP for manufacturers.
* Valid Drug Sale License for importers
* In case of expired / in valid, valid renewal application must be attached
 |  |
| Certificates | * NTN, GST, Valid Professional Tax
 |  |
| Free sale certificate (Where Applicable) | * Embassy Attested Free Sale Certificate of the Product
* Documents Pakistan embassy attested be attached and duly notarized / verified from country of origin.
 |  |
| Authorized Agency Agreement (for Sole Agents / Distributors) | * Affidavit from the sole agent / Distributor that their product(s) are freely available with same brand name in the country of manufacturer & is safe for human consumption
 |  |
| Drug Registration Certificate (DRC) | * Valid Drug Registration Certificate issued by DRAP
* Valid product enlistment certificate issued by DRAP (where applicable)
 |  |
| Product Quality Certificate | * Valid quality certificate of quoted product
 |  |
| Good Manufacturing Practices (GMP) Certificate | * Valid Good Manufacturing Practices (GMP) certificate issued by the Drug Regulatory Authority Pakistan (DRAP)
* In case of imported product, valid GMP certificate issued by the regulatory authority of manufacturer’s country will be considered.
 |  |
| Income Tax Return | * Clearly showing annual sale/turnover of the bidder for the year 2022-23.
 |  |
| Undertaking Regarding Non Cancellation / Suspension of Drug Registration of quoted product of the bidder by Drug Regulatory Authority of Pakistan within last two years | * Undertaking on judicial paper
 |  |
| Undertaking Regarding Non Declaration of Spurious / Adulterated batch by DTLs of Punjab / any Competent Lab of quoted item within last two years | * Undertaking on judicial paper
 |  |
| Specification quoted in the technical offer will be verified from samples provided with the bid product that comply 100% with the advertised specification and fulfill the requirement as per labeling and packing Rules 1986 shall be considered for evaluation | * Samples of quoted items.
 |  |

## ORDINARY PARAMETERS:

Failure to comply with any compulsory parameter will result in disqualification of bidder.

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| **S. #** | **Parameters** | **Detail** | **Total Marks**  | **Remarks** |
| 1 | Past Performance of the Bidder (Last two years)  | Major institutions (Government / Semi Government) Served:

|  |  |  |
| --- | --- | --- |
|  | 1 | 02 |
|  | 2 to 3 | 04 |
|  | 4 to 5 | 06 |
|  | 6 to 7 | 08 |
|  | 8 & above | 10 |

 | **10** | The claim requires documentation (Purchase orders, Receipts / Completion Certificates and Delivery Challans etc) of the institution(s) |
| 2 | Market Experience of quoted product |

|  |  |  |
| --- | --- | --- |
|  | Market Availability of quoted items in leading chain stores & pharmacies for last 02 years | 05 |
|  | 2-4 years | 7 |
|  | 5-6 years | 10 |

 | **15** | * Market availability in leading Chain Stores & pharmaceuticals / institution of quoted item will be calculated from the date of commercial invoice.
* The firm will attach purchase orders of the quoted item of any Government/ Semi-Government Institution / Private Institution registered with Income Tax Department.
 |
| 3 | Credibility & Certification of Manufacturer |

|  |  |  |
| --- | --- | --- |
|  | WHO / US FDA / FDA-Japan / CE certification/ WHO prequalification / prequalification by Provincial or Federal institutes (July 2018 to onward) | 07 |
|  | Valid ISO Certifications | 03 |

 | **10** | Valid copies of certificates / letters required |
| 4 | Financial Status of Bidders |

|  |  |  |
| --- | --- | --- |
|  | Last Year Audited Accounts | 05 |
|  | Tax Return last three year  | 05 |

 | **10** | Bank Statements are also required for last year bidder can provide more than one Bank certificate. |
| 5 | Past performance of quoted product for last three years |

|  |  |  |
| --- | --- | --- |
|  | No Complaint during last three year of the quoted item from any statutory lab. | 05 |
|  | No Complaint during last two year of the quoted item from any statutory lab. | 03 |

 | **05** | The firm will provide undertaking in this regard. The purchaser reserves the rights to verify the claim. |
| 6 | Institution Experience Rawalpindi Institute of Cardiology, Rawalpindi. |

|  |  |  |
| --- | --- | --- |
|  | Bidder who supplied items as order within stipulated delivery period. | 15 |
|  | The bidder who supplied the ordered items after due date/expiration of stipulated delivery period.  | 10 |
|  | The bidder who could not supply items within stipulated time / after due date/expiration of stipulated delivery period.  | 0 |

 | **15** | Delivery challan with receiving of the stores is required; the provided documents should be verifiable, not applicable for new bidders. |

* **Total Marks of Ordinary Parameters: 65**
* **Qualifying marks in Ordinary parameters: 65% (42.25 and above)**

**NOTE:**

The institute will procure DRAP registered medical devices only through open tender under PPRA rules but in case of any item, if no DRAP registered item is being offered then for smooth functioning of hospital, to deliver the uninterrupted services to deserving cardiac patients and for the sake of stop gap arrangements in order to save the precious human lives / public exchequer, the procuring agency will only accept NON DRAP registered medical devices after getting approval from DRAP. Moreover, if that turndown, item will not be accepted by the procuring agency.

# SECTION V

# BID FORM

## BID COVER SHEET

Bid Ref. No \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ Date: \_\_\_\_\_\_\_\_\_\_\_\_\_

Name of the supplier / Firm Contractor: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

Address: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

E-mail: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

Phone: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

Facsimile: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

Bid Security.

Bid Security attached with financial bid YES NO

Bid for

Selected items for Schedule of Requirements

|  |  |  |  |  |
| --- | --- | --- | --- | --- |
| **Item / Tender Enquiry Number** | **Name of the Item** | **Batch Capacity of the Drug / Medicine / Product** | **Trade Price** | **MRP (Maximum Retail Price)** |
|  |  |  |  |  |
|  |  |  |  |  |
|  |  |  |  |  |
|  |  |  |  |  |
|  |  |  |  |  |
|  |  |  |  |  |

Signed: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

Date: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

Official Stamp: \_\_\_\_\_\_\_\_\_\_\_\_

Attachment: Original Receipts for the purchase of the bidding documents.

## BID FORM 1

**LETTER OF INTENTION**

*Bid Ref No. RIC/PO/\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_*

*Date of the Opening of Bids \_\_\_\_\_\_\_\_\_\_\_\_\_*

*Name of the Contract :{ Supply of Drugs and Medicines etc}*

To: Rawalpindi Institute of Cardiology, Rawalpindi

Dear Sir/Madam,

Having examined the bidding documents including Addenda Nos *[insert* ***numbers & Date of individual Addendum]***, the receipt of which is hereby acknowledged, we, the undersigned, offer to supply and deliver the Goods under the above-named Contract in full conformity with the said bidding documents and at the rates/unit prices described in the price schedule or such other sums as may be determined in accordance with the terms and conditions of the Contract. The above amounts are in accordance with the Price Schedules attached herewith and are made part of this bid.

We undertake, if our bid is accepted, to deliver the Goods in accordance with the delivery schedule specified in the schedule of requirements.

If our bid is accepted, we undertake to provide a performance security/guaranty in the form, in the amounts, and within the times specified in the bidding documents.

We agree to abide by this bid, for the Bid Validity Period specified in the Bid Data Sheet and it shall remain binding upon us and may be accepted by you at any time before the expiration of that period.

Until the formal final Contract is prepared and executed between us, this bid, together with your written acceptance of the bid and your notification of award, shall constitute a binding Contract between us.

We understand that you are not bound to accept the lowest or any bid you may receive.

We undertake that, in competing for (and, if the award is made to us, in executing) the above contract, we will strictly observe the laws against fraud and corruption in force in Pakistan.

We confirm that we comply with the eligibility requirements as per ITB clauses 18 &19 of the bidding documents.

Dated thi*s [insert: number****]*** day of *[insert: month****]***, *[insert: year].*

Signed: In the capacity of *[insert:* ***title or position]***

Duly authorized to sign this bid for and on behalf of *[insert :****name of Bidder]***

## BID FORM 2

**AFFIDAVIT**

I/We, the undersigned solemnly state that:

1. I/We have read the contents of the Bidding Documents and have fully understood it.
2. The Bid being submitted by the undersigned complies with the requirements enunciated in the bidding documents.
3. The Goods that we propose to supply under this contract are eligible goods within the meaning of Clause 18 of the ITB.
4. The undersigned are also eligible Bidders within the meaning of Clause 19 of the ITB.
5. The undersigned are solvent and competent to undertake the subject contract under the Laws of Pakistan.
6. The undersigned have not paid nor have agreed to pay, any Commissions or Gratuities to any official or agent related to this bid or award or contract.
7. The undersigned are not blacklisted or facing debarment from any Government, or its organization or project.
8. That the prices offered are not more than trade price.
9. I/We, further undertake that the prices given are reasonable and not given more than in any Government/Autonomous/District Government institutions during the current financial year. If any difference detected, the firm is bound to refund the difference in price.

I/We affirm that the contents of this affidavit are correct to the best of our knowledge and belief.

Signed: In the capacity of *[insert: T****itle or position]***

Duly authorized to sign this bid/affidavit for and on behalf of *[insert: N****ame of Bidder]***

## BID FORM 3(A)

Name of the Firm \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

Bid Reference No: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

Date of opening of Bid. \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

Documentary Evidence: Eligibility of the Bidders and Goods

|  |  |  |  |
| --- | --- | --- | --- |
| **Required****Documentation****(*To Be Filled by the******Procuring Agency*)** | **Checklist1*****(To be initialed******by the Bidder******against each******document)*** | **Relevant Page****Number2 in the****Bid *(To be filled******by the Bidder)*** | **Supporting Documents3*****(To be filled by the Bidder******with name of the******documents that are******submitted to meet the******requirement)*** |
| **Column: 1** | **Column: 2** | **Column: 3** | **Column: 4** |
| Valid Drug ManufacturingLicense |  |  |  |
| Valid Drug Registrationcertificate of quoteditems |  |  |  |
| Valid Drugs SaleLicense for Importer4 |  |  |  |
| Valid Good ManufacturingPractices Certificate |  |  |  |
| Valid Import License(where applicable) |  |  |  |
| Letter of Manufacturer'sauthorization |  |  |  |
| Partnership Deed(where applicable) |  |  |  |
| NTN Certificate |  |  |  |
| GST Certificate |  |  |  |
| Letter of Intention |  |  |  |
| **Affidavit** |  |  |  |
| Past Performanceevidence |  |  |  |
| Child Labor FreeCertificate5 |  |  |  |
| Original Receipt ofpurchase of Bidding |  |  |  |

1. Bidders should only initial against those requirements that they are attaching with the form 3(a). In case they do not have any document to attach the corresponding cell in column 2 should be left blank.
2. Bidders are required to mention the exact page number of relevant document placed in the Bid.
3. Bidders are advised to attach all Supporting documents with this form in the order of the requirement as mentioned in column 1.
4. In case of Sole Agent
5. Bidders are required to furnish a certificate to the effect that their firm is free from child labor and having standard child labor free policy

## BID FORM 3(B)

**MANUFACTURER’S AUTHORIZATION**

**To:** ***[Name & Address of the Procuring Agency]***

WHEREAS *[name of the Manufacturer]* who are established and reputable Manufacturers of

*[name and/or description of the goods]* having factories at *[address of factory]* do hereby authorize *[name and address of Supplier/ Agent]* to submit a bid, and subsequently negotiate and sign the Contract with you against the Invitation for Bids (IFB) No. *[Reference of the Invitation to Bid] for* the goods manufactured by us.

We hereby extend our full guarantee and warranty as per Clause 14 of the General Conditions of Contract for the goods offered for supply by the above firm against this Invitation for Bids.

Signature: -------------------------------------.

Designation: -------------------------------------

Official Stamp: ----------------------------------

1. This letter of authority should be on the letterhead of the Manufacturer and should be signed by a person competent and having the power of attorney to bind the Manufacturer. It should be included by the Bidder in its bid.

## BID FORM 4

**Firm’s Past Performance**

Name of the Firm:

Bid Reference No:

Date of opening of Bid: \_\_\_\_\_\_\_\_\_\_\_\_\_\_

Assessment Period: (as per Evaluation Criteria)

|  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- |
| **Name of the****Purchaser/Institution** | **Purchase****Order No.** | **Description****Of Order** | **Value of****Order** | **Date of****Completion** | **Purchaser’s8****Certificate** |
|  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |

1. Bidders may use additional Sheets if required.
2. All certificates are to be attached with this form.

## BID FORM 5(A)

**Price Schedule**

**(User Note):** *This form is to be filled by the Bidder for each individual quoted item and shall submit with Financial Proposal.*

**Name of Firm: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**

**Bid Reference No. \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**

**Tender Inquiry No. \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**

**Date of Opening of Bid. \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**

|  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- |
| **S. #** | **Name of Item** | **Unit Price****(inclusive all****applicable taxes****+ transportation****charges)** | **No. of****Units** | **Total****Price** | **Discounts*****(if any)*** | **Final Total Price****(Inclusive of all taxes)** |
| 1 | 2 | 3 | 4 | 5 (3\*4) | 6 | 7 (5-6) |
|  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |
|  | **TOTAL** |  |  |  |

1. **FINAL TOTAL PRICE: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**
2. **DISCOUNT: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**
3. **FINAL QUOTED PRICE: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**

**(C=A-B)**

**Signature: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**

**Designation: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**

**Date: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**

**Official Stamp: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**

9 If a Bidder does not wish to offer an item wise discount but intends to offer an overall discount to its quoted price that should be mentioned here.

## BID FORM 5(B)

**Price Schedule**

(Price Analysis)

**(User Note):** *This form is to be filled by the Bidder for each individual quoted item and shall submit with Financial Proposal*

**Name of Firm: \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**

**Bid Reference No. \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**

**Tender Inquiry No. \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**

**Date of Opening of Bid. \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**

|  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- |
| **Item / Tender****Enquiry****No.** | **Name****of the****Item** | **Unit Price** | **Total****Price/Unit** | **No. of****Units** | **Total****Price** |
|
| **Ex-factory, Ex****Ware house,****Ex-Show****Room, Off the****Shelf** | **Sales****and****Income****Tax** | **Other Levies****and Duties *(if******any)*** | **Packagin****g** | **Transportatio****n Costs****incidental to****delivery** | **Other****Incidental****Costs as****defined in the****Schedule of****Requirement** |  |  |  |
|
|
|
|
|
| A | b | c | D | E | F | Gg=a+b+c+d+e+f | h | ii = g\*h |
|
|
|
|  |  |  |  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |  |  |  |

Signature: ---------------------------------------------------

Designation: ------------------------------------------------

Date: ----------------------------------------------------------

Official Stamp: ---------------------------------------------

## BID FORM 6

**Performance Guarantee**

To: *Executive Director*

 *Rawalpindi Institute of Cardiology*

 *Rawal Road Rawalpindi.*

Whereas *[Name of Supplier]* (hereinafter called “the Supplier”) has undertaken, in pursuance of

Contract No. *[Number]* dated *[date]* to supply *[description of goods]* (hereinafter called “the Contract”).

And whereas it has been stipulated by you in the said Contract that the Supplier shall furnish you with a Bank Guarantee by a scheduled bank for the **sum of 5% of the total** Contract amount as a Security for compliance with the Supplier’s performance obligations in accordance with the Contract.

And whereas we have agreed to give the Supplier a Guarantee:

Therefore we hereby affirm that we are Guarantors and responsible to you, on behalf of the Supplier, 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 Supplier 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\_\_\_\_\_\_\_\_\_\_\_\_ day of\_\_\_\_\_\_\_\_\_, 201\_\_\_

Signature and Seal of the Guarantors/ Bank

Address

Date

# SECTION VI

# DRAFT STANDARD CONTRACT

## Contract Form

AGREEMENT

**THIS CONTRACT** is made at on day of 201\_\_,

between the **(Rawalpindi Institute of Cardiology, Rawalpindi)**, (hereinafter referred to as the “Purchaser”) of the First Part; and M/s *(firm name)* a firm registered under the laws of Pakistan and having its registered office at *(address of the firm)* (hereinafter called the “Supplier”) of the Second Part (hereinafter referred to individually as “Party” and collectively as the “Parties”).

**WHEREAS** the Purchaser invited bids for procurement of goods, in pursuance whereof M/s*(firm name)* being the Manufacturer/ authorized sole agent /Supplier of (item name) in Pakistan and ancillary services offered to supply the required item (s); and

Whereas, the Purchaser has accepted the bid by the Supplier as per following detail;

|  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- |
|  |  |  |  |  |  |  |
|  |  |  | **Unit** |  |  |  |
| **Tender** |  | **Approved** | **Price in** |  | **Total Cost** |  |
| **Enquiry/** | **Item Name** | **PKR** | **Quantity** |  |
| **Specifications** | **(PKR)** |  |
| **Item No.** |  | **(As per** |  |  |
|  |  |  |  |  |
|  |  |  | **contract)** |  |  |  |
|  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |
| **NOW THE PARTIES TO THIS CONTRACT AGREE TO THE FOLLOWING;** |  |  |

1. **The Contract:** The following documents shall be deemed to form and be read and construed as integral part of this Contract , viz:-

|  |  |  |
| --- | --- | --- |
| **a.** | This Contract Form |  |
| **b.** | The Schedule of Requirements | **Annex- A** |
| **c.** | Special Conditions of Contract & the Technical Specifications | **Annex- B** |
| **d.** | Original Price Schedule along with unsolicited discount offered by the firm (if any) |
|  | Submitted by the Bidder. | **Annex- C** |
| **e.** The Purchaser’s Notification of Award(AAT) | **Annex- D** |
| **f.** | Purchase Order | **Annex-E** |
| **g.** | Payment Schedule | **Annex-F** |
| **h.** | The General Conditions of Contract | **Annex-G** |
| **i.** | Performance Guarantee/Security | **Annex-H** |

1. Manufacturer’s certificate of warranty under Drugs Act 1976/DRAP Act 2012 &

rules framed there under **Annex-I**

**k.** The bidding document of Procuring Agency **Annex-J**

1. **Interpretation:** In this Contract words and expressions shall have thesame meanings as are respectively assigned to them in the General Conditions of this

Contract hereinafter referred to as “Contract”:

1. **The Term of the Contract:** This contract shall remain valid for one year from the dateof signing, unless amended by mutual consent.
2. The Supplier declares as under:
	1. *[Name of the 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 Punjab or any administrative subdivision or agency thereof or any other entity owned or controlled by it (Government of Punjab) through any corrupt business practice.
	2. Without limiting the generality of the foregoing, [the Supplier] represents and warrants that it has fully declared the brokerage, commission, fees 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 consultation fee or otherwise, with the object of obtaining or including the procurement of a Contract, right interest, privilege or other obligation or benefit in whatsoever form from Government of Punjab, except that which has been expressly declared pursuant hereto.
	3. *[The Supplier]* certifies that has made and shall make full disclosure of all agreements and arrangements with all persons in respect of or related to the transaction with Government of Punjab and has not taken any action or shall not take any action to circumvent the above declaration, representation or warranty.
	4. *[The Supplier]* accepts full responsibility and strict liability for making any false declaration, not making full disclosure, misrepresenting facts 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 Procuring Agency under any law, Contract or other instrument, be void able at the option of Procuring Agency.
	5. Notwithstanding any rights and remedies exercised by Procuring Agency in this regard, *[The Supplier]* agrees to indemnify Procuring Agency for any loss or damage incurred by it on account of its corrupt business practices and further pay compensation to Procuring Agency in an amount equivalent to ten time the sum of any commission, gratification, bribe, finder’s fee or kickback given by *[The 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 Procuring Agency.
	6. In case of any dispute concerning the interpretation and/or application of this Contract shall be settled through arbitration. The \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

(**Name of Authority to be inserted here**) or his nominee shall act as sole arbitrator.

The decisions taken and/or award made by the sole arbitrator shall be final and binding on the Parties.

1. **Items to be Supplied & Agreed Unit Cost:**
	1. The Supplier shall provide to the Purchaser the items on the agreed cost more specifically described in the Price Schedule Submitted by the Bidder (Annex C).
	2. Each Items supplied shall strictly conform to the Schedule of Requirements (Annex A) and to the Technical Specification (Annex B) prescribed by the Purchaser against each item
	3. The Unit Cost agreed in the Price Schedule (Annex C), is inclusive of all taxation and costs associated with transportation and other agreed incidental costs.
2. **Payments:**The Purchaser hereby covenants to pay the Supplier in consideration ofthe provision of the Goods and Services, as specified in the Schedule of Requirements and Technical Specification in accordance with the Price Schedule submitted by the Supplier, the amount against the delivered items or such other sum as may become payable under the provisions of this Contract at the time and in the manner prescribed by this Contract.
3. **Mode of Payment:**All payments to the Supplier shall be made through CrossedCheques issued in the name of [supplier’s name]
4. **Payment Schedule**: All payments to the Supplier shall be made in accordance with theagreed Payment Schedule at Annex: F, upon satisfactory completion of delivery and fulfillment of documentary and codal formalities highlighted in the Payment Schedule at Annex F.
5. **Performance Guarantee/Security:**
	1. The Supplier, within 07 days of signing of this contract, shall provide to the Purchaser a Performance Security in the form of an Irrevocable Bank Guarantee equivalent to**5%**of the total Contract amount having validity of one year from its date of issuance from any scheduled bank on the prescribed format and in prescribed manner. This Performance Guarantee/Security shall be released to the Supplier upon successful completion of the Contract.
	2. Supplier’s Bid Security already submitted with the Bid shall only be released upon satisfactory submission of a Performance Guarantee/Security in accordance with sub-clause (i) above.
	3. Failure to submit a Performance Guarantee/Security shall result into forfeiture of Bid Security and Cancellation of Contract. Failure to furnish the required Performance Guarantee/Security shall constitute a breach of the contract and the procuring agency shall be entitled to make other arrangement at risk and expenses of firm without any notice.

1. **Penalties/ Liquidated Damages**
	1. Wherein the Supplier fails to make deliveries as per signed contract & purchase order and within the stipulated time frame specified in the Schedule of Requirement, the Contract to the extent of non-delivered portion of supplies shall stand cancelled.
	2. After the cancellation of the Contract no supplies shall be accepted and the amount of Performance Guaranty/Security to the extent of non–delivered portion of supplies shall be forfeited.
	3. If the Supplier fails to supply the whole consignment and not able to deliver to consignee’s end, the entire amount of Performance Guaranty/Security shall be forfeited to the Government account and the firm shall be blacklisted minimum for two years for future participation.
	4. The exact time frame for making supplies with and without penalty shall be indicated in subsequent purchase order.
	5. In case of late delivery of goods beyond the periods specified in the Schedule of Requirements and subsequent purchase order, **a penalty @ 0.067% per day of thecost of late delivered supply shall be imposed upon the Supplier.**
2. **Notices:**All notices and correspondences incidental to this contract shall be inEnglish language and shall be addressed to:

***Rawalpindi Institute of Cardiology, Rawalpindi***

**For the Supplier:**



IN WITNESS Whereof the Parties hereto have caused this Contract to be executed at\_\_\_\_\_\_\_\_\_\_\_\_\_(the place) and shall enter into force on the day, month and year first above mentioned.

**Sealed & singed on behalf of Sealed / Sealed for the Manufacturer**

**Procuring Agency Authorized Supplier / Authorized Agent**

 Signature of Owner of Firm---------------------

 Name ------------------------------------------------

Medical Superintendent Father Name----------------------------------------

Rawalpindi Institute of Cardiology Designation-----------------------------------------

Rawalpindi CNIC#------------------------------------------

**Witnesses (Procuring Agency): Witnesses:**

Signature\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ Signature\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

CNIC#\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ CNIC#\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

Name \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ Name \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

Designation\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_ Designation\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

## ANNEX: A

### SCHEDULE OF REQUIREMENTS:

The supplies shall be delivered in accordance with the Purchase Orders as per following schedule of requirements:-

**Respective Consignee’s End: Rawalpindi Institute of Cardiology, Rawal Road, Rawalpindi**

Free Delivery to Consignee’s end (DDP) Basis:

|  |  |
| --- | --- |
| **MODE OF PENALTY** | **DELIVERY OF 100% QUANTITY AS PER PURCHASE ORDER** |
| Without Recovery of Late Delivery Charges | 45 days or earlier(to be determined by the Procuring Agency) |
| With Recovery of Late Delivery Charges @0.067% per day | After 45 (Forty Five) days or earlier (to be determined by the Procuring Agency) and decided by concerned consignee on the formal request of supplier with proper justification. |
| Maximum Rate of Late Delivery Charges | Maximum limit of Late Delivery Charges is 10% after which contract will be cancelled with all legal and codal formalities |
| Risk Purchase | After expiry of prescribed delivery period the Procuring Agency may proceed for risk purchase (at the risk and cost of defaulter) to ensure the un-interrupted healthcare service to the patients |

## ANNEX. B

### Special Conditions of the Contract & Technical Specifications

**a).** **Product Specifications.**

*(Detailed technical specifications, given in Section III, will be followed)*

**b).** **Labeling and Packing**

* + 1. The manufacturer shall follow the Drugs (Labelling and Packing) Rules 1986, framed under the Drugs Act, 1976.
		2. However, the name of Drug / Medicine (Generic & Brand), equally prominent, should be printed/ written in indelible ink both in English and Urdu on the outer cartons and on each Pack, Bottle, Strip/ Blister, Tubes etc. Besides the name and principal place of business of the Manufacturer, the drug manufacturing license no., manufacturing date, expiry date, registration No., batch No., retail price, and Urdu version namely: name of drug, dosage and instructions, should also be written on the outer carton and on the most inner container in bold letters. All tablets shall be supplied in strip / blister pack (one side aluminum and other side PVC/PVD). Expiry date must be printed on each strip / blister. The syrup should be supplied in glass / pet bottle with sealed caps.
1. **Additional instructions for packing**
	1. The suppliers are required to furnish the Warranty certificate with regard to the potency and stability (Including coloration of medicines) of the Drug for human consumption etc. in accordance with the Drugs Act, 1976/DRAP Act 2012 & rules framed the reunder on judicial paper.
	2. The bidder shall supply the Drugs/Medicines/Items in special green packing with Logo of the Government of Punjab. The following wording/insignia shall be printed in bold letters both in Urdu & English in indelible red color ink on each carton, pack, bottle, strip / blister, tubes, vial / ampoule etc. In combo Packs the sterilized water for injection / solvent shall bear the wording/insignia on the vial/ampoules etc. In case of items supplied by the foreign manufacturer the mentioned condition may be relaxed by the Procuring Agency.

**(Name of Procuring agency)**

**“PUNJAB GOVERNMENT PROPERTY”**

**“NOT FOR SALE”**

1. After signing of the Contract, the Supplier shall submit the samples of finished medicines in accordance with the above instructions for approval of the **(Name & Address ofProcuring Agency is to be inserted here).** The approved samples will be shared with the

Consignee/End User and all subsequent supplies must be in accordance with the approved samples.

**d).** **Shelf life**

1. The shelf life must be **at least 50% or 45% with penalty**
2. The lower limit of the shelf life must be up to **45% with imposition of 1%penalty** charges of actual shortfall in shelf life below prescribed limit.
3. **Supplier firm shall be bound that the near expiry / unused disposable could have to be timely replaced by the supplier firm upon request of the procuring agency.**

**e).** **Testing/Verification Procedures**

* + - 1. After delivery of drugs and medicines at the Purchaser’s premises, the Purchaser shall send the samples from all batches of each consignment of the supplied store to the Drugs Testing Laboratory, Punjab, for testing. The Inspection Committee constituted by the Purchaser shall inspect the quantity, specifications of goods after receipt of standard quality report of each batch of supplied store issued by DTL concerned under Drugs Act 1976/DRAP Act 2012 & rules framed there under. **The cost of the lab tests** shall be borne by the Supplier.
	1. In case of **substandard/failure** report of any batch, the Supplier has the right to go for appellate laboratory. If it is again declared substandard, the Supplier will be intimated and they will be bound to re-supply the **entire fresh stock** of that batch **free of cost** within the reasonable time period to be intimated by the purchaser but not later than **21days (three weeks**) from the date of intimation, which will be subject to completion of all testing and verification formalities. At the parallel, the case will also be forwarded to the concerned authority for **legal action** as per Drugs Act 1976 and **disposal of substandard stocks**.
	2. The Inspection Committee will carry out detailed physical examination of stocks and can reject, even if it is declared of standard quality by DTL, if found not according to the approved sample and other technical specifications like packaging, labeling, printing and quantity etc. Moreover, the Supplier will also be responsible to replace the unconsumed expired stores without any further charges.
1. **Transportation/Delivery Requirements**
	* 1. The Supplier shall arrange such transportation of the drugs and medicines as is required to prevent their damage or deterioration during transit to their final destination and in accordance with the terms and manner prescribed in the Schedule of Requirement.
		2. All costs associated with the transportation including loading/unloading of drugs and medicines and road taxes shall be borne by the Supplier.
2. All **cold chain (perishable**) items must be delivered in a safe and proper manner, prescribed for such types of items.

## ANNEX. C

### PRICE SCHEDULE SUBMITTED BY THE BIDDER

*(The approved price schedule submitted by the Bidder will be attached)*

## ANNEX. D

### PURCHASER’S NOTIFICATION OF AWARD

*(****Advance Acceptance of Tender issued by the Procuring Agency will be attached****)*

## ANNEX. E

### PURCHASE ORDER

*(Specimen Sample of PO)*



No.\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

Dated \_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_

|  |  |  |  |  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- | --- |
|  |  |  |  |  |  |  |  |  |  |  |  |
| **1** | **Purchase Order No** |  |  |  |  |  |  |  |  |
| **Date** |  |  |  |  |  |  |  |  |
| **2** | **Supplier/Firm Name** |  |  |  |  |  |  |  |  |
| **3** | **Supplier/Firm’s Address** |  |  |  |  |  |  |  |
| **4** | **Firm Contact No** |  |  |  |  |  |  |  |  |
| **5** | **Conditions of the Contract:** |  | As already communicated in the BiddingDocument & Signed Contract |  |  |
|  |  |  |
| **6** | **Particulars of Stores:** |  |  | As per detail given below |  |  |
|  |  |  |  |  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |  |  |  |  |
| **Item****No.** | **Item Name** | **Approved****Specifications** | **Unit****Price in****PKR****(As per****contract)** | **Quantity** | **Total Cost (PKR)** |  |  |
|  |  |
|  |  |
|  |  |
|  |  |
|  |  |
|  |  |
|  |  |  |  |  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |  |  |  |  |
|  |  |  |  |  |  |  |  |  |  |  |  |

Additional instructions (if any):

1.

2.

3.

**\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_**

## ANNEX. F

### PAYMENT SCHEDULE

1. *100% Payment to the Suppliers will be made*
	1. *against satisfactory performance and upon submission of required documents and in accordance with the procedure mentioned in Rule 64 and other relevant rules of PPR-2014.*
	2. *On production of Inspection Certificate and receipt certificate from Consignee, after recovery of Government dues (if any) including Professional Tax.*
2. *Part Supply and Part Payment is allowed, but the Payment will only be made after the receipt of next installment within due time.*

*(However, if there is any alternate payment schedule, agreed by the Procuring Agency and Supplier, it will be annexed*

## ANNEX. G

### GENERAL CONDITIONS OF CONTRACT (GCC)

|  |  |  |  |  |  |  |  |
| --- | --- | --- | --- | --- | --- | --- | --- |
| **1.** | **Definitions** | 1.1 | In this Contract, the | following | terms shall be | interpreted | as |
|  |  |  | indicated: |  |  |  |  |  |  |  |  |
|  |  |  | (a) “The | Contract” | means | the | agreement | entered | into |
|  |  |  |  | between the Purchaser (\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_\_) |
|  |  |  |  | and the Supplier, as recorded in the Agreement signed by |
|  |  |  |  | the Parties, including all attachments and appendices |
|  |  |  |  | thereto and all documents incorporated by reference |
|  |  |  |  | therein. |  |  |  |  |  |  |  |
|  |  |  | (b) “The | Contract | Price” | means | the price payable to the |
|  |  |  |  | Supplier under the Contract for the full and proper |
|  |  |  |  | performance of its Contractual obligations. |  |  |  |
|  |  |  | (c) | “The Goods” means all those supplies which the Supplier |
|  |  |  |  | is required to supply to the Purchaser under the Contract. |
|  |  |  | (d) “The | Services” | means those | services ancillary to | the |
|  |  |  |  | supply of above goods, such as printing of special |
|  |  |  |  | instructions on the label and packing, design and logo of |
|  |  |  |  | the Government of Punjab, transportation of goods upto |
|  |  |  |  | the desired destinations and other such obligations of the |
|  |  |  |  | Supplier covered under the Contract. |  |  |  |
|  |  |  | (e) “GCC” means | the | General | Conditions | of | Contract |
|  |  |  |  | contained in this section. |  |  |  |  |  |
|  |  |  | (f) | “SCC” means Special Conditions of the Contract. |  |  |
|  |  |  | (g)  | “The Purchaser” means The Rawalpindi Institute of Cardiology Rawalpindi |  |
|  |  |  |  |  |  |  |
|  |  |  | (h) “The Supplier” means the individual or firm supplying the |
|  |  |  |  | goods under this Contract. |  |  |  |  |  |
|  |  |  | (i) | “Day” means calendar day. |  |  |  |  |
| **2.** | **Application** | 2.1 | These General Conditions shall apply to the extent that they |
|  |  |  | are not superseded by provisions of other parts of the Contract. |

|  |  |  |
| --- | --- | --- |
| **3. Source of** | 3.1 | All goods and related services to be supplied under the |
|  | **Import** |  | contract that are required to be imported in Pakistan shall have |
|  |  |  | their origin in eligible source countries as prescribed by the |
|  |  |  | commercial policies of the Federal Government of Pakistan and |
|  |  |  | all expenditures made under the contract shall be limited to |
|  |  |  | such goods and services. |
|  |  | 3.2 | For purposes of this clause, “origin” means the place where the |
|  |  |  | goods are produced, or the place from which the related |
|  |  |  | services are supplied. Goods are produced when, through |
|  |  |  | manufacturing or processing. |
| **4.** | **Standards** | 4.1 | The goods supplied under this Contract shall conform to the |
|  |  |  | standards mentioned in the Technical Specifications. |
|  |  | 4.2 | In consideration of the payments to be made by the Purchaser |
|  |  |  | to the Supplier as hereinafter mentioned, the Supplier hereby |
|  |  |  | covenants with the Purchaser to provide the Goods and |
|  |  |  | Services and to remedy defects therein in conformity in all |
|  |  |  | respects with the provisions of this Contract. |
|  |  | 4.3 | If the Supplier provide substandard item and fail to provide the |
|  |  |  | fresh supply, the payment of risk purchase (which will be |
|  |  |  | purchased by the (**RIC Rawalpindi)** the price difference shall be paid by the supplier. |
|  |  | 4.4 | In case of supply of substandard product the cost associated |
|  |  |  | with disposal/destruction or associated handling shall be borne |
|  |  |  | by the Supplier i.e., removal from purchaser’s premises, |
|  |  |  | burning, dumping, or incineration. |
| **5.** | **Use of Contract** | 5.1 | The Supplier shall not, without the Purchaser’s prior written |
|  | **Documents and** |  | consent, disclose the Contract, or any provision thereof, or any |
|  | **Information.** |  | specification, plan, drawing, pattern, sample, or information |
|  |  |  | furnished by or on behalf of the Purchaser in connection |
|  |  |  | therewith, to any person other than a person employed by the |
|  |  |  | Supplier in the performance of the Contract. Disclosure to any |
|  |  |  | such employed person shall be made in confidence and shall |

extend only so far as may be necessary for purposes of such performance.

1. The Supplier shall not, without the Purchaser’s prior written consent, make use of any document or information enumerated in GCC Clause 5.1 except for purposes of performing the contract.

1. Any document, other than the Contract itself, enumerated in GCC Clause 5.1 shall remain the property of the Purchaser and

shall be returned (all copies) to the Purchaser on completion of the Supplier’s performance under the Contract if so required by the Purchaser.

|  |  |  |  |
| --- | --- | --- | --- |
|  |  |  5.4 | The Supplier shall permit the Purchaser to inspect the |
|  |  |  | Supplier’s accounts and records relating to the performance of |
|  |  |  | the Supplier. |
| **6.** | **Patent Rights** | 6.1 | The Supplier shall indemnify the Purchaser against all third- |
|  |  |  | party claims of infringement of patent, trademark, or industrial |
|  |  |  | design rights arising from use of the Goods or any part thereof |
|  |  |  | in the country. |
| **7. Submission of** | 7.1 | Before commencing supplies, the Supplier shall provide |
|  | **Samples** |  | samples free of cost, if and as specified in the Schedule of |
|  |  |  | Requirements of the product to the designated office or staff, |
|  |  |  | as the case may be. |
| **8. Ensuring** | 8.1 | To ensure storage arrangements for the intended supplies, the |
|  | **storage** |  | Supplier shall inform the Purchaser at least **0ne (01) week** in |
|  | **arrangements** |  | advance. However, in case no space is available at the |
|  |  |  | Purchaser’s premises at the time of supply, the Purchaser |
|  |  |  | shall, at least **02 days** prior to such situation, shall inform the |
|  |  |  | Supplier, in writing, of the possible time frame of availability of |
|  |  |  | space by which the supplies can be made. In case the Supplier |
|  |  |  | abides by the given time frame it shall not be penalized for |
|  |  |  | delay. |
| **9.** | **Inspections and** | 9.1 | The Purchaser or its representative shall have the right to |
|  | **Tests** |  | inspect and / or to test the goods in accordance with the |
|  |  |  | procedure given in the SCC to confirm their conformity to the |
|  |  |  | Contract specifications at no extra cost to the Purchaser. |
|  |  | 9.2 | All costs associated with testing shall be borne by the Supplier. |

1. The Purchaser’s right to inspect, test and, where necessary, reject the goods after the goods either at Supplier’s premises or upon arrival at Purchaser’s destinations shall in no way be limited or waived by reason of the goods having previously been inspected, tested, and passed by the Purchaser or its representative prior to the goods delivery from the point of Supply or manufacturing.

|  |  |  |
| --- | --- | --- |
|  |  | Nothing in GCC Clause 9 shall in any way release the Supplier |
|  |  | from any warranty or other obligations under this Contract. |
| **10. Delivery and** | 10.1 | The Supplier in accordance with the terms and manner |
| **Documents** |  | specified in the Schedule of Requirements shall make delivery |
|  |  | of the goods. |
|  | 10.2 |  |
|  |  | The Supplier shall furnish all necessary documentation |
|  |  | necessary for completion of the delivery, at the time of delivery |
|  |  | and in the manner prescribed. |
|  | 10.3 |  |
|  |  | The goods supplied under the Contract shall be delivered on |
|  |  | free delivery of consignee’s end basis under which risk is |
|  |  | transferred to the buyer after the Goods having been delivered; |
| **11. Insurance** | 11.1 | The supplier shall be solely responsible for Insurance of the |
|  |  | Goods subject to the contract. |
| **12. Transportation** | 12.1 | The Supplier shall arrange such transportation of the goods as |
|  |  | is required to prevent their damage or deterioration during |
|  |  | transit to their final destination and in accordance with the |
|  |  | terms and manner prescribed in the Schedule of Requirement |
|  | 12.2 | All costs associated with the transportation of the goods subject |
|  |  | to this contract shall be borne by the Supplier. |
| **13. Incidental** | 13.1 | The Supplier shall be required to provide the incidental services |
| **Services** |  | as specified in the SCC and the cost of which is included in the |
|  |  | total bid price. |
| **14. Warranty** | 14.1 | All goods subject to this contract shall be accompanied by the |
|  |  | necessary warranty in the manner prescribed in the SCC. |
|  | 14.2 | The Purchaser shall promptly notify the Supplier in writing of |
|  |  | any claims arising under this warranty. |

|  |  |  |
| --- | --- | --- |
| **15. Payment** | 15.1 | The purchaser shall make payments to the Supplier in |
|  |  | accordance with the conditions set forth in the Payment |
|  |  | Schedule agreed and annexed to this contract. |
|  | 15.2 | The currency of payment shall be Pakistan Rupee. |
| **16. Prices** | 16.1 | Prices charged by the Supplier for goods delivered under the |
|  |  | Contract shall not vary from the prices quoted by the Supplier |
|  |  | in its bid and shall remain the same till the expiry of the contract |
|  |  | unless the Parties to this contract mutually agree to vary the |
|  |  | prices. |
| **17. Contract** | 17.1 | No variation in or modification of the terms of the Contract shall |
| **Amendments** |  | be made except by written amendment signed by the Parties. |
| **18. Assignment** | 18.1 | The Supplier shall not assign, in whole or in part, its obligations |
|  |  | to perform under this Contract, except with the Purchaser’s |
|  |  | prior written consent. |
| **19. Subcontracts** | 19.1 | The Supplier shall not be allowed to sublet and award |
|  |  | subcontracts under this Contract. |
| **20. Delays in the** | 20.1 | Delivery of the goods shall be made by the Supplier in |
| **Supplier’s** |  | accordance with the time schedule prescribed by the Purchaser |
| **Performance** |  | in the Schedule of Requirements. In case the contractor fails to |
|  |  | adhere to the prescribed time schedule, the purchaser is at |
|  |  | liberty to make risk purchases at the risk & cost of the |
|  |  | contractor in the best public interest. |
|  | 20.2 | If at any time during performance of the Contract, the Supplier |
|  |  | encounters conditions impeding timely delivery of the goods, |
|  |  | the Supplier shall promptly notify the Purchaser in writing of the |
|  |  | fact of the delay, its likely duration and its cause(s). As soon as |
|  |  | practicable after receipt of the Supplier’s notice, the Purchaser |
|  |  | shall evaluate the situation and may at its discretion extend the |
|  |  | Supplier’s time for performance, with liquidated damages, in |
|  |  | which case the extension shall be ratified by the Parties by an |
|  |  | amendment to the Contract. |
|  | 20.3 | Except as provided under GCC Clause 20, a delay by the |
|  |  | Supplier in the performance of its delivery obligations shall |

render the Supplier liable to the imposition of liquidated damages as prescribed in the SCC, unless the parties to this contract mutually agree for extension of time.

|  |  |  |  |
| --- | --- | --- | --- |
| **21. Termination for** | 21.1 The Purchaser, without prejudice to any other | remedy | for |
| **Default** | breach of Contract, by written notice of default | sent to | the |
|  | Supplier, may terminate this Contract in whole or in part: |  |

1. if the Supplier fails to deliver any or all installments of the goods within the period(s) specified in the signed contract, and subsequent Purchase order or within any extension thereof granted by the Purchaser pursuant to GCC Clause 20; or
2. if the Supplier fails to perform any other obligation(s) under the Contract.
3. if the Supplier, in the judgment of the Purchaser has engaged in corrupt or fraudulent practices in competing for or in executing the Contract.

For the purpose of this clause Corrupt and fraudulent practices means:

*“the offering, giving, receiving, or soliciting of anything of value to influence the action of a public official or the contractor in the procurement process or in contract execution to the detriment of the procuring agency; or misrepresentation of facts in order to influence a procurement process or the execution of a contract, collusive practices among bidders (prior to or after bid submission) designed to establish bid prices at artificial, non-competitive levels and to deprive the procuring agency of the benefits of free and open competition and any request for, or solicitation of anything of value by any public official in the course of the exercise of his duty; it may include any of the following practices:*

1. *coercive practice by impairing or harming, or threatening to impair or harm, directly or indirectly, any party or the property of the party to influence the actions of a party to achieve a wrongful gain or to cause a wrongful loss to another party;*
2. *collusive practice by arrangement between two or more parties to the procurement process or contract execution, designed to achieve with or without the knowledge of the procuring agency to establish prices at artificial, noncompetitive levels for any wrongful gain;*
3. *corrupt practice by offering, giving, receiving or*

*soliciting, directly or indirectly, of anything of value to influence the acts of another party for wrongful gain;*

* 1. *fraudulent practice by any act or omission, including a misrepresentation, that knowingly or recklessly misleads, or attempts to mislead, a party to obtain a financial or other benefit or to avoid an obligation;*
	2. *obstructive practice by harming or threatening to harm, directly or indirectly, persons or their property to influence their participation in a procurement process, or affect the execution of a contract or deliberately destroying, falsifying, altering or concealing of evidence material to the investigation or making false statements before investigators in order to materially impede an investigation into allegations of a corrupt, fraudulent, coercive or collusive practice; or threatening, harassing or intimidating any party to prevent it from disclosing its knowledge of matters relevant to the investigation or from pursuing the investigation, or acts intended to materially impede the exercise of inspection and audit rights;*
1. Indulgence in corruption and fraudulent practices is liable to result in rejection of Bids, cancellation of contracts, debarring and blacklisting of the Bidder, for a stated or indefinite period of time.
2. The following are the events which would lead to initiate under Rule 21 of PPRA Rules 2014 Blacklisting / Debarment process;
	* 1. Submission of false fabricated / forged documents for procurement in tender.
		2. Not attaining required quality of work.
		3. Inordinate tardiness in accomplishment of assigned/agreed responsibilities / contractual obligations resulting loss to procuring agency / Government.
		4. Non execution of work as per terms & condition of contract.
		5. Any unethical or unlawful professional or business behavior detrimental to good conduct and integrity of the public procurement process.
		6. Involvement in any sort of tender fixing.
		7. Persistent and intentional violation of important conditions of contract

1. Non-adherence to quality specification despite being importunately pointed out.
2. Security consideration of the State i.e., any action that jeopardizes the security of the State or good repute of the procuring agency.

**PROCEDURE:** A notice will be issued to theagency/individual seeking it/his explanation for the lapses committed by it/him. The explanation will be required within \_\_\_\_\_ days from the date of issue, (time will be fixed depending upon the intensity of lapses). In case its/his explanation is found unsatisfactory, a show cause notice shall be issued providing an opportunity of being heard followed by decision for blacklistment for a maximum period of three years depending upon the intensity of lapses. The letter for debarring the agency/individual will be published on PPRA website. Once the blacklisting order is issued it shall not be revoked ordinarily unless as provided under Rule-21 of the procurement Rules 2014.

*“the offering, giving, receiving, or soliciting of anything of value to influence the action of a public official or the contractor in the procurement process or in contract execution to the detriment of the procuring agency; or misrepresentation of facts in order to influence a procurement process or the execution of a contract, collusive practices among bidders (prior to or after bid submission) designed to establish bid prices at artificial, non-competitive levels and to deprive the procuring agency of the benefits of free and open competition and any request for, or solicitation of anything of value by any public official in the course of the exercise of his duty; it may include any of the following practices:*

1. *coercive practice by impairing or harming, or threatening to impair or harm, directly or indirectly, any party or the property of the party to influence the actions of a party to achieve a wrongful gain or to cause a wrongful loss to another party;*
2. *collusive practice by arrangement between two or more parties to the procurement process or contract execution, designed to achieve with or without the knowledge of the procuring agency to establish prices at artificial, noncompetitive levels for any wrongful gain;*

1. *corrupt practice by offering, giving, receiving or soliciting, directly or indirectly, of anything of value to influence the acts of another party for wrongful gain;*
2. *fraudulent practice by any act or omission, including a misrepresentation, that knowingly or recklessly misleads, or attempts to mislead, a party to obtain a financial or other benefit or to avoid an obligation;*
3. *obstructive practice by harming or threatening to harm, directly or indirectly, persons or their property to influence their participation in a procurement process, or affect the execution of a contract or deliberately destroying, falsifying, altering or concealing of evidence material to the investigation or making false statements before investigators in order to materially impede an investigation into allegations of a corrupt, fraudulent, coercive or collusive practice; or threatening, harassing or intimidating any party to prevent it from disclosing its knowledge of matters relevant to the investigation or from pursuing the investigation, or acts intended to materially impede the exercise of inspection and audit rights;*

|  |  |
| --- | --- |
| **22. Force Majeure** | 22.1 Notwithstanding the provisions of GCC Clauses 20 and 21, the |
|  | Supplier shall not be liable for forfeiture of its Performance |
|  | Guaranty, or termination/ blacklisting for default if and to the |
|  | extent that it’s delay in performance or other failure to perform |
|  | its obligations under the Contract is the result of an event of |
|  | Force Majeure. For the purposes of this clause Force Majeure |
|  | means an act of God or an event beyond the control of the |
|  | Supplier and not involving the Supplier’s fault or negligence |
|  | directly | or | indirectly | purporting | to | mis-planning, |

mismanagement and/or lack of foresight to handle the situation. Such events may include but are not restricted to acts of the Purchaser in its sovereign capacity, wars or revolutions, fires, floods, earthquakes, strikes, epidemics, quarantine restrictions and freight embargoes.

1. If a Force Majeure situation arises, the Supplier shall promptly notify the Purchaser in writing with sufficient and valid evidence of such condition and the cause thereof. The Purchaser shall examine the merits of the case and all reasonable alternative means for completion of the purchase order under the signed contract and inform the Supplier of its findings promptly.

1. Unless Purchaser informs the Supplier in writing of its agreement on the application of force majeure, the Supplier shall continue to perform its obligations under the Contract as far as is reasonably practical and shall seek reasonable alternative means for performance not prevented by the Force Majeure event.

|  |  |  |
| --- | --- | --- |
| **23. Termination for** | 23.1 | The Purchaser may at any time terminate the Contract by |
| **Insolvency** |  | giving written notice of one month time to the Supplier if the |
|  |  | Supplier becomes bankrupt or otherwise insolvent. In this |
|  |  | event, termination shall be without compensation to the |
|  |  | Supplier, provided that such termination shall not prejudice or |
|  |  | affect any right of action or remedy which has accrued or shall |
|  |  | accrue thereafter to the Parties. |  |
| **24. Arbitration and** | 24.1 | The Purchaser and the Supplier shall make every effort to |
| **Resolution of** |  | resolve amicably by direct informal negotiation any |
| **Disputes** |  | disagreement or dispute arising between them under or in |
|  |  | connection with the Contract. |  |
|  | 24.2 | If, after thirty (30) days from the commencement of suchinformal negotiations, the Purchaser and the Supplier have been unable to resolve amicably a Contract dispute, either party may require that the dispute be referred to the Arbitrator for resolution through arbitration. |  |
|  | 24.3 | In case of any dispute concerning the interpretation and/orapplication of this Contract shall be settled through arbitrationunder the Arbitration Act of 1940 (As amended from time to time). |
|  |  |
|  |  |
| **25. Governing** | 25.1 | The Contract shall be written in English language. Subject to |
| **Language** |  | GCC Clause 26, the version of the Contract written in the |
|  |  | specified language shall govern its interpretation. | All |
|  |  | correspondence and other documents pertaining to the |
|  |  | Contract, which are exchanged by the Parties, shall be written |
|  |  | in English. |  |
| **26. Applicable** | 26.1 | This Contract shall be governed by the Laws of Pakistan and |
| **Law** |  | the courts of Pakistan shall have exclusive jurisdiction. |  |
| **27. Notices** | 27.1 | Any Notice given by one party to the other pursuant to this Contract shall be sent to the other party in writing and on the others address specified in SCC. |
|  | 27.2 | A notice shall be effective when delivered or on the notice’s effective date, whichever is later. |
| 1. **Taxation**
 | 28.1 | All taxation, whether International, Federal, Provincial or Local, shall be borne by the Supplier. |