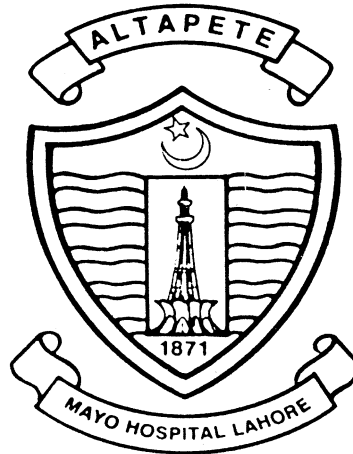


BIDDING DOCUMENTS



**BIDDING DOCUMENT APPLICABLE TO PROCUREMENT OF GOODS
(DRUGS / MEDICINES / SURGICAL DISPOSABLES / IMPLANTS / CARDIAC
SURGERY / CARDIOLOGY DISPOSABLES / BME / DENTAL / X-RAY & C.T
SCAN/ CONSUMABLES / THERAPEUTIC GOODS / STATIONARY /
GENERAL STORE / MIR / ELECTRIC / LAB KITS AND CHEMICAL /
BEDDING CLOTHING AND LINEN / ORTHOPEDIC RAW MATERIAL /
GLASS WARES / SANITATION ETC. EXCEPT ELECTRO MEDICAL
EQUIPMENTS ON FRAMEWORK CONTRACT BASIS DURING FINANCIAL
YEAR 2023-2024.**

Name of Procuring Agency	MAYO HOSPITAL LAHORE
Corresponding Address:	Chief Executive Officer / D.D.O., Mayo Hospital Lahore Near by Nila Gumbad Lahore
Telephone No(s)	+92-42-99211139-110,117,378 & 381
Fax No.	+92-42-99211115
Email	mayohospital@gmail.com

IMPORTANT NOTICE FOR BIDDERS

All the bidders who intend to participate, are hereby informed that the bidding document is prepared carefully by the committee, however following under mentioned references will also be applicable / binding upon bidders if and when required.

1. Procurement will be governed by PPRA Rules 2014 amended (till date). (<https://ppra.punjab.gov.pk/>).
2. Policies, Guidelines & Instructions (if and when notified / circulated) by the Government of the Punjab, Specialized Healthcare & Medical Education Department / Mayo Hospital (Authority), shall be applicable / binding upon the bidders.
3. Offered Product specification shall match with the advertised specification as well as notified specification by DRAP and Drug ACT 1976. (<http://www.dra.gov.pk/>)

Bid Data Sheet

Description	Detail
Language of bid	English or Urdu
Bid currency	Pak Rs. On free delivery to Consignee's end basis including all Ex-work, Transportation, Storage charges till the destination.
Bid Security	2% of estimated cost in the shape of Bank Guarantee / CDR from any scheduled bank The bid security shall be in the shape of Bank Guarantee / CDR and item wise. However if any bidder opts to bid for more than one item and opts to submit bid security in lump sum, in such cases if the bid security amount is found lesser than the required amount , then the total bid of the said bidder shall be rejected.
Bid validity period	180 Days
Address for communication:	
Name of Procuring Agency	MAYO HOSPITAL LAHORE
Corresponding Address:	Chief Executive Officer / Mayo Hospital Lahore Near by Nila Gumbad Lahore
Telephone No(s)	+92-42-99211129-110,117,378 & 381
Fax No.	+92-42-99211115
Email	mayohospital@gmail.com
<u>Performance Guarantee:</u> It will be 5% of the Contract Value in the shape of Bank Guarantee / CDR from any scheduled bank.	



INVITATION TO BIDS

MAYO HOSPITAL LAHORE

Bid Reference No.

Subject: FRAMEWORK CONTRACT FOR PROCUREMENT OF GOODS (DRUGS / MEDICINES / SURGICAL DISPOSABLES / IMPLANTS / CARDIAC SURGERY / CARDIOLOGY DISPOSABLES / BME / DENTAL / X-RAY & C.T SCAN/ CONSUMABLES / THERAPEUTIC GOODS / STATIONARY / GENERAL STORE / MIR / ELECTRIC / LAB KITS AND CHEMICAL / BEDDING CLOTHING AND LINEN / ORTHOPEDIC RAW MATERIAL / GLASS WARES / SANITATION ETC. EXCEPT ELECTRO MEDICAL EQUIPMENTS ON FRAMEWORK CONTRACT BASIS DURING FINANCIAL YEAR 2023-2024.

Dated: _____

Dear Sir /Madam

Mayo Hospital Lahore invites sealed bids/tenders (Technical & Financial) to conclude the Framework contract for the supply of Goods on free delivery to Consignee's end Basis **Mayo Hospital, Lahore.**

1. Pharmaceutical Manufacturers /Importer /authorized Agents .General Order supplier (for general store related items) may participate in bid for the year **2023-24**. The contract shall be valid for one (1) year from the date of issuance of advance acceptance letter/ notification of Award.
2. Authority letter shall be in the name of Chief Executive Officer, Mayo Hospital Lahore by the manufacturer regarding items to participate in the Bid.
3. The firm must participate in the bidding process for their **items / section** against the total quantity.
4. Bidding shall be conducted through 38(2) (a) single stages-two envelopes bidding procedure of Punjab Public Procurement Rules, 2014 (amended). The bids shall clearly be **marked with Bid Enquiry No.** for each the proposal is submitted.
5. Bidder can download the Bidding Documents containing tender's item specifications, quantity and terms & conditions from PPRA website of www.mayohospital.gop.pk until the closing date for the submission of bids.
6. Sealed bids are required to be submitted by the Bidders as per advertised schedule positively in **the Committee Room**, of the **Mayo Hospital Lahore**. The bids received till stipulated date & time shall be opened on same day at (**as per schedule**) in the presence of the bidders or their authorized representatives who choose to attend. Late bids shall not be entertained.
7. All bids should be submitted in Tape Binding. All documents should contain proper page marking, attached in sequence as indicated for evaluation in the Bidding Documents and Signatures of authorized person. Moreover, signing and stamping of each page of bidding documents/ form is mandatory.
8. In case the date of opening is declared as 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 submission and opening of tenders accordingly. The time and venue shall be remained the same.
9. The intending bidders may get the tender/ bidding document and detail specification from the Almoner Office during office hours along with the payment of non-refundable fee of Rs. 5000 (five thousand only) for each set of bidding document and detail specification.

Note: in case of pre bid meeting it will be held in committee room while the time & date of meeting and minutes of meeting will be mentioned in the tender notice. The minutes of the pre bid meeting will be uploaded on official website of Mayo Hospital Lahore, and will the part of bidding document.


**CHIEF EXECUTIVE OFFICER,
MAYO HOSPITAL, LAHORE.**

REQUIRED ESSENTIAL DOCUMENTS / CHECK LIST (TO BE FILLED BY THE BIDDER)

Sr.No.	Description	Documents Attached	Page No.
01	Attested copy of CNIC of authorized bidding Signatory person of the Bidder	Yes / No.	
02	Tender purchase / sale Receipt	Yes / No.	
03	Bid Security (Bank Guarantee / CDR)	Yes / No.	
04	Drug Reg. Certificate	Yes / No.	
05	Drug Sale License	Yes / No.	
06	Authority Letter from Manufacturer / Importer / Local Manufacturer / authorization certificate.	Yes / No.	
07	c.G.M.P./ final inspection report (if applicable)	Yes / No.	
08	Undertaking that bidder is not blacklisted	Yes / No.	
09	Undertaking that DML/DRC is not canceled / suspended by DRAP. (if applicable)	Yes / No.	
10	Undertaking that the DTL has declared None of the offered product as spurious / Adulterated / Substandard etc.	Yes / No.	
11	FBR Registration Certificate / Income Tax Registration	Yes / No.	
12	Company Registration Certificate	Yes / No.	
13	List of Sample of the quoted items	Yes / No.	
14	All requisite documents must be attached according to evaluation criteria i.e. compulsory parameters as well as ordinary parameters.	Yes / No.	
15	The documents required for ordinary Parameters should be attached according to the detail mentioned against each requirement.	Yes / No.	
Note. The Bidder (s) is directed to follow the above mentioned sequence while preparing their bid.			

INSTRUCTIONS TO BIDDERS

1. Scope of Bid

Mayo Hospital, Lahore invites bids for supply of goods as per enclosed list along with Technical Specifications.

2. Source of Funds

The Government of Punjab, allocated funds to Mayo Hospital, Lahore for the purpose of procurement of advertised goods/ items etc. for the Financial Year 2023-24.

3. Eligible bidders

3.1 This Invitation for Bids is open to all original Manufacturers / Importer/ Authorized Agent General Order Supplier (for General Store Related Items).

3.2 Government-owned enterprises in Pakistan may participate only if they are legally and financially autonomous and authorized to participate in bidding.

3.3 The Agent /supplier /importer must possess valid authorization form 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 manufacture, they should have a documentary prove as prescribed in section V , bid form, to the effect that they are the original manufacturer of the required specifications of goods.

3.4 Bidders under a declaration of ineligibility for corrupt and fraudulent practices, issued by any Government (Federal/Provincial/Local), or a Public Sector Organization are **Not Eligible**.

4. Corrupt and Fraudulent Practices and mechanism to debar /Black list the defaulted bidder .

4.1 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 contractor in the procurement process or in contract execution to the detriment of the Procuring Agencies; or misrepresentation of facts in order to influence a procurement process or the execution of 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 agencies 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.

(i) 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;

(ii) 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;

(iii) 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;

(iv) 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;

(v) 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;

4.2 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.

4.3 The following are the events which would lead to initiate under Rule 21 of PPRA Rules 2014 Blacklisting/Debarment process:

- i. Submission of false fabricated / forged documents for procurement in tender.
- ii. Not attaining required quality of work.
- iii. Inordinate tardiness in accomplishment of assigned/agreed responsibilities / contractual obligations resulting loss to procuring agency / Government.
- iv. Non execution of work as per terms & condition of contract.
- v. Any unethical or unlawful professional or business behavior detrimental to good conduct and integrity of the public procurement process.
- vi. Involvement in any sort of tender fixing.
- vii. Persistent and intentional violation of important conditions of contract
- viii. Non-adherence to quality specification despite being importunately pointed out.
- ix. 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 by the agency to the bidder seeking it/his explanation for the lapses committed by it/him. The explanation will be required within 7 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 blacklisting 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.

5. **Eligible Goods and Services:** All goods and related services to be supplied under the contract shall conform to the policies of the Government of the Punjab in vogue. All expenditures made under the contract shall be limited to such goods and services. For purpose of this clause, (a) the term "Goods" includes any Goods that are the subject of this Invitation for Bids and (b) the term "Services" includes related ancillary services such as transportation, insurance after sale services etc.
6. **Cost of Bidding:** The bidder shall bear all costs associated with the preparation and submission of its bid, and Mayo Hospital, Lahore shall in no case be responsible or liable for those costs, regardless of the conduct or outcome of the bidding process.
7. **Bidding for Selective Items.**

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

However, Bidders cannot bid for partial quantities of an item as mentioned in the Schedule of Requirement & Technical Specifications. The bid must be for the whole quantity of an item required in the schedule of requirement & Technical Specifications.

THE BIDDING PROCEDURE

8. **The Governing Rules.**
The Bidding procedure shall be governed by the Punjab Procurement Rules, 2014 amended of the Government of Punjab.
9. **Applicable Bidding Procedure.**
 - 9.1 The bidding procedure is governed by Rule 38 "Procedures for selection of contractors" sub-rule (2) (a) "Single stage-- two envelop 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 below.

Single Stage two envelopes Bidding Procedure

Single Stage two envelope Bidding Procedure shall be used for procurement of such goods where the bids to be evaluated on technical & financial ground and the procedure for single stage two envelopes shall be.

- i. The bid shall be a single package consisting of two separate envelopes, containing separately the Financial & the Technical Proposals;
- ii. The Envelopes shall be marked as “FINANCIAL PROPOSAL” and “TECHNICAL PROPOSAL”
- iii. In the first Instance, the "TECHNICAL PROPOSAL" shall be opened; and the enveloped marked as financial proposal shall be retained unopened in the custody of the procuring agency.
- iv. 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;
- v. During the technical evaluation, no amendments in the technical proposal shall be permitted;
- vi. After the evaluation and approval of the technical proposal, the procuring agency shall open the financial proposal of the technically accepted bids, publically at time, date and venue announced and communicated to the bidder in advance, within the bid validity period.
- vii. The financial bids found technically non-responsive shall be returned unopened to the respective bidders;
- viii. The lowest evaluated bidder shall be awarded the contract.

THE BIDDING DOCUMENTS

10 Contents of Bidding Documents

10.1. The goods required, applicable bidding procedures, and Contract terms are prescribed in the bidding documents. In addition to the Invitation for Bids, the bidding documents include:-

- a. Instructions to bidders;
- b. Schedule of Requirements & Technical Specifications
- c. Evaluation Criteria
- d. Bid Forms
 - i) Letter of Intension,
 - ii) Affidavit,
 - iii) Technical Forms,
 - iv) Financial Forms)
- e. Draft Standard Contract
 - i) Contract Form
 - ii) General Conditions of Contract;
 - iii) 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 said Bidding Documents shall take precedence.

10.3. 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.

11. Clarification(s) on Bidding Documents

A prospective bidder requiring any clarification(s) on the bidding documents may notify the Chief Executive Officer, Mayo Hospital, Lahore, in writing. The procuring Agency shall respond in writing to any request for clarification(s) of the bidding documents, which it receives no later than Ten (10) days, prior to the deadline for the submission of bids prescribed in the Invitation for Bids. Written copies of the procuring agency’s response (including an explanation of the query but without identifying the source of inquiry) shall be sent to all prospective bidders that have received the bidding documents.

12. Amendment(s) to Bidding Documents

12.1 At any time prior to the deadline for submission of bids, 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 in writing amendment(s). (If required)

- 12.2 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.
- 12.3 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

13. Language of Bid

All correspondence, communications associated with preparation of Bids, clarifications, amendments, submissions, shall be written 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.

14. Documents Comprising the Bids

- 14.1 The bid shall comprise of the Bid Forms of this Bidding Documents and all those ancillary documentations that are prescribed for the eligibility of the bidders and goods and ancillary services that are found necessary and highlighted in bid forms.
- 14.2 The bidder shall complete the bid forms and an appropriate price schedule 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.

15. Bid Price

- 15.1 The Bidder shall indicate on the appropriate form, prescribed in this Bidding Document, the unit prices and total bid price of the goods, it proposes to supply under the Contract.
- 15.2 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.
- 15.3 The Bidder should quote the prices of goods according to the technical specifications as provided in of this document. The technical specifications of goods, different from the required specifications, shall straightway be rejected.
- 15.4 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.
- 15.5 The benefit of exemption from or reduction in the taxes and duties shall be passed on to the Procuring Agency. (where applicable).
- 15.6 Prices offered should be for the entire quantity of an item demanded in the Schedule of Requirement & technical specifications; partial quantity offers shall straightaway be rejected. Conditional offer shall also be considered as non-responsive Bid.
- 15.7 No request for increase in price due to market fluctuation in the cost of goods and services shall be entertained.

16. Bid currencies:

Prices shall be quoted in Pak Rupees.

17. Samples

- 17.1 The bidder shall provide the requisite number of samples (prescribed in schedule of requirement) of the quoted product along with bid.

18. Documentations on Eligibility of bidders

- 18.1 The bidder shall furnish, as part of its technical bid (Bid form) as specified, documents establishing the bidder's eligibility to bid and its qualifications to perform the Contract if its bid is accepted.
- 18.2 The documentary evidence of the bidder's eligibility to Bid shall be established to the Procuring Agency's satisfaction that the bidder, at the time of submission of its bid, is an eligible as defined under instruction to the bidders clause 3 above .

19. Documentations on Eligibility of goods

The bidder shall furnish as part of its Bid (Bid form) as specified, documents establishing the eligibility and conformity to the bidding documents of all goods, which the bidder proposes to supply under the Contract.

20. Bid Security

20.1 The bidder shall furnish separately against each quoted item, as part of its Financial Bids, in the shape of **only Bank Guarantee / CDR from any scheduled Bank** in the name of Chief Executive Officer, Mayo Hospital Lahore .Failure to furnish the prescribed bid security / lesser bid security than prescribed shall result in the rejection of bids .The bid security must have a minimum validity period of 180 days from the last date for submission of the bids or until furnishing of the performance security , whichever is later.

20.2 The Bid Security shall be forfeited by the purchaser, on the occurrence of any / all of the following conditions:

- i). if the bidder withdraws its bid during the period of bid validity specified in bidding documents or
- ii) If the bidder does not accept the correction in his total bid price or.
- iii) If the bidder , having been notified for the acceptance of the bid by the purchasers during the period of bid validity, fails or refuses to furnish the performance security, in accordance with the bidding documents.

20.3 Unsuccessful bidders, 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.

20.4 The bid security shall be in the shape of Bank Guarantee / CDR and item wise. However any bidder opts to bid for more than one item and opts to submit bid security in lump sum, in such cases if the bid security amount is found lesser than the required amount, then the total bid of the said bidder shall be rejected.

21. Bid Validity

21.1 Bids shall remain valid for the period of 180 Days after the date of opening of Technical Bid. A bid valid for a shorter period shall be rejected by the Procuring Agency as non-responsive.

21.2 The Procuring Agency, shall ordinarily be under an obligation to process and evaluate the bid within the stipulated bid validity period, But under exceptional circumstance 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.

21.3 A bidder who,-

- (a) Agree to the extension of bid validity period shall also extend the validity of bid security for the extended period of the bid validity.
- (b) Agrees to the procuring Agency's request for extension of bid validity period shall not be permitted to change the substance of the bid
- (c) Does not agree to an extension of bid validity period shall be allowed to withdraw the bid without forfeiture of the bid security.

22. Format and Signing of Bid:

22.1 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.

22.2 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.

22.3 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 documents and the whole bid must be signed and stamp by the authorized person. Moreover, signing and stamping of each page of bidding document/form is mandatory.

22.4 Any interlineations, erasures, or overwriting shall be valid only if they are initialed by the person or persons signing the bid.

SUBMISSION OF BIDS

23. Sealing and Marking of Bids

- 23.1 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.
- 23.2 The inner and outer envelopes shall:
- a. be addressed to the Procuring Agency at address given in the invitation for bids and .
 - b. Bid reference number indicated in the Bid data sheet, Tender Enquiry No. Indicated in Section III , Schedule of Requirements & Technical Specification and statement: DO NOT OPEN BEFORE “ , the time and the date specified in the Bid data sheet for opening of bids.
- 23.3 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".
- 23.4 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.

24. Deadline for Submission of Bids:

- 24.1 All bids should be submitted in tape binding. 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 at its discretion, extend the prescribed deadline for the submission of bids by amending the bidding documents in accordance with IBT 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.

25. Late Bids:

- 25.1 Any bid received by the Procuring Agency after the deadline for submission of bids prescribed Procuring Agency Pursuant to ITB Clause 24 shall be rejected and returned unopened to the bidder.

26. Withdrawal of Bids:

- 26.1 The bidder may withdraw its bid after the bid's submission and prior to the deadline prescribed for submission of bids.
- 26.2 No bid may be withdrawn in the period between the 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

27. Opening of Bids by Procuring Agency

- 27.1 **All bids received**, shall be open 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 Bids data sheet.
- 27.2. The opening of the bids shall be subject to the bidding procedure prescribed in the Bid Data Sheet and elaborated in ITB clause of above.
- 27.3 All Bidders in attendance shall sign on attendance sheet.
- 27.4 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 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 PPR-2014, specifically Rule 30 (Opening of Bids).
- 27.5 The Procuring Agency have the minutes of the bid opening (Technical & When Applicable, Financial) recorded.
- 27.6 No bid shall be rejected at technical proposal / bid opening, except for late bids, which shall be returned un-opened to the bidder.
- 27.7 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 statement / reason on such bids.

- 27.8 No Tenderer or its representative will be allowed to keep any digital device during tender opening meeting at given time and location.

28. Clarification of Bids:

- 28.1 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.

29. Preliminary Examination

- 29.1 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.
- 29.2 In the financial bids the arithmetical errors shall be rectified on the following basis:
- a. 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.
- b. If the bidder does not accept the correction of the errors, its bid shall be rejected, and its bid Security may be forfeited.
- c. If there is a discrepancy between words and figures, the amount in words shall prevail.
- 29.3 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 Law, Taxes & Duties and internationally recognize best practices shall be deemed to be a material deviation for technical proposals and Bid Security for financial proposals. The Procuring Agency determination of a bid's responsiveness is to be based on the contents of the bid itself without recourse to extrinsic evidence.
- 29.4 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.

30. Evaluation of Bids

- 30.1 The Procuring Agency shall evaluate and compare the bids, which have been determined to be substantially responsive in accordance with ITB Clause 29 above.
- 30.2 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.
- 30.3 The Technical Evaluation Committee (TEC) will submit the Evaluation report (including the status of the bidder in all parts of technical evaluation criteria that is part A, B and C) and all parameters will be checked simultaneously.
- 30.4 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.
- 30.5 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.

31. Qualification of Bidder

- 31.1 The Procuring Agency, at any stage of the procurement proceedings, having credible reasons for or prime 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.
- 31.2 Such qualification shall only be laid down after recording reasons thereof in writing. They shall form part of the records of that procurement proceeding.
- 31.3 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.
- 31.4 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.

31.5 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.

31.6 The Procuring Agency shall disqualify a contractor on the ground that he had provided false, fabricated or materially incorrect information.

32. Rejection of Bids

32.1 The Procuring Agency may reject any or all bids at any time prior to the acceptance of a bid or proposal as prescribed in Rule 35 of Punjab Procurement Rules -2014. The Procuring Agency shall upon request communicate to any bidder who submitted a bid, the grounds for its rejection of any or all bids, but shall not be required to justify those grounds.

32.2 The Procuring Agency incurs no liability, solely by virtue of its invoking Clause 32.1 towards Bidders who have submitted bids.

32.3 Notice of the rejection of any or all bids shall be given promptly to the concerned bidders which submitted bids.

33. Re-Bidding

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.

34. Announcement of Evaluation Report

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

35. Contacting the Procuring Agency.

35.1 Subject to IBT 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 Report. If a Bidder wishes to bring additional information to the notice of the Procuring Agency, it should do so in writing.

35.2 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.

Award of Contract

36. Acceptance of Bid and Award Criteria

The bidder, 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.

37. Procuring Agency's Right to vary quantities at the time of Award

The Procuring Agency reserves the right at the time of award of contract to increase or decrease, the quantity of goods originally specified in schedule of Requirements & Technical Specifications without any change in unit price and other terms & conditions.

38. Notification of Award.

38.1 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.

38.2 The notification of award shall constitute the formation of the contract between the Procuring Agency and the successful bidder.

38.3 The enforcement of the contract shall be governed by Rule 63 of the PPR-2014(Amended).

39. Limitation on Negotiations.

Save and otherwise provided in PPRA Rules 2014(Amended), Procuring Agency shall not negotiate with any bidder.

40. Signing of Contract.

40.1 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.

40.2 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.

40.3 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.

40.4 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 and relevant laws/Rules.

40.5 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 Fiancé Act 1995 (Act-VI of 1995) Notification No. JAWH/HD /8-21/77(PG) dated 1st January, 2014.

41. Performance Guarantee.

41.1 Successful Bidder shall furnish a Performance Guarantee in the shape of Bank Guarantee / CDR in the name of Chief Executive Officer, Mayo Hospital, Lahore.

41.2 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.

41.3 Failure to provide the 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.

42. Price Reasonability Certificate

The supplier shall certify on judicial stamp paper that the prices quoted are not more than the trade prices as per MRP (Maximum Retail Price) fixed by the Federal Government under Drugs Act, 1976 /DRAP Act, 2012 (If applicable).

43. All supplies will comply with the provision of Drugs Act 1976/DRAP Act, 2012 and rules framed there under / notifications issued by the Federal /Punjab Govt. (If applicable)

TORs FOR EVALUATION/PROCUREMENT OF ORTHOPEDIC IMPLANTS .

- i. All branded supplies (Local / imported) should be randomly evaluated by testing from PCSIR laboratories and cost of testing will be paid by related firm.
- ii. In case of supply of damaged or poor quality implants, the item will be returned back and penalty will be in accordance to the TORs of contract/ bidding document.
- iii. Demand will generate every three month (as per requirements of the end user).
- iv. All vendors will ensure to provide tools / instrumentation of the same brand and technical staff for the safe application of implants.
- v. Technical material specification either stainless steel or titanium should be provided and rate contracted.

SCHEDULE OF REQUIREMENTS & TECHNICAL SPECIFICATIONS:

Schedule of Requirements:

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

Respective Consignee's End: *Name & Address of Procuring Agency*

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 as described in purchase order.
With recovery of late delivery charges @ 0.067 % per day.	After 45 days or earlier (as described in purchase order), decided by the concerned Consignee on formal request of supplier with proper justification.
Maximum Rate of Late Delivery Charges	Maximum limit of <i>Late Delivery Charges</i> is 10% of the contract after which contract will be cancelled with all legal and codal formalities
Risk Purchase	After expiry of prescribed delivery period the Procuring Agency will proceed for risk purchases (at the risk & cost of defaulter) to ensure the un-interrupted healthcare services to the patients. The risk purchase will be automatically done if no request of bidder is received for grace period or stock is nil.

- Separate Purchase order(s) shall be issued, out of the total advertised quantities as per schedule mentioned in list of requirements during the contract period as per storage capacity, consumption of the previous stock to avoid any untoward situation regarding short fall in shelf life, efficacy, etc. Payment shall be made on production of inspection certificate and receipt certificate from the consignee (Store Department) after recovery of Government dues including professional taxes. Part supply and part payment may be allowed.

LIST OF REQUIRED PRODUCTS WITH QUANTITIES AND TECHNICAL SPECIFICATIONS ETC

BID ENQUIRY NO.	GENERIC	SPECIFICATIONS	ESTIMATED COST PER UNIT	TOTAL QTY	2% OF THE ESTIMATED TOTAL COST (Required for Bid Security)	REQUIRED SAMPLE
LIST ATTACHED						

NOTE: -

- The bidder shall provide samples in No. as mentioned in the list of demand of required products. However in case, the technical evaluation committee needs samples (Medicines/Drugs only), the bidder is bound to provide the requisite number of samples as defined by the Chairman TEC/End-user with in the period specified. Failure to provide sample in time shall lead to disqualification of the firm.
- For thermo-labile drugs for which storage temperature is 2-8 degree centigrade. The firm shall be bound to reduce batch wise cold chain data from the source of origin & thermo log data from factory to Consignee's end.
- Sensitivity discs for antibiotics shall be provided by the successful bidder along with supply of Antibiotics free of cost.
- Samples for evaluation shall be submitted on the same day of submission of bid by the bidder at relevant Store and store will retain 01 sample (with sign & date) throughout the validity of contract agreement.

EVALUATION CRITERIA FOR DRUGS / MEDICINE

Part-A.

COMPULSORY PARAMETERS:

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

Sr. No.	PARAMETERS	DOCUMENTS REQUIRED	REMARKS
1	CNIC	Copy of valid CNIC of signatory person	
2	Bid Security	In the shape of Bank Guarantee / CDR in the name of Chief Executive Officer Mayo Hospital, Lahore.	
3	Receipt	Original Tender Purchase Receipt issued by Almoner Office, Mayo Hospital, Lahore.	
4	Drug Manufacturing /Drug Sale License	Copy of Drug Manufacturing License / valid Drug sale License applicable for importers.	
5	Drug Registration Certificate (DRC)	Copy of Drug Registration Certificate of quoted drugs.	
6	Authority letter for participation in tender at MHL	i) Manufacturer ii) Importer iii) Sole Agent iv) Authorized sole distributor for Mayo Hospital	
7	Product Experience	Products (locally manufactured) having less than ONE year experience shall be ineligible (Experience shall be calculated from the date of registration of the Product with the DRAP. (For drugs and medicines).	
8	Current Good Manufacturing Practices (cGMP) Certificate	1. Copy of cGMP/ copy drug registration certificate of quoted product (for Local Manufacturer). 2. In case of imported product valid GMP certificate issued by the regulatory authority of manufacturer's country will be considered.	
9	Undertaking on Stamp Paper worth Rs:100 (Minimum)	Regarding i. NON Cancellation / Suspension of Drug Registration of quoted product of the bidder by Drug Regulatory Authority of Pakistan within last two years. ii. Non Declaration of Spurious/Adulterated batch by DTLs of the Punjab/any iii. Non blacklisting from any public procuring agency of Pakistan to the bidder. iv. Quoted Rates are not more than Market Rates and Rates quoted / approved by other Hospitals.	
10	Other Documents Required	NTN No. / Income tax registration certificate / sale tax registration certificate.	
11	Income Tax Returns	Latest tax return showing annual sale.	
12	Specification quoted in the technical offer will be verified from samples provided along with the bid.	Samples of quoted items.	

	Product that comply 100% with the advertised specifications and fulfill the requirements as per Labeling and Packing Rules 1986 shall be considered for evaluation		
13	Batch Capacity of the manufacturer for the Quoted Item / Product	Certificate(s) provided by the manufacturer.	

Part-B

TECHNICAL EVALUATION PARAMETERS:

The bid complying with compulsory parameter shall be evaluated for below mentioned Parameters:

Sr. #.	Parameters	Detail			Total Marks	Documents required	MARKS OBTAINED
1	Bidder Performance (Last two years)	Government / Semi-Government institutions served:			10	The Claim requires documentation (Purchase Orders/ Delivery Challans /Frame work agreements etc.) of the institution(s).	
		i.	1	2			
		ii.	2 to 3	4			
		iii.	4 to 5	6			
		iv.	6 to 7	8			
		v.	8 & above	10			
2	Product experience	i.	Sale/ supply of quoted item in leading Chain Pharmacies / Pharmacies / institutions for last 02 years	7	15	Commercial invoices for chain Pharmacies/ Purchase orders of any Government/ Semi-Government institution.	
		Ii	More than 02 up to 04 years	10			
		Iii	More than 04 years	15			
3	Quality Certificates of manufacturer	i.	US FDA registration / CE certification / WHO Prequalification / Pre-qualification from any Provincial / Federal Govt. Institution / Department	7	10	Valid copies of certificates/letters Required.	
		ii.	Valid ISO / equivalent certification.	3			
4	Credibility and certification of API / product	i.	Source of API approved by USFDA/FDA	10	10	Certificate of analysis of API from source of manufacturer for each product. The copies of the certificates will be provided by local manufacturer.	
		ii.	Certificate of analysis of finish product from the country of origin.	5			
5	Batch history last year	Annual production batch history advertised quantity:			10	Verifiable invoices, Certificate from the Incharge QA department of	
		I	More than 20 Number of batches of quoted item	10			

			manufactured during last 12 months			concerned manufacturer (for local manufacturer)	
		ii	At-least 16-20 Number of batches of quoted item manufactured during last 12 months	5			
		iii	At-least 10-15 Number of batches of quoted item manufactured during last 12 months	3			
		iv	Less than 10 Number of batches of quoted items manufactured during last 12 months	0			
6	Batch quality (on Stamp paper worth : 100 Rupees Minimum)	i.	No batch failed during last (03) three year of the quoted item from any Statutory lab.	5	5	The firm will provide undertaking in this regard. The purchaser reserves the right to verify the claim.	
		ii.	No Batch failed during last (02) year of the quoted item from any Statutory lab.	3			
7	FINANCIAL CAPACITY OF THE BIDDER (ANNUAL TURNOVER OF LAST FINANCIAL YEAR)						
	2000 Million or above					20	
	Between 1,000 Million to – 2000 Million					15	
	Between 500 Million to – 1000 Million					10	
	Less than 500 Million					05	
	The bidder will provide requisite documents i.e Federal Board Of Revenue documents showing the annual sale of the firm						
	Marks				80		

Overall recommendation _____

Total Marks = 80

Minimum Qualifying Marks = 65% of Total Marks = 52

EVALUATION CRITERIA FOR SURGICAL DISPOSABLE /MEDICAL DEVICES/ LAB KITS & CHEMICAL (REAGENTS ONLY).

Part-A COMPULSORY PARAMETERS:

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

Sr.No.	PARAMETERS	DOCUMENTS REQUIRED	STATUS
1	Product Registration Certificate	<ul style="list-style-type: none"> i. Valid Product Registration certificate issued by the DRAP where applicable. ii. Valid Product enlistment certificate issued by DRAP. (where applicable) (Exemption shall be granted in the light of SRO 224(I)/ 2023 issued by DRAP.) 	
2	Firm Establishment Certificate	Valid License to import/ Manufacturing and sale certificate issued by DRAP (where applicable)	
3	Notarized letter of authorization from manufacturer	Valid manufacturer's authorization from the Foreign Manufacturer with indication of manufacturing site and its location (For Importer/ Sole Agent / Authorize sole Distributor) for Mayo Hospital	
4	Product Quality Certificate	Valid quality certification of US FDA/JpMHLW/MDD/ of the quoted product	
5	Undertaking on Stamp Paper worth Rs:100 (Minimum)	<ul style="list-style-type: none"> Regarding i. Non Cancellation / Non Suspension of Registration of quoted product of the bidder by Drug Regulatory Authority of Pakistan within last two years. ii .Non blacklisting from any public procuring agency of Pakistan of the bidder. iii. Non declaration of spurious / adulterated by the DTL of the Punjab/ any competent lab of quoted items within last two years. 	
6	Other Documents Required	<ul style="list-style-type: none"> i. NTN No. / Income tax registration certificate / sale tax registration certificate. ii. Original Receipt of Tender Fee. iii. Copy of Bank Guarantee / CDR in the name of Chief Executive Officer Mayo Hospital Lahore in technical Bid iii. CNIC of signatory of the Bid. iv. Signed terms & conditions of bidding documents and acceptance of bid validity period (180 days) 	
7	Product Related Free Sale Certificate issued by the Regulatory Body of manufacturer country	<ul style="list-style-type: none"> I. The bidder will submit Pakistan Embassy attested "free sale certificate of the product" (Medical devices) bearing the brand name of the product in country of manufacturer(where applicable) ii. Affidavit of the sole agent that their product(s) are freely available with same brand name in the country of the manufacture for at least/ last two (02) years and is safe for human use (where applicable) 	

8	Specification quoted in the Technical offer will be verified from sample provided with the bid (Product that complies 100 % with the advertised specification and full fill the requirements as per Medical Devices rules will be considered for evaluation.	Sample of quoted item.	
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Recommendation for Part-A: _____

Part-B

ORDINARY PARAMETERS:

The bid complying with compulsory parameter shall be evaluated for below mentioned parameters:

Sr. #.	Parameters	Detail			Total Marks	Remarks	MARKS OBTAINED
1	Past Performance of the Bidder (Last two years)	Major institutions (Government / Semi-Government) served:			10	The Claim requires documentation (Purchase Orders, Receipt Certificates & Delivery Challans etc.) of the institution(s).	
		i.	1	2			
		ii.	2 to 3	4			
		iii.	4 to 5	6			
		iv.	6 to 7	8			
		v.	8 & above	10			
2	Market / institutional Experience of quoted Product.	i.	Market Availability of quoted product in leading Chain Stores/ Pharmacies / institutions for last 02 years	7	15	Market availability in leading Chain Stores, Pharmacies of quoted item will be calculated from the date of commercial invoice. The firm will attach purchase orders / delivery Challan of the quoted item of any Government/ Semi-Government institution.	
		ii	More than 02 up to 04 years	10			
		iii	More than 04 years	15			
3	Credibility & Certification of Manufacturer	i.	WHO / US FDA / CE certification / WHO Prequalification / Prequalification by Provincial or Federal Institutes.	7	10	Valid copies of certificates/letters Required.	
		ii.	Valid ISO certification. (Notarized ISO) /international reputed certificate.	3			
4	Batch quality For Last Three Years.	i.	No batch failed during last three year of the quoted item from any Statutory lab.	5	5	The firm will provide undertaking in this Regard. The purchaser reserves the right to	

		ii.	No Batch failed during last two year of the quoted item from any Statutory lab.	3		verify the claim.																			
5	Financial status of Bidders	<table><tr><td>i</td><td>Last year audited balance sheet</td><td>03</td></tr><tr><td>ii</td><td>Tax returned (last 03 year)</td><td>02</td></tr></table>			i	Last year audited balance sheet	03	ii	Tax returned (last 03 year)	02	05	Acknowledgement of Tax Return must be attached.													
i	Last year audited balance sheet	03																							
ii	Tax returned (last 03 year)	02																							
6	Technical Staff	<table><tr><td rowspan="2">i</td><td rowspan="2">Regional Manager / Head of Concerned Department</td><td>Graduation in concerned field/B. pharm/ pharm. D</td><td>2</td></tr><tr><td>Post-graduation in concerned field</td><td>4</td></tr><tr><td rowspan="2">ii</td><td rowspan="2">Institutional Manager</td><td>Graduation in concerned field/B. pharm/ pharm. D</td><td>2</td></tr><tr><td>Post-graduation in concerned field</td><td>3</td></tr><tr><td rowspan="2">iii</td><td rowspan="2">Territory Managers / Quality Assurance Manager</td><td>Graduation in concerned field/B. pharm/ pharm. D</td><td>2</td></tr><tr><td>Post-graduation in concerned field</td><td>3</td></tr></table>			i	Regional Manager / Head of Concerned Department	Graduation in concerned field/B. pharm/ pharm. D	2	Post-graduation in concerned field	4	ii	Institutional Manager	Graduation in concerned field/B. pharm/ pharm. D	2	Post-graduation in concerned field	3	iii	Territory Managers / Quality Assurance Manager	Graduation in concerned field/B. pharm/ pharm. D	2	Post-graduation in concerned field	3	10	The bidder is required to attach attested copies of the relevant degrees and appointment letters of concerned technical staff. (Bank salary transaction statement of concerned staff)	
i	Regional Manager / Head of Concerned Department	Graduation in concerned field/B. pharm/ pharm. D	2																						
		Post-graduation in concerned field	4																						
ii	Institutional Manager	Graduation in concerned field/B. pharm/ pharm. D	2																						
		Post-graduation in concerned field	3																						
iii	Territory Managers / Quality Assurance Manager	Graduation in concerned field/B. pharm/ pharm. D	2																						
		Post-graduation in concerned field	3																						
	Total Marks				55																				

Part –B

Minimum Qualifying Marks = 65% of Total Marks = 35.75

PART C

- Satisfactory performance report by Government Teaching Hospitals of the quoted product is the prerequisite of Part-C
- Submission of the sample is mandatory.

EVALUATION OF SAMPLES AS PER ADVERTISED SPECIFICATON

BID ENQUIRY NO.	NAME OF ITEMS(Advertised)	Offered Specification	OFFERED BRAND NAME	MANUFACTURER / COUNTRY OF ORIGIN	COUNTING UNIT	SAMPLE STATUS	REMARKS (RESPONSIVE / NON RESPONSIVE WITH VALID REASON)

Recommendation for part (C) _____

Overall recommendation _____

EVALUATION CRITERIA FOR MEDICAL DEVICES/ LAB KITS REAGENTS & EQUIPMENT PLACEMENT BASIS.

Part-A COMPULSORY PARAMETERS:

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

Part =A Compulsory Parameters for all Categories

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

- Valid Computerized National Identity Card
- N. T. N. Certificate
- G. S. T. Reg. Certificate (where applicable)
- Bid Security (Estimated price as mentioned in tender documents, indicating the number, date and Bank (Copy must be attached with the technical bid).
- Under Taking Certificate Regarding Black Listing on the legal stamp paper of Rs.100/- or more.
- Bid Validity Period of 180 days
- Signed Terms & Conditions / Bidding Documents
- Authorization Letter from Manufacturer/Sole Agent in case of Sole Distributor
- Tender Sale Receipt.

Part-B

ORDINARY PARAMETERS:

The bid complying with compulsory parameter shall be evaluated for below mentioned parameters:

Sr. #.	Parameters	Detail			Total Marks	Remarks	MARKS OBTAINED
1	Past Performance of the Bidder (Last two years)	Major institutions (Government / Semi-Government) served:			10	The Claim requires documentation (Purchase Orders, Receipt Certificates & Delivery Challans etc.) of the institution(s).	
		i.	1	2			
		ii.	2 to 3	4			
		iii.	4 to 5	6			
		iv.	6 to 7	8			
		v.	8 & above	10			
2	Market / institutional Experience of quoted Product.	i.	Market Availability of quoted product in leading Chain Stores/ Pharmacies / institutions for last 02 years	7	15	Market availability in leading Chain Stores, Pharmacies of quoted item will be calculated from the date of commercial invoice. The firm will attach purchase orders / delivery Challan of the quoted item of any Government/Semi-Government institution.	
		ii	More than 02 up to 04 years	10			
		iii	More than 04 years	15			
3	Credibility & Certification of Manufacturer	i.	WHO / US FDA / CE certification / WHO Prequalification / Prequalification by Provincial or Federal Institutes.	7	10	Valid copies of certificates/letters Required.	
		ii.	Valid ISO certification. (Notarized ISO) /international reputed certificate.	3			
4	Batch	i.	No batch failed during	5	5	The firm will provide	

	quality For Last Three Years.		last three year of the quoted item from any Statutory lab.			undertaking in this Regard. The purchaser reserves the right to verify the claim.																			
		ii.	No Batch failed during last two year of the quoted item from any Statutory lab.	3																					
5	Financial status of Bidders	<table><tr><td>i</td><td>Last year audited balance sheet</td><td>03</td></tr><tr><td>ii</td><td>Tax returned (last 03 year)</td><td>02</td></tr></table>			i	Last year audited balance sheet	03	ii	Tax returned (last 03 year)	02	05	Acknowledgement of Tax Return must be attached.													
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6	Technical Staff	<table><tr><td rowspan="2">i</td><td rowspan="2">Regional Manager / Head of Concerned Department</td><td>Graduation in concerned field/B. pharm/ pharm. D</td><td>2</td></tr><tr><td>Post-graduation in concerned field</td><td>4</td></tr><tr><td rowspan="2">ii</td><td rowspan="2">Institutional Manager</td><td>Graduation in concerned field/B. pharm/ pharm. D</td><td>2</td></tr><tr><td>Post-graduation in concerned field</td><td>3</td></tr><tr><td rowspan="2">iii</td><td rowspan="2">Territory Managers / Quality Assurance Manager</td><td>Graduation in concerned field/B. pharm/ pharm. D</td><td>2</td></tr><tr><td>Post-graduation in concerned field</td><td>3</td></tr></table>			i	Regional Manager / Head of Concerned Department	Graduation in concerned field/B. pharm/ pharm. D	2	Post-graduation in concerned field	4	ii	Institutional Manager	Graduation in concerned field/B. pharm/ pharm. D	2	Post-graduation in concerned field	3	iii	Territory Managers / Quality Assurance Manager	Graduation in concerned field/B. pharm/ pharm. D	2	Post-graduation in concerned field	3	10	The bidder is required to attach attested copies of the relevant degrees and appointment letters of concerned technical staff. (Bank salary transaction statement of concerned staff)	
i	Regional Manager / Head of Concerned Department	Graduation in concerned field/B. pharm/ pharm. D	2																						
		Post-graduation in concerned field	4																						
ii	Institutional Manager	Graduation in concerned field/B. pharm/ pharm. D	2																						
		Post-graduation in concerned field	3																						
iii	Territory Managers / Quality Assurance Manager	Graduation in concerned field/B. pharm/ pharm. D	2																						
		Post-graduation in concerned field	3																						
	Total Marks				55																				

Minimum Qualifying Marks = 65% of Total Marks = 35.75

PART-C

EVALUATION AS PER ADVERTISED SPECIFICATON

BID ENQUIRY NO.	NAME OF ITEMS/ Tests (Advertised)	Offered Specification	OFFERED BRAND NAME	MANUFACTURER / COUNTRY OF ORIGIN	COUNTING UNIT	SAMPLE STATUS	REMARKS (RESPONSIVE / NON RESPONSIVE WITH VALID REASON)

Recommendation for part (C) _____

Overall recommendation _____

EVALUATION CRITERIA FOR B.M.E ITEMS

PART =A COMPULSORY PARAMETERS FOR B.M.E. ITEMS

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

Sr.No.	Parameter	Status
1.	Attested Copy of Computerized National Identity Card (CNIC) of authorized bidding signatory person of the bidder.	
2.	N. T. N. Certificate	
3.	G. S. T. Reg. Certificate (where applicable)	
4.	Bid Security (Estimated price as mentioned in tender documents, indicating the number, date and Bank (Copy must be attached with the technical bid).	
5.	Under Taking Certificate Regarding Black Listing on the legal stamp paper of Rs.100/- or more.	
6.	Bid Validity Period of 180 days	
7.	Signed Terms & Conditions / Bidding Documents	
8.	Original Tender Purchase Receipt	
9.	Intuitional Performance	

PART B EVALUATION CRITERIA (B.M.E. ITEMS.)

MARKING CRITERIA

Sr. No.	Parameters	Detail	Total Marks	Remarks												
1-	Performance of Last three years of the item being quoted(attach relevant documents)	<div>Major institutions Served Past Performance</div> <table><tr><td>i.</td><td>No institution served</td><td>0</td></tr><tr><td>Ii.</td><td>Institution served 1 to 4</td><td>5</td></tr><tr><td>Iii.</td><td>Institution served 5 to 9</td><td>10</td></tr><tr><td>Iv.</td><td>Institution served 10 or above</td><td>15</td></tr></table>	i.	No institution served	0	Ii.	Institution served 1 to 4	5	Iii.	Institution served 5 to 9	10	Iv.	Institution served 10 or above	15	15	The claims require documentation purchase order, receipt certificates, delivery challans, etc. from concerned institution.
i.	No institution served	0														
Ii.	Institution served 1 to 4	5														
Iii.	Institution served 5 to 9	10														
Iv.	Institution served 10 or above	15														
2-	Market experience of quoted products (attach supporting documents as proof)	<table><tr><td>i.</td><td>02 years</td><td>5</td></tr><tr><td>ii.</td><td>More than 02 up to 04 years</td><td>10</td></tr><tr><td>iii.</td><td>More than 04 years</td><td>15</td></tr></table>	i.	02 years	5	ii.	More than 02 up to 04 years	10	iii.	More than 04 years	15	15	Less than 2 year experience is in eligible.			
i.	02 years	5														
ii.	More than 02 up to 04 years	10														
iii.	More than 04 years	15														
3-	Compliance of Quality	<table><tr><td>i.</td><td>FDA / CE certification</td><td>10</td></tr><tr><td>ii.</td><td>Valid ISO Certificate</td><td>10</td></tr></table>	i.	FDA / CE certification	10	ii.	Valid ISO Certificate	10	20	Attach valid Certificates						
i.	FDA / CE certification	10														
ii.	Valid ISO Certificate	10														
4-	Financial Status	<table><tr><td>i.</td><td>10 Million or above</td><td>10</td></tr><tr><td>ii.</td><td>5 Million or above</td><td>05</td></tr><tr><td>iii.</td><td>Below 05 Million</td><td>02</td></tr></table>	i.	10 Million or above	10	ii.	5 Million or above	05	iii.	Below 05 Million	02	10	FBR tax returns showing sale of last financial year is required.			
i.	10 Million or above	10														
ii.	5 Million or above	05														
iii.	Below 05 Million	02														
5-	Valid letter of Authorization from Principal/manufacturer	<table><tr><td>i.</td><td>Sole Distributor certificate</td><td>10</td></tr></table>	i.	Sole Distributor certificate	10	10	Attach valid certificates									
i.	Sole Distributor certificate	10														

6-	Company Profile				10	Attach the attested copies of their CVs,their valid PEC No., attested set of relevant degrees along with their appointment letter and salary certificates.
		i.	B.Sc / B-Technical Engineers 4 or more	10		
		ii.	DAE Technical Engineers 4 or more	05		
7-	Registration, Tax and Audit Certificate				20	
		i.	Tax Return Last 3-years	10		
		ii.	Audit Report Last Three Years	10		
Total Marks					100	

Total marks: 100

Qualifying marks: 65% (65) and above

PART C

EVALUATION OF SAMPLES AS PER ADVERTISED SPECIFICATION

BID ENQUIRY NO.	NAME OF ITEMS	BRAND NAME	COUNTING UNIT	SAMPLE STATUS	REMARKS

Recommendation of par (C) _____

Over all Recommendation with justification _____

Category:- DENTAL MATERIAL ITEMS

EVALUATION CRITERIA

FOR DENTAL MATERIAL ITEMS,

Part-A COMPULSORY PARAMETERS:

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

Part =A COMPULSORY PARAMETERS FOR DENTAL MATERIAL ITEMS

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

Sr.No.	Parameter	Status
i.	Computerized National Identity Card	
ii.	N. T. N. Certificate	

iii.	G. S. T. Reg. Certificate (where applicable)	
iv.	Bid Security (Estimated price as mentioned in tender documents, indicating the number, date and Bank (Copy must be attached with the technical bid).	
v.	Under Taking Certificate Regarding Black Listing on the legal stamp paper of Rs.100/- or more.	
vi.	Bid Validity Period of 180 days	
vii.	Signed Terms & Conditions / Bidding Documents	
viii.	Authorization Letter from Manufacturer/Sole Agent in case of Sole Distributor) if applicable.	
ix.	Tender Sale Receipt	

EVALUATION CRITERIA (DENTAL MATERIAL ITEMS)

MARKING CRITERIA PART –B

Sr. No.	Parameters	Detail		Total Marks	Remarks									
1	Past Performance	Major institutions served, Past performance, contract execution: <table><tr><td>i</td><td>1</td><td>5</td></tr><tr><td>ii</td><td>2 to 3</td><td>15</td></tr><tr><td>iii</td><td>4 and above</td><td>20</td></tr></table>		i	1	5	ii	2 to 3	15	iii	4 and above	20	20	The claim requires documentation) Purchase Orders, Receipt Certificates & Delivery Challans, etc.) from the concerned institution.
i	1	5												
ii	2 to 3	15												
iii	4 and above	20												
2	Market / Institution experience of quoted product.	<table><tr><td>i</td><td>Market availability of quoted item in dental Store for last 01 year</td><td>10</td></tr><tr><td>ii</td><td>1 -2 years institution experience</td><td>110</td></tr></table>		i	Market availability of quoted item in dental Store for last 01 year	10	ii	1 -2 years institution experience	110	20	The market availability of quoted item will be calculated from the date of commercial invoice for parameters (i) the product having less than one year experience is ineligible and market availability of quoted items relates to availability in open market other than dental stores. Items experience shall be confirmed from 1 st market launch of the product with documentary proof / institution.			
i	Market availability of quoted item in dental Store for last 01 year	10												
ii	1 -2 years institution experience	110												
3	Compliance of Quality Standards	<table><tr><td>i</td><td>FDA/WHO approved</td><td>20</td></tr><tr><td>ii</td><td>Others</td><td>10</td></tr></table>		i	FDA/WHO approved	20	ii	Others	10	20	Valid copies of certificates / letters required.			
i	FDA/WHO approved	20												
ii	Others	10												

4	Financial status of Bidders	<table><tr><td>i</td><td>1 Million or above</td><td>20</td></tr><tr><td>ii</td><td>0.5 Million or above</td><td>10</td></tr></table>	i	1 Million or above	20	ii	0.5 Million or above	10	20	FBR Tax Return showing sale of last financial year is required.
i	1 Million or above	20								
ii	0.5 Million or above	10								
5	Contract Execution	<table><tr><td>i</td><td>Supply orders executed in two Tertiary Care Hospitals Punjab including Mayo Hospital Lahore within due delivery period</td><td>10</td></tr><tr><td>ii</td><td>Supply order executed in Tertiary Care Hospitals Punjab</td><td>05</td></tr></table>	i	Supply orders executed in two Tertiary Care Hospitals Punjab including Mayo Hospital Lahore within due delivery period	10	ii	Supply order executed in Tertiary Care Hospitals Punjab	05	10	The bidder is required to attach contract execution certificate from concerned institution
i	Supply orders executed in two Tertiary Care Hospitals Punjab including Mayo Hospital Lahore within due delivery period	10								
ii	Supply order executed in Tertiary Care Hospitals Punjab	05								
6	Technical Staff	<table><tr><td>i</td><td>Metric or equivalent in Any field</td><td>10</td></tr></table>	i	Metric or equivalent in Any field	10	10	The bidder is required to attach attested copies of the relevant degrees and appointment letters of concerned technical staff.			
i	Metric or equivalent in Any field	10								

Total marks: 100

Qualifying marks: 65% (65) and above

PART C

EVALUATION OF SAMPLES AS PER ADVERTISED SPECIFICATON

BID ENQUIRY NO.	NAME OF ITEMS	BRAND NAME	COUNTING UNIT	SAMPLE STATUS	REMARKS

Recommendation of par (C) _____

Over all Recommendation with justification _____

Part =A Compulsory Parameters for CT-Scan & X-Ray Films and its chemicals

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

- Computerized National Identity Card
- N. T. N. Certificate
- G. S. T. Reg. Certificate (where applicable)
- Bid Security (Estimated price as mentioned in tender documents, indicating the number, date and Bank (Copy must be attached with the technical bid).
- Under Taking Certificate Regarding Black Listing on the legal stamp paper of Rs.100/- or more.
- Bid Validity Period of 180 days
- Signed Terms & Conditions / Bidding Documents
- Authorization Letter from Manufacturer/Sole Agent in case of Sole Distributor) if applicable.
- Tender Sale Receipt

PART B= EVALUATION CRITERIA (X-RAY / CX.T.SCAN FILMS & ITS CHEMICALS, ITEMS)

MARKING CRITERIA

Sr. No.	Parameters	Detail			Total Marks	Remarks															
1	Past Performance (Last two years) As per Bid Form 4	Major institutions served, Past performance, contract execution: <table><tr><td>i</td><td>1</td><td>4</td></tr><tr><td>ii</td><td>2 to 3</td><td>8</td></tr><tr><td>iii</td><td>4 to 5</td><td>12</td></tr><tr><td>iv</td><td>6 to 7</td><td>16</td></tr><tr><td>v</td><td>8 and above</td><td>20</td></tr></table>			i	1	4	ii	2 to 3	8	iii	4 to 5	12	iv	6 to 7	16	v	8 and above	20	20	The claim requires documentation) Purchase Orders, Receipt Certificates & Delivery Challans, etc.) from the concerned institution.
i	1	4																			
ii	2 to 3	8																			
iii	4 to 5	12																			
iv	6 to 7	16																			
v	8 and above	20																			
2	Market / Institution experience of quoted product.	<table><tr><td>i.</td><td>Market availability of quoted product in leading chain stores / Pharmacies / Institutions from 02 years</td><td>7</td></tr><tr><td>ii.</td><td>More than 02 up to 04 years</td><td>10</td></tr><tr><td>iii.</td><td>More than 04 years</td><td>15</td></tr></table>			i.	Market availability of quoted product in leading chain stores / Pharmacies / Institutions from 02 years	7	ii.	More than 02 up to 04 years	10	iii.	More than 04 years	15	15	The market experience will not go beyond the date of registration (for registered items). less than Two year experience is ineligible and market availability of quoted items relates to availability in open market. Items experience shall be confirmed from 1 st market launch of the product with documentary proof / institution.						
i.	Market availability of quoted product in leading chain stores / Pharmacies / Institutions from 02 years	7																			
ii.	More than 02 up to 04 years	10																			
iii.	More than 04 years	15																			
3	Compliance of Quality Standards	<table><tr><td>i</td><td>FDA/WHO approved</td><td>20</td></tr><tr><td>ii</td><td>Others</td><td>10</td></tr></table>			i	FDA/WHO approved	20	ii	Others	10	20	Valid copies of certificates / letters required.									
i	FDA/WHO approved	20																			
ii	Others	10																			
4	Financial status of Bidders	<table><tr><td>i</td><td>2 Million or above</td><td>20</td></tr><tr><td>ii</td><td>1 Million or above</td><td>10</td></tr><tr><td>Iii</td><td>0.5 Million or above</td><td>05</td></tr></table>			i	2 Million or above	20	ii	1 Million or above	10	Iii	0.5 Million or above	05	20	FBR Tax Return showing sale of last financial year is required.						
i	2 Million or above	20																			
ii	1 Million or above	10																			
Iii	0.5 Million or above	05																			
5	Contract Execution	<table><tr><td>i</td><td>Supply orders executed in two Tertiary Care Hospitals Punjab including Mayo Hospital Lahore within due delivery period</td><td>10</td></tr><tr><td>ii</td><td>Supply order executed in one Tertiary Care Hospitals Punjab</td><td>05</td></tr></table>			i	Supply orders executed in two Tertiary Care Hospitals Punjab including Mayo Hospital Lahore within due delivery period	10	ii	Supply order executed in one Tertiary Care Hospitals Punjab	05	10	The bidder is required to attach contract execution certificate from concerned institution									
i	Supply orders executed in two Tertiary Care Hospitals Punjab including Mayo Hospital Lahore within due delivery period	10																			
ii	Supply order executed in one Tertiary Care Hospitals Punjab	05																			

Total marks: 85

Qualifying marks: 65% (55.25) and above

PART C EVALUATION OF SAMPLES AS PER ADVERTISED SPECIFICATION

BID ENQUIRY NO.	NAME OF ITEMS	BRAND NAME	MANUFACTURER / COUNTRY OF ORIGIN	ACCOUNTING UNIT	SAMPLE STATUS	REMARKS (RESPONSIVE / NON RESPONSIVE WITH VALID REASON)

Recommendation of part (C) _____

Overall recommendation _____

TECHNICAL EVALUATION PERFORMAS FOR PROCUREMENT OF GENERAL STORES, ELECTRIC STORE, STATIONERY STORE, SANITATION STORE, M.I.R WORKSHOP, BEDDING CLOTHING & LINEN STORE, ORTHO RAW MATERIAL ITEMS.

Part =A Compulsory Parameters for all Categories

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

- Attested Copy of Computerized National Identity Card (CNIC) of authorized bidding signatory person of the bidder
- N. T. N. Certificate
- G. S. T. Reg. Certificate (where applicable)
- Bid Security (Estimated price as mentioned in tender documents, indicating the number, date and Bank (Copy must be attached with the technical bid).
- Under Taking Certificate Regarding Black Listing on the legal stamp paper of Rs.100/- or more.
- Bid Validity Period of 180 days.
- Signed Terms & Conditions / Bidding Documents
- Original Tender Sale Receipt
- Institutional performance of bidder reference to quoted product from any Government institute.

Part =B Technical Evaluation Parameters

S. No.	Parameters	Total Marks	Marks Obtained
1	Company Profile: i. Storage Capacity (10) ii. other facilities i.e vehicles (05) iii. Human resource (05)	20	
2	Relevant Experience: (Government, Semi Government) i. More than 05 Institutes (20) ii. 02- 05 Institutes (10) iii. Less Than 02 (05)	20	

3	Certificate of satisfactory past performance issued by Competent Authority of relevant procuring agency for each year. For last 02 years		5	20
	More than 02 up to 04 years		10	
	More than 04 years		20	
4	Financial Status / Soundness:			20
	i. Turn over i.e. Bank Account or through Bank Certificate			
	50 Million or above		(20)	
	30 Million or above		(10)	
	10 Million or above		(05)	
	ii. Tax Paid for the last			
	Last 3 Years =		(10)	
	Last 2 Years =		(05)	
	iii. Audit Report			
	Company Audit Report			
Last 3 Years =		(10)	10	
Last 2 Years =		(05)		
Total Marks				100

Total marks: 100

Qualifying marks: 65% (65) and above

PART C EVALUATION OF SAMPLES AS PER ADVERTISED SPECIFICATON

BID ENQUIRY NO.	NAME OF ITEMS	BRAND NAME	ACCOUNTING UNIT	SAMPLE STATUS	REMARKS (RESPONSIVE / NON RESPONSIVE WITH VALID REASON)

Recommendation of part (C) _____

Overall recommendation _____

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:

☐: All Items mentioned in the Schedule of Requirements.

☐: Selected Items from the Schedule of Requirements¹.

List of Selected Items: (In case the Bidder has opted to bid for Selected Items, please type the Serial No². and the name of the Items selected for Bidding. Use additional Sheets if Required)

Sr. No.	Name of the Item

Signed:

Dated:

Official Stamp:

Attachment³: ☐ Original receipt for the purchase of the bidding documents.

BID FORM 1

Letter of Intention

Bid Ref No.

Date of the Opening of Bids

Name of the Contract : { Add name e.g. Supply of Drugs and Medicines etc }

To: [Name and address of Procuring Agency]

Dear Sir,

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

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

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

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 this [insert: number] day of [insert: month], [insert: year].

Signed:

BID FORM 2

AFFIDAVIT

I/We, the undersigned solemnly state that:

- 1) I/We have read the contents of the Bidding Document 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 department from any Government, or its organization or project.
- 8) The undersigned is ready to all by the charges sample(s)(1 or more depending upon the ordered quantity amounting to Rs.10 million or more) tested by any international WHO accredited Laboratory , collected by Department's Inspection Committee which will be paid directly to the International Lab and will accept the results.
- 9) I/We further undertake that I /we will ready to pay the standard charges of testing samples by DTLs Punjab alongwith sample cost. The procuring agency reserved the rights to send the samples of Medical Devices to verify the claims of sterilization, under such circumstances the DTL fee; sample cost etc should be the responsibility of the bidder.
- 10) I/We further under take to provide the Batch Release Laboratory Test Reports of each batch of the product on its delivery.
- 11) The price offered to Mayo hospital is not more than any institution.
- 12) In case offered product is not consumed with in prescribed shelf life, I / we undertake to replace the same without any extra charges.
- 13) Incase product is declared spurious, adulterated, counterfeit, misbranded or substandard; I /we undertake to provide fresh stock without any extra charges.
- 14) I / we undertake to adhere with the polices of the hospital / Government / DRAP for disposal of such product (mentioned above).

15) I / we undertake that prices coated by us are not more than the prices charge by us from any other procuring agency. In case of price difference I have no objection for such deduction.

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

Signed _____

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)	Checklist ⁴ (To be initialed by the Bidder against each document)	Relevant Page Number ⁵ in the Bid (To be filled by the Bidder)	Supporting Documents (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
Letter of Manufacturer's authorization			
Partnership Deed (where applicable)			
NTN Certificate			
GST Certificate			
Letter of Intention			
Affidavit			
One year experience evidence			
Original Receipt of purchase of Bidding Documents			

BID FORM 3(B)

MANUFACTURER'S AUTHORISATION⁶

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 15 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:-----

⁴ 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.

⁵ Bidders are required to mention the exact page number of relevant document placed in the Bid.

⁶ 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 Performance⁷.

Name of the Firm:

Bid Reference No:

Date of opening of Bid:

Assessment Period: (One Year as per Evaluation Criteria)

Name of the Purchaser/Institution	Purchase Order No.	Description Of Order	Value of Order	Date of Completion	Purchaser's ⁸ Certificate

BID FORM 5

Offered Item (s) (Technical Bid)

User Note: This form is to be filled by the Bidder for each individual item and shall submit in envelope marked as Technical Proposal.

Name of the Firm:

Bid.Ref.No:

Date of opening of Bid.

T/E No.	Name of the Item	Offered Specifications with make / manufacturer, country of origin etc.	Brand Name	Pack Size
1				
2				
3				

Signature: -----

Designation: -----

Date: -----

Official Stamp: -----

BID FORM 6

⁷ Bidders may use additional Sheets if required.

⁸ All certificates are to be attached with this form.

Price Schedule
(Financial Bid)

User Note: This form is to be filled by the Bidder for each individual item and shall submit in envelope marked as Financial Proposal.

Name of the Firm:

Bid.Ref.No:

Date of opening of Bid.

T/E No.	Name of the Item	Offered Specifications with make / manufacturer, country of origin etc.	Brand Name	Pack Size	Trade Prize	Retail Price	Offered Unit Price (Inclusive of all taxes)	Total price in figure	Total price in words
1									
2									
3									

Signature: -----

Designation: -----

Date: -----

Official Stamp: -----

Special Conditions of the Contract

AGREEMENT

THIS CONTRACT is made at _____ on _____ day of _____ 20____, between The Chief Executive Officer Mayo Hospital, Lahore (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 Supplier/ authorized Agent 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;

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. The Schedule of Requirements.
 - b. The Technical Specifications.
 - c. The Price Schedule submitted by the Bidder.
 - d. The Purchaser’s Notification of Award.
 - e. The Purchase Order
 - f. The General Conditions of Contract
 - g. The acceptance of bid.
2. **Interpretation:** In this Contract words and expressions shall have the same meanings as are respectively assigned to them in the General Conditions of this Contract hereinafter referred to as “Contract”:
3. **The Term of the Contract:** This contract shall remain valid for one year from the date of issuance of advance acceptance / notification of award.
4. The Supplier declares as under:
 - i. [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 Government of Punjab through any corrupt business practice.
 - ii. Without limiting the generality of the foregoing, [the Seller/ 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 Mayo Hospital, Lahore, except that which has been expressly declared pursuant hereto.
 - iii. [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 Mayo Hospital, Lahore and has not taken any action or shall not take any action to circumvent the above declaration, representation or warranty.
 - iv. [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 voidable at the option of Procuring Agency.
- v. 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 times 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.
 - vi. In case of any dispute concerning the interpretation and/or application of this Contract shall be settled through arbitration. The Secretary Health 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.

5. Items to be Supplied & Agreed Unit Cost: (i) 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).

(ii) Each Item 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

(iii) 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.

6. Payments: The Purchaser hereby covenants to pay the Supplier in consideration of the 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.

7. Mode of Payment: All payments to the Supplier shall be made through Crossed Cheques issued in the name of [supplier's name]

8. Payment Schedule: All payments to the Supplier shall be made in accordance with the agreed 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.

9. Performance Guarantee: (i) The Supplier shall within 10 days of issuance of advance acceptance, shall provide to the Purchaser a Performance Guarantee equivalent to 5% of the total Contract amount in the shape of Bank Guarantee / CDR. This Performance Guarantee shall be released to the Supplier upon successful completion of the Contract.

ii) Supplier's Bid Security already submitted with the Bid shall only be released upon satisfactory submission of a Performance Guarantee in accordance with sub-clause (i) above.

iii) Failure to submit a Performance Guarantee shall result into forfeiture of Bid Security and Cancellation of Contract / withdrawal of advance acceptance.

10. Penalties/ Liquidated Damages.

- i) Wherein the Supplier fails to make deliveries as per 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.
- ii) 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.
- iii) If the Supplier fails to supply the whole consignment, the entire amount of Performance Guaranty/ Security shall be forfeited to the Hospital account and the firm shall be blacklisted minimum for two years for future participation.
- iv) The exact time frame for making supplies with and without penalty shall be indicated in subsequent

Purchase orders.

- v) 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 the cost of late delivered supply shall be imposed upon the Supplier.
- vi) If Drug Testing is failed due to Misbranding, 25% Penalty of all contract amounts will be deducted and action will be taken against the firm. (As per letter No SO(DCP)H/9-2/2020)
- vii) Risk Purchase of the same item declared as Misbranded will be implemented.
- viii) Process of black listing of such companies causing inconvenience may be initiated by procuring agency.

11. Notices: All notices and correspondences incidental to this contract shall be in English language and shall be addressed to:

For the Purchaser:

[insert: name of office]

[insert: name of officer]

[insert: postal address]

[insert: telephone number, indicate country and city code]

[insert: facsimile or cable number or e-mail address]

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.

Signed/ Sealed: For the Purchaser.

Signature: _____
Name: _____
Designation: _____

**Signed/ Sealed: For the Manufacturer/
Authorized Supplier/ Authorized Agent.**

Signature: _____
Name: _____
Designation: _____

Technical Specifications and Ancillary Services

a). Product Specifications.

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

b). Labeling and Packing

- i. The manufacturer shall follow the Drugs (Labelling and Packing) Rules 1986, framed under the Drugs Act, 1976. (Read with guideline issued by Government of the Punjab vide No.PQCB/PRW-G-01/2019, Dated 27th May, 2019)
- ii. 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 aluminum strip / blister pack. Expiry date must be printed on each aluminum strip / blister. The syrup should be supplied in glass / pet bottle with sealed caps.

c) **Additional instructions for packing**

- i. 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 Drug Act, 1976 on judicial paper.
- ii. The bidder shall supply the drugs/medicines 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 case of items supplied by the foreign manufacturer the mentioned condition may be relaxed by the Procuring Agency.

**“NOT FOR SALE” “PUNJAB GOVERNMENT/
MAYO HOSPITAL, LAHORE PROPERTY”**

d). **Shelf life**

- i. The shelf life must be up to 85% for the locally manufactured drugs and 75% for the imported drugs.
- ii. The lower limit of the shelf life must be up to 80% and 70% with imposition of 1% penalty charges of actual shortfall in shelf life below prescribed limit for locally manufactured and imported medicines respectively.
- iii. In case of vaccines & other biotechnical products, the stores with the shelf life up to 70% will be accepted without penalty charges and up to 60% with imposition of 1% penalty charges of actual shortfall in shelf life below prescribed limit”

e). **Testing/Verification Procedures**

- x. After delivery of drugs and medicines at the Purchaser’s premises, the Purchaser shall send the samples from each batch 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 from DTL concerned as per law. The cost of the lab tests shall be borne by the Supplier.
- xi. In case of substandard 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 specified time period to be intimated by the purchaser but not later than 21 days (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 Drugs Regulatory Authority for legal action as per Drugs Act 1976 and disposal of substandard stocks. The supplies found to contravene the section 23 of Drugs Act, 1976 will not be returned to the supplier till the decision of the Provincial Quality Control Board.
- xii. 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.
- xiii. Provided item (s) were required may be tested from concerned laboratory / institute etc and the cost of sample (s) and deposit fee shall be the responsibility of vender / supplier.

f). **Transportation/Delivery Requirements**

- i. 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
- ii. All costs associated with the transportation including loading/unloading of drugs and medicines and road taxes shall be borne by the Supplier.
- iii. The firm/ contractor shall be bound to receive / collect the warrantor portion of sample for onward submission through the authorized person dedicated for the financial year 2023-24 for Hospital supplies, correspondence etc, at their own cost and risk.
- iv. All cold chain (perishable) items must be delivered in a safe and proper manner, prescribed for such types of items.

PAYMENT SCHEDULE

(Payment to the Suppliers will be made against satisfactory performance and upon submission of required documents and in accordance with the procedure mentioned in the PPR-2014. However, if there is any alternate payment schedule, agreed by the Procuring Agency and Supplier, will be annexed here)

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 Supplier" means the individual or firm supplying the goods under this Contract.
 - (h) "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 Import** 3.1 All goods and related services to be supplied under the 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 Hospital 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 Documents and Information

- 5.1 The Supplier shall not, without the Purchaser's prior written consent, disclose the Contract, or any provision thereof, or any 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.
- 5.2 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.
- 5.3 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 Samples

- 7.1 Before commencing supplies, the Supplier shall provide 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 Storage Arrangements

- 8.1 To ensure storage arrangements for the intended supplies, the Supplier shall inform the Purchaser at least One (01) week in 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 Tests

- 9.1 The Purchaser or its representative shall have the right to 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.
- 9.3 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.
- 9.4 Nothing in GCC Clause 9 shall in any way release the Supplier from any warranty or other obligations under this Contract.

10. Delivery of Documents

- 10.1 The Supplier in accordance with the terms and manner 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 Duty Paid (DDP) 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 Services	13.1	The Supplier shall be required to provide the incidental 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 as per Drugs Act, 1976.
	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. 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 Amendments	17.1	No variation in or modification of the terms of the Contract shall 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 Supplier's Performance	20.1	Delivery of the goods shall be made by the Supplier in accordance with the time schedule prescribed by the Purchaser in the Schedule of Requirements.
	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 or without 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 Default	21.1	<p>The Purchaser, without prejudice to any other remedy for breach of Contract, by written notice of default sent to the Supplier, may terminate this Contract in whole or in part:</p> <p>(a) if the Supplier fails to deliver any or all installments of the goods within the period(s) specified in the Contract and subsequent purchase order, or within any extension thereof granted by the Purchaser pursuant to GCC Clause 20; or</p> <p>(b) if the Supplier fails to perform any other obligation(s) under the Contract.</p> <p>(c) if the Supplier, in the judgment of the Purchaser has engaged in corrupt or fraudulent practices in competing for or in executing the Contract.</p> <p>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 supplier or contractor in the procurement process or in contract execution to the detriment of the Procuring agencies; 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 agencies 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"</p>

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.
	22.2	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 purchase order under the Contract and inform the supplier of its findings promptly.
	22.3	Unless Purchaser informs the Supplier in writing of tis 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 Insolvency	23.1	The Purchaser may at any time terminate the Contract by giving written notice of one month time to the Supplier if the Supplier becomes bankrupt or otherwise insolvent. In this even, 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 Resolution of Disputes	24.1	The purchaser and the supplier shall make every effort to resolve amicably by direct informat negotiation any disagreement or dispute arising between them under or in connection with the Contract.
	24.2	If, after thirty (30) days from the commencement of such informal 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 / or application if this contract shall be settled through arbitration under the Arbitration Act of 1940 (As amended from time to time.
	24.4	Redressal of Grievances and Arbitration shall be in accordance with Rules 67 and 68 of PPRA, 2014 respectively. The Redressal grievance committee shall investigate and decide upon the complaint submitted by the bidder.
25. Governing Language	25.1	The Contract shall written in English language. Subject to 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 Law	26.1	This contract shall be governed by the Laws of Pakistan and the courts of Pakistan shall have exclusive jurisdiction.
27. Notices	27.1	Any Notice given by one party to the others 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 notices or on the notices, s effective date, whichever is later.

- 28. Taxation** 28.1 All taxation, whether International, Federal, Provincial or Local, shall be borne by the supplier.
- 29. Risk Purchase** 29.1 Risk purchases shall be proceeded as options:
- Option i- Risk purchase (on account of bulk procurement) shall be made from the descending lowest bidders where available /applicable.
- Option ii- Petty purchase through Sanction/quotation shall made as per financial limits, in case of failure against option i.
- Option iii- Local /Day to Day purchase shall be adopted for urgent requirements only till the availability in stores by above mentioned modes. Stores must be kept up to cope with the urgent needs of patients.
- If the first lowest contractor failed to supply the product in prescribed time of delivery, risk purchase will be made the extra amount will be deducted from the CDR/Bills/Performance Guarantee of bidder lying in this hospital.
- NOTE= Any matter not prescribed in the said bidding documents, or the detail off which is which is not present herein, shall be dealt as mentioned in Health Department standard Bidding Documents and governed under PPR ,2014 (amended) upto date.


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TENTATIVE ANNUAL DEMAND OF MEDICINES (R-01) FOR THE FINANCIAL YEAR 2023-24

T/E No	Name of Store	Quantity Required	Estimated Cost (per unit) LPR / Retail Price (Rs)	Total Cost (Rs)	Delivery Schedule	2% Bid Security
1	Inhalation Isoflurane 100ml (Pack in carton with leaflet)(with brand new Tech-five vaporizer with calibration certificate , back up services and key filler as per requirements of theatres)(Pack in carton with leaflet)	6000	3250	19500000	2	390000
2	Inhalation Sevoflurane 250ml (with brand new Tech-five vaporizer with calibration certificate, back up services, and key filler as per requirements of theatres)(Pack in carton with leaflet)	800	16400	13120000	2	262400
3	Inj 20% Albumin (Human) vial of 50ml (Pack in carton with leaflet)	12000	7117	85404000	3	1708080
10	Inj Heparin 5000IU/ml (Vial of 5ml) (Pack in carton with leaflet)	50000	887	44350000	2	887000
13	Inj Octreotide acetate 0.1mg/ml (Amp/Vial) (Pack in carton with leaflet) with thermolog data.	24000	260	6240000	2	124800
15	Inj Streptokinase 1.5 MIU (Vial) (Pack in carton with leaflet)	3000	5650	16950000	2	339000
17	Infusion 0.45% Sodium Chloride + 5% Dextrose (Bottle of 500 ml) (Pack in carton)	30000	39.65	1189500	2	23790
18	Infusion 0.9% Sodium Chloride (Bottle of 1000 ml) (Pack in carton)	400000	48	19200000	2	384000
19	Infusion 0.9% Sodium Chloride (Bottle of 500ml) (Pack in carton)	100000	36.21	3621000	2	72420
20	Infusion 0.9% Sodium Chloride (Bottle of 100ml) (Pack in carton)	25000	33	825000	2	16500
21	Infusion 10% Dextrose Water 1000ml (Pack in carton)	12000	63.22	758640	1	15172.8
22	Infusion 10% Dextrose Water 500ml (Pack in carton)	10000	43.1	431000	1	8620
23	Infusion 20% Fat Emulsion ,Vial of 250ml (Pack in carton)	4000	386.75	1547000	2	30940
24	Infusion 25% Dextrose water (Bottle of 1000ml) (Pack in carton)	4000	108	432000	1	8640
25	Infusion 5% Aminoacids (Bottle of 500ml) (Pack in carton)	12000	328	3936000	3	78720

26	Infusion 5% Dextrose Saline (Bottle of 1000ml) (Pack in carton)	30000	56.9	1707000	2	34140
27	Infusion 5% Dextrose Saline (Bottle of 500ml) (Pack in carton)	18000	24	432000	2	8640
28	Infusion 5% Dextrose water (Bottle of 1000ml) (Pack in carton)	40000	56.32	2252800	2	45056
29	Infusion 5% Dextrose water (Bottle of 500ml) (Pack in carton)	30000	39.65	1189500	2	23790
30	Infusion Dextrose 4.3% + Sodium Chloride 0.18% w/v (Bottle of 500ml) (1/5 NS) (Pack in carton)	12000	39.08	468960	2	9379.2
31	Infusion Hydroxy ethyl starch 3% (Bottle of 500ml) (Pack in carton)	15000	224.54	3368100	2	67362
32	Infusion Mannitol 20% (Bottle of 500ml) (Pack in carton)	20000	111.89	2237800	2	44756
33	Infusion Modified Fluid Gelatin 4% (Bottle of 500ml) (Pack in carton)	5000	396.59	1982950	1	39659
34	Infusion Polygeline 3.5% (Bottle of 500ml) (Pack in carton)	7000	334.87	2344090	1	46881.8
35	Infusion Ringer Lactate (Bottle of 1000ml) (Pack in carton)	150000	50.75	7612500	2	152250
36	Infusion Ringer Lactate (Bottle of 500ml) (Pack in carton)	80000	31	2480000	2	49600
37	Inj 25 % Dextrose water (Amp of 25ml or less) (Pack in carton with leaflet)	100000	5.5	550000	2	11000
38	Inj Potassium Chloride 7.46% or Less (Amp of 20ml) (Pack in carton with leaflet)	90000	8.45	760500	2	15210
39	Inj Sodium Bicarbonate 8.4% or Less (Pack in carton with leaflet)	60000	16.1	966000	2	19320
43	Inj Ketamine HCl 50mg/ml, Amp/Vial of 2ml (Pack in carton with leaflet)	10000	25	250000	1	5000
45	Inj Lignocaine HCl 2% (Amp of 2ml) (Pack in carton with leaflet)	200000	1.6	320000	1	6400
46	Inj Lignocaine 2% with adrenaline 0.001%, amp of 2ml (Pack in carton with leaflet)	60000	2.41	144600	1	2892
50	Inj Propofol 10mg/ml, amp of 20 ml (pack in carton with leaflet)	45000	370	16650000	2	333000
52	Gel Lignocaine 2%, 15grm (Sealed tube with applicator) (Pack in carton with leaflet)	100000	16.5	1650000	2	33000
53	Solution Lignocaine 4% 50ml (Pack in carton with leaflet)	500	42.75	21375	1	427.5
57	Inj Amphotericin B 50mg (Vial) (Pack in carton with leaflet)	2000	525.9	1051800	2	21036
58	Inj Artemether 80mg/ml (Amp of 1ml) (pack in carton with leaflet)	1000	19	19000	1	380

59	Inj Benzyl Penicillin 1000000 Units (Vial) (Pack in carton with leaflet)	200000	8.49	1698000	2	33960
61	Inj Cefotaxime Sodium 1G IV,IM (Vial) with water for injection (Pack in carton with leaflet)	60000	75	4500000	1	90000
62	Inj Ceftazidim 1G (Vial) with water for injection (Pack in carton with leaflet)	20000	121	2420000	1	48400
64	Inj Cephadrine 500mg (Vial) (Pack in carton with leaflet)	5000	35.95	179750	2	3595
65	Inj Ciprofloxacin 200mg/100ml (Vial) (Pack in carton with leaflet)	36000	84	3024000	2	60480
66	Inj Clarithromycin 500mg (Vial) with water for injection (pack in carton with leaflet)	2400	262.2	629280	1	12585.6
68	Inj Fluconazole 2mg/ml (Vial of 50ml) (Pack in carton with leaflet)	3000	297	891000	1	17820
69	Inj Gentamycin Sulphate 80mg/2ml, amp of 2ml (Pack in carton with leaflet)	15000	17.47	262050	1	5241
70	Inj Moxifloxacin 400mg/250ml (Vial) (Pack in carton with leaflet)	120000	127.55	15306000	3	306120
71	Inj Streptomycin 1gm (Vial) (Pack in carton with leaflet)	10000	6	60000	1	1200
76	Inj. Linezolid 600mg Vial of 300ml (Pack in carton with leaflet)	24000	183.22	4397280	2	87945.6
77	Cap/Tab. Amoxycillin (as trihydrate) 500mg + Clavulanic (as potassium) Acid 125mg (Pack in carton with leaflet)	80000	16.69	1335200	2	26704
79	Cap/Tab. Cephadrine 500 mg (Blister Pack) (Pack in carton with leaflet)	5000	6.83	34150	1	683
80	Cap/Tab. Clindamycin 300mg (Blister Pack) (Pack in carton with leaflet)	8000	21.56	172480	1	3449.6
81	Cap/Tab. Doxycycline 100 mg (Blister Pack) (Pack in carton with leaflet)	10000	2.89	28900	1	578
82	Cap/Tab. Fluconazole 150mg (Blister Pack) (Pack in carton with leaflet)	6000	59	354000	2	7080
83	Cap/Tab. Itraconazole 100 mg (Blister Pack) (Pack in carton with leaflet)	5000	11.43	57150	1	1143
84	Cap/Tab. Rifampicin 450mg (Blister Pack) (Pack in carton with leaflet)	12000	6.75	81000	1	1620
85	Tab. Acyclovir 400mg (Blister Pack) (Pack in carton with leaflet)	12000	13	156000	1	3120
86	Tab. Artemether 80mg + Lumefantrine 480mg (Blister Pack) (Pack in carton with leaflet)	2400	21.17	50808	1	1016.16
90	Tab. Ethambutol 400mg (Blister Pack) (Pack in carton with leaflet)	20000	5	100000	1	2000

91	Tab. Ethambutol=275mg / 300mg, Rifampicin=150mg, Isoniazide=75mg (Blister Pack) (Pack in carton with leaflet) 1. Bioavailability certificate/ Bioequivalence studies by WHO recommended laboratory available on WHO web site must be attached. 2. The supplier will submit a certificate/affidavit that company is having GMP and has not changed source of raw material/active ingredients since the certificate of bioavailability/bioequivalence.	600000	6.25	3750000	2	75000
92	Tab. Ethambutol=275mg, Rifampicin=150mg, Isoniazide=75mg, Pyrazinamide =400mg (Blister Pack) (Pack in carton with leaflet) 1. Bioavailability certificate/ Bioequivalence studies by WHO recommended laboratory available on WHO web site must be attached. 2. The supplier will submit a certificate/affidavit that company is having GMP and has not changed source of raw material/active ingredients since the certificate of bioavailability/bioequivalence.	360000	9.8	3528000	2	70560
93	Tab. Isoniazid 100mg (Blister Pack) (Pack in carton with leaflet)	18000	0.35	6300	1	126
95	Tab. Linezolid 600mg (Blister Pack) (Pack in carton with leaflet)	6000	22.9	137400	1	2748
96	Tab. Metronidazole 400mg (Blister Pack) (Pack in carton with leaflet)	60000	1.63	97800	1	1956
98	Tab. Pyrazinamide 500mg (Blister Pack) (Pack in carton with leaflet)	20000	2.05	41000	1	820
99	Tab. Rifaximin 550mg (Blister Pack) (Pack in carton with leaflet)	20000	27	540000	2	10800
101	Susp. Amoxicillin 250mg (Bottle of 90ml or less) (Pack in carton with leaflet)	5000	86.9	434500	1	8690
102	Susp. Amoxycillin (as trihydrate) 250mg + Clavulanic (as potassium) Acid 62.50mg (Pack in carton with leaflet)	24000	107	2568000	2	51360
103	Susp. Artemether 15mg/5ml + Lumefantrine 90mg Bottle of 90ml or less (Pack in carton with leaflet)	1200	135	162000	1	3240
106	Susp/Syp. Clarithromycin 125mg/5ml (Pack with teaspoon in carton with leaflet)	12000	169	2028000	2	40560
107	Susp. Metronidazole 200mg/5ml (Bottle of 60ml) (Pack in carton with leaflet)	12000	20	240000	2	4800
108	Susp. Nystatin 100000 IU/ml with dropper Bottle of 30ml Individually packed in carton with dropper and leaflet	3000	57	171000	1	3420

109	Inj 5-Flourauracil 250mg (Pack in carton with leaflet)	24000	79	1896000	2	37920
110	Inj Bleomycin 15mg (Pack in carton with leaflet)	800	850	680000	1	13600
113	Inj Cyclophosphamide 500mg(Vial) (Pack in carton with leaflet)	6000	240	1440000	2	28800
114	Inj Cyterabine 500mg (Vial) with water for injection (Pack in carton with leaflet)	3600	579	2084400	2	41688
116	Inj Dactinomycin 0.5mg(Vial) (Pack in carton with leaflet)	240	240	57600	2	1152
117	Inj Daunorubicin 20mg (Vial) (Pack in carton with leaflet)	1800	484	871200	2	17424
120	Inj Doxorubicin 50mg (Vial) (Pack in carton with leaflet)	7200	1142	8222400	2	164448
121	Inj Etoposide 100mg/5ml(Vial of 5ml) (Pack in carton with leaflet)	2400	467	1120800	2	22416
123	Inj Folinic Acid 15mg (Vial) with water for injection (Pack in carton with leaflet)	20000	246	4920000	2	98400
126	Inj Goserelin 3.6mg (Vial) (Pack in carton with leaflet)	300	13669.74	4100922	2	82018.44
127	Inj Ifosphamide 1G (Vial) (Pack in carton with leaflet)	1200	430	516000	2	10320
129	Inj L-Asparaginase 10000 IU (Vial) (Pack in carton with leaflet)	300	1550	465000	2	9300
130	Inj MesNa 400mg with water for injection (Pack in carton with leaflet)	9000	38.81	349290	2	6985.8
131	Inj Methotrexate 500mg (Vial) with water for injection (Pack in carton with leaflet)	720	1400	1008000	2	20160
138	Inj Vinblastine Sulphate 10mg (Vial) (Pack in carton with solvent & leaflet)	120	340	40800	2	816
156	Tab. Tamoxifen 10mg (Blister Pack) (Pack in carton with leaflet)	6000	4.67	28020	1	560.4
157	Saccharomyces Boulardii 250mg/ Sachet (Pack in carton with leaflet)	5000	25	125000	1	2500
158	Inj Anti Diphteria Serum 10000iu (Pack in carton with leaflet)	1000	3430	3430000	2	68600
162	Inj Enoxaparin 40mg Prefilled Syringe (Pack in carton with leaflet)	20000	378.47	7569400	2	151388
163	Inj Enoxaparin 60mg Prefilled Syringe (Pack in carton with leaflet)	50000	509.87	25493500	2	509870
164	Inj Human Immunoglobulin (IgG, IgA, IgM) 0.5gm Vial in 10ml (Pack in carton with leaflet)	2000	15290	30580000	2	611600
165	Inj Human Immunoglobulin IgG 2.5gm 5% Vial in 50ml (Pack in carton with leaflet)	1000	15250	15250000	2	305000

166	Inj Human Immunoglobulin IgG intravenous 5% 0.5gm in 10ml (Pack in carton with leaflet)	5000	8640	43200000	2	864000
167	Inj Human Insulin 70/30 100 IU/ml, (vial of 10ml) with Bioequivalence studies and thermolog data (Pack in carton with leaflet)	36000	415	14940000	2	298800
168	Inj Human Insulin NPH 100 IU/ml, (vial of 10ml) with Bioequivalence studies and thermolog data (Pack in carton with leaflet)	4000	415	1660000	2	33200
169	Inj Human Insulin Regular 100 IU/ml, (vial of 10ml) with Bioequivalence studies and thermolog data (Pack in carton with leaflet)	8000	415	3320000	2	66400
170	Inj Somatropin 5mg (with thermolog data) (Pack in a carton with leaflet)	600	6844	4106400	2	82128
172	Inj Tetanus Immunoglobulin 250 IU (Human) (Pack in carton with leaflet)	2000	1921	3842000	2	76840
173	Inj Tetanus Toxoid 0.5ml (Amp of 0.5ml) (Pack in carton with leaflet)	120000	47.15	5658000	2	113160
174	Inj. Adenosin 3mg/ml (Pack in carton with leaflet)	500	1020	510000	2	10200
175	Inj. Adrenaline 0.1% w/v (1:1000) (Amp of 1ml) (Pack in carton with leaflet)	120000	5.08	609600	2	12192
176	Inj. Aminophylline 250mg/10ml (Amp of 10ml) (Pack in carton with leaflet)	2400	8.84	21216	1	424.32
178	Inj. Atropine 1mg/ml (Amp of 1ml) (Pack in carton with leaflet)	120000	3.3	396000	2	7920
180	Inj. Calcium Chloride 10ml (Pack in carton with leaflet)	1200	20	24000	1	480
181	Inj. Calcium Gluconate 10% (Amp of 10ml) (Pack in carton with leaflet)	120000	10	1200000	2	24000
183	Inj. Diazepam 10mg/2ml (Amp of 2ml) (Pack in carton with leaflet)	100000	48.46	4846000	2	96920
185	Inj. Digoxin 0.5mg/ml (Amp of 1ml) (Pack in carton with leaflet)	1000	10	10000	1	200
186	Inj. Dimenhydrinate 50mg/ml amp of 1ml (Pack in carton with leaflet)	10000	6.4	64000	1	1280
189	Inj. Drotaverine HCl 40mg/2ml , Ampoule of 2ml(Pack in carton with leaflet)	100000	8.3	830000	2	16600
190	Inj. Ephedrine 30mg (Pack in carton with leaflet)	3000	50	150000	2	3000
191	Inj. Flumazenil 1mg, Vial (Pack in carton with leaflet)	1000	4000	4000000	2	80000
194	Inj. Haloperidol (5mg/ml) (Amp of 1ml) (Pack in carton with leaflet)	5000	8.81	44050	1	881
195	Inj. Hydralazine 20mg (Pack in carton with leaflet)	5000	23.38	116900	1	2338

197	Inj. Iopromide 300 Epo 0.623gm 300mg I/ml (Vial of 50ml) (Pack in carton with leaflet) or Inj Iohexol 300mg/ml (Vial of 50ml) (Pack in carton with leaflet)	6000	1249	7494000	1	149880
198	Inj. Iopromide 370/100cc 768.86mg/ml Vial or Inj Iohexol 755 mg/ml eq to 350mg/ml (Pack in carton with leaflet)	20000	2125	42500000	2	850000
202	Inj. Labetalol 50mg/10ml (Amp of 10ml) (Pack in carton with leaflet)	10000	29.923	299230	2	5984.6
203	Inj. Levetriacetam 100mg/ml (Amp of 5ml) (Pack in carton with leaflet)	20000	85	1700000	2	34000
204	Inj. L-Ornithine L-Aspartate 10ml (Pack in a carton with leaflet)	2000	284	568000	2	11360
205	Inj. Magnesium sulphate 500mg/ml (Amp of 10ml) (Pack in carton with leaflet)	5000	20.01	100050	2	2001
206	Inj. Methyl Prednisolone Sodium Succinate 1G (Vial) (Pack in carton with leaflet)	6000	1229	7374000	2	147480
209	Inj. Nalbuphine HCl 10mg/ml (Amp of 1ml) (Pack in carton with leaflet)	240000	25.45	6108000	2	122160
214	Inj. Pheniramine Maleate 25mg/ml (Amp of 2ml) (Pack in carton with leaflet)	150000	1.59	238500	1	4770
215	Inj. Phenobarbitone 130mg/ml (Amp of 1ml) (Pack in carton with leaflet)	500	25	12500	1	250
217	Inj. Pralidoxime 200mg (Pack in carton with leaflet)	1500	196.41	294615	2	5892.3
218	Inj. Protamine Sulphate 10mg/ml (Amp of 5ml) (Pack in carton with leaflet)	2400	200	480000	1	9600
219	Inj. Sodium Amidotrizoate 0.1gm, Meglumin amidotrizoate 0.66gm, 370mg/ml (Amp of 20ml) (Pack in carton with leaflet)	1000	182.72	182720	1	3654.4
220	Inj. Thiamine 100mg, Pyridoxine 100mg, Cyanocobalamin 1000mcg/3ml, ampoule of 3ml (Pack in carton)	6000	3.1	18600	1	372
224	Inj. Triamcinolone 40mg (Pack in carton with leaflet)	2000	53.55	107100	2	2142
225	Inj. Valproate Sodium 100mg/ml (Amp of 5ml) (Pack in carton with leaflet)	40000	145.97	5838800	2	116776
226	Inj. Verapamil 2.5mg/ml (Amp of 2ml) (Pack in carton with leaflet)	600	19.6	11760	1	235.2
229	Cap/Tab. Gabapentin 100mg (Blister Pack) (Pack in carton with leaflet)	60000	3.49	209400	1	4188
230	Cap/Tab. Nifedipine 10mg (Blister Pack) (Pack in carton with leaflet)	5000	1.233	6165	1	123.3
233	Cap/Tab. Tramadol 50mg (Blister Pack) (Pack in carton with leaflet)	60000	5.89	353400	1	7068
235	Rotacap Budesonide and Formoterol 200/6 mcg (Pack in carton with leaflet)	12000	9.36	112320	1	2246.4
237	Tab. Acetazolamide 250mg (Blister Pack) (Pack in carton with leaflet)	4000	1.13	4520	1	90.4

238	Tab. Alprazolam 0.5mg (Blister Pack) (Pack in carton with leaflet)	15000	6.23	93450	1	1869
239	Tab. Aluminium Hydroxide 250mg + Magnesium Trisilicate 500mg (Blister Pack) (Pack in carton with leaflet)	50000	0.72	36000	1	720
241	Tab. Ascorbic Acid 500mg (Pack in carton)	3600	2.08	7488	1	149.76
242	Tab. Aspirin (Enteric coated) 75 mg (Blister Pack) (Pack in carton with leaflet)	100000	1.07	107000	1	2140
243	Tab. Aspirin 300mg (Blister Pack) (Pack in carton with leaflet)	40000	1.54	61600	1	1232
244	Tab. Atenolol 100mg (Blister Pack) (Pack in carton with leaflet)	16000	0.55	8800	1	176
246	Tab. Azathioprine 50mg (Blister Pack) (Pack in carton with leaflet)	5000	9	45000	1	900
247	Tab. Baclofen 10mg (Blister Pack) (Pack in carton with leaflet)	10000	4.16	41600	1	832
248	Tab. Bisoprolol 5mg (Blister Pack) (Pack in carton with leaflet)	20000	4	80000	1	1600
249	Tab. Bromazepam 3mg (Blister Pack) (Pack in carton with leaflet)	12000	2.5	30000	1	600
250	Tab. Calcium acetate containing 667 mg of calcium acetate equivalent to 169 mg of calcium (Blister Pack) (Pack in carton with leaflet)	12000	2.6	31200	1	624
251	Tab. Calcium Carbonate eq to Elemental Calcium 268mg or more + Vit D 125 IU or more/tab (Pack in carton with leaflet)	36000	14.9	536400	1	10728
252	Tab. Captopril 25 mg (Blister Pack) (Pack in carton with leaflet)	60000	5.0425	302550	1	6051
253	Tab. Carbamazepine 200 mg (Blister Pack) (Pack in carton with leaflet)	60000	3.6	216000	1	4320
256	Tab. Clonazepam 0.5mg (Blister Pack) (Pack in carton with leaflet)	20000	1.36	27200	1	544
260	Tab. Deferasirox 400mg (Pack in carton with leaflet)	36000	175	6300000	2	126000
261	Tab. Diazepam 5mg (Blister Pack) (Pack in carton with leaflet)	10000	2	20000	1	400
263	Tab. Digoxin 0.25mg (Pack in carton with leaflet)	6000	1.4	8400	1	168
264	Tab. Divalproex Sodium 500mg (Blister Pack) (Pack in carton with leaflet)	100000	9.71	971000	2	19420
265	Tab. Domperidone 10mg (Blister Pack) (Pack in carton with leaflet)	60000	0.98	58800	1	1176
266	Tab. Dosulepin 25mg (Blister Pack) (Pack in carton with leaflet)	2000	1.1704	2340.8	1	46.816
267	Tab. Drotaverine 40mg (Blister Pack) (Pack in carton with leaflet)	12000	4.57	54840	1	1096.8
269	Tab. Febuxostat 40 mg (Blister Pack) (Pack in carton with leaflet)	6000	4	24000	1	480

270	Tab. Ferrous Fumarate /Sulphate 150mg +Folic Acid 0.5mg	12000	5.06	60720	1	1214.4
271	Tab. Fexofenadine 120mg (Blister Pack) (Pack in carton with leaflet)	36000	6.2	223200	1	4464
272	Tab. Frusemide 40 mg (Blister Pack) (Pack in carton with leaflet)	24000	2.22	53280	1	1065.6
273	Tab. Glibenclamide 5mg (Blister Pack) (Pack in carton with leaflet)	6000	1.75	10500	1	210
274	Tab. Gliclazide MR 60mg (Blister Pack) (Pack in carton with leaflet)	30000	7.68	230400	1	4608
276	Tab. Glyceryl Trinitrate (Sub Lingual) 0.5mg (Blister Pack) (Pack in carton with leaflet)	18000	0.43	7740	1	154.8
277	Tab. Haloperidol 5mg (Blister Pack) (Pack in carton with leaflet)	5000	0.57	2850	1	57
278	Tab. Hyoscine-N- Butyl Bromide 10mg + Paracetamol 500mg (Blister Pack) (Pack in carton with leaflet)	6000	5.11	30660	1	613.2
280	Tab. Levodopa 250mg+ Carbidopa 25mg (Blister Pack) (Pack in carton with leaflet)	6000	5.78	34680	1	693.6
283	Tab. Losartan Potassium 50mg+Hydrochlorothiazide 12.5mg (Blister Pack) (Pack in carton with leaflet)	6000	4.24	25440	1	508.8
285	Tab. Mefenamic Acid 500mg (Blister Pack) (Pack in carton with leaflet)	100000	1.95	195000	1	3900
286	Tab. Metformin 500 mg (Blister Pack) (Pack in carton with leaflet)	50000	1.29	64500	1	1290
287	Tab. Metoclopramide 10mg (Blister Pack) (Pack in carton with leaflet)	20000	0.59	11800	1	236
288	Tab. Metoprolol 100mg (Blister Pack) (Pack in carton with leaflet)	20000	3.65	73000	1	1460
290	Tab. Multivitamin (Vit A, Vit D, Thiamine, Riboflavin, Nicotinamide) with minerals (Pack in carton)	30000	3.52	105600	1	2112
294	Tab. Potassium Chloride 500mg (Pack in carton with leaflet)	30000	0.5	15000	1	300
295	Tab. Prednisolone 5mg (Pack in carton)	240000	1.43	343200	1	6864
296	Tab. Procyclidine 5mg (Blister Pack) (Pack in carton with leaflet)	12000	0.65	7800	1	156
297	Tab. Propranolol 10mg (Blister Pack) (Pack in carton with leaflet)	10000	1.47	14700	1	294
298	Tab. Risperidone 2 mg (Blister Pack) (Pack in carton with leaflet)	3000	3.2	9600	1	192
300	Tab. Serratiopeptidase 10mg (Blister Pack) (Pack in carton with leaflet)	20000	7.5	150000	1	3000
301	Tab. Silymarin 200mg (Pack in carton with leaflet)	18000	7.8	140400	1	2808
302	Tab. Spironolactone 100mg (Blister Pack) (Pack in carton with leaflet)	10000	6	60000	1	1200

303	Tab. Spironolactone 50mg + Frusemide 40mg (Blister Pack) (Pack in carton with leaflet)	50000	5.33	266500	1	5330
305	Tab. Trimetazidine 35mg (Blister Pack) (Pack in carton with leaflet)	20000	7.59	151800	1	3036
306	Tab. Vitamin B6 (Pyridoxine) 50mg (Blister Pack) (Pack in carton with leaflet)	60000	1.48	88800	2	1776
307	Tab. Warfarin 5mg (Blister Pack) (Pack in carton with leaflet)	3000	9	27000	1	540
308	Susp. Aluminium hydroxide 215mg or more, Magnesium Hydroxide 80mg, Simethicon 25mg or more/5ml (Bottle of 120ml) (Pack in carton with leaflet)	60000	47	2820000	2	56400
310	Susp. Paracetamol 120mg/5ml (Bottle of 60ml) (Pack in carton)	50000	35.5	1775000	2	35500
311	Susp. Sucralfate 1gm/5ml (Bottle of 60ml) (Pack in carton with leaflet)	6000	52	312000	1	6240
313	Syp. Calcium Phosphate 210mg, Vitamin D3 350IU, Bottle of 120ml or less (Pack in carton)	2000	47.33	94660	2	1893.2
315	Syp. Dextromethophan HBr 10mg or more, Chlorpheniramine Maleate 2mg or more, Pseudoephedrine HCl 30mg or more/5ml (Bottle of 120ml or less) (Pack in carton with leaflet)	20000	13.5	270000	2	5400
316	Syp. Divalproex Sodium 250mg/5ml (Bottle of 120ml or less) (Pack in carton with leaflet)	3000	154.42	463260	2	9265.2
317	Syp. Lactulose 3.35G/5ml (Bottle of 120ml with measuring cup) (Pack in carton with leaflet)	24000	130	3120000	2	62400
318	Syp. L-Ornithine L-Aspartate 60mg, Nicotinamide 4.8mg, Riboflavin 5, Phosphate Sodium 0.153mg/ml (Bottle of 120ml) (Pack in carton with leaflet)	3000	194	582000	1	11640
319	Syp. Multivitamin + Mineral (Bottle of 120ml) (Pack in carton)	5000	69.43	347150	2	6943
320	Syp. Zinc Sulphate Monohydrate (Bottle of 60ml) (Pack in carton)	7000	35	245000	2	4900
321	Oral Rehydration Salt Sodium Chloride 2.6gm, Sodium Citrate 2.9gm, Potassium Chloride 1.5gm, Dextrose Anhydrous 13.5gm / Sachet. (Pack in carton) (Low Osmolar)	18000	5.09	91620	1	1832.4
322	Acyclovir Cream 10 gm (Pack in carton with leaflet)	2000	56.53	113060	1	2261.2
323	Aerosol Salbutamol (Pack in carton with leaflet)	1200	132	158400	1	3168
325	Betamethasone + Gentamycin Ointment 15gm (Pack in carton with leaflet)	2000	48	96000	1	1920

326	Betamethasone 17 valerate 0.1% Lotion 60ml (Pack in carton with leaflet)	2000	63	126000	1	2520
327	Betamethasone 17 valerate 0.1% Ointment 5gm (Pack in carton with leaflet)	1000	14	14000	1	280
328	Clobetasole Propionate 0.05% Ointment 10gm (Pack in carton with leaflet)	1600	15	24000	1	480
330	Clotrimazole 1% Skin Cream (10gm) (Pack in carton with leaflet)	1200	49	58800	1	1176
333	Mupirocin Ointment 2% w/w (Tube of 15gm or less) (Pack in carton with leaflet)	500	110	55000	1	1100
334	Permethrin 0.25% Lotion 60ml (Pack in carton with leaflet)	3000	60	180000	1	3600
335	Polymyxin Beta sulphate 10,000 or less units/gm, Bacitracin 500 units/gm Skin Ointment 20gm or Less (Pack in carton with leaflet)	72000	45.43	3270960	2	65419.2
336	Polymyxin Beta sulphate 10,000 or less units/gm, Bacitracin 500 units/gm, Lignocaine 40mg/gm (20gm or Less) (Pack in carton with leaflet)	20000	31.39	627800	1	12556
337	Silversulphadiazine 1% Cream 50G (Pack in carton with leaflet)	80000	70	5600000	2	112000
338	Acetylcysteine 200mg (Sachet) (Pack in carton with leaflet)	20000	14.06	281200	2	5624
339	Eneema (sodium-biphosphate 19.2 gm, Sodium phosphate 7.2 gm/120ml (Bottle 135ml or Less) with Nozzle (Pack in carton with leaflet)	8000	65.75	526000	2	10520
341	Salbutamol Nebulizing Solution (20ml) (Pack in carton with leaflet)	12000	31.65	379800	2	7596
342	Inj Hydroxy propyl methyl cellulose 2% (vial/Amp 2.5/3ml) (Pack in carton with leaflet)	10000	130	1300000	2	26000
343	Inj. Carbachol 0.01% Solution (vial/Amp 1.5/2ml) (Pack in carton with leaflet)	3000	306	918000	2	18360
344	Eye Drop Atropine 1% (Bottle of 15ml or less) (Pack in carton with leaflet)	1000	22.83	22830	1	456.6
345	Eye Drop Cyclopentolate 1% (Bottle of 15ml or less) (Pack in carton with leaflet)	1000	50	50000	1	1000
346	Eye Drop Ciprofloxacin 0.3% (Bottle of 5ml) (Pack in carton with leaflet)	5000	51.6	258000	2	5160
347	Eye Drop Dexamethasone 0.1%, Neomycin Sulphat 0.35%, Polymyxin B Sulphate 6,000 iu (Bottle of 5ml) (Pack in carton with leaflet)	3000	50.00	150000	1	3000
348	Eye Drop Dexamethasone 0.1%, (Bottle of 5ml) (Pack in carton with leaflet)	2000	58.00	116000	1	2320
349	Eye Drop Diclofenac Sodium 0.1% (Bottle of 5ml) (Pack in carton with leaflet)	2000	35	70000	1	1400

350	Eye Drop Dorzolamide 2% + Timolol 0.5% (Bottle of 5ml or less) (Pack in carton with leaflet)	3600	240	864000	2	17280
351	Eye Drop Fluorometholone 0.1% (Bottle of 5ml) (Pack in carton with leaflet)	1000	75	75000	1	1500
352	Eye Drop Levobunolol HCl 0.5% (Bottle of 5ml) (Pack in carton with leaflet)	3000	80	240000	2	4800
354	Eye Drop Pilocarpine 2% (Bottle of 5ml) (Pack in carton with leaflet)	1500	66.98	100470	2	2009.4
355	Eye Drop Polyvinyl Alcohol (Bottle of 15ml or less) (Pack in carton with leaflet)	4000	53.55	214200	2	4284
356	Eye Drop Proparacaine HCl (Bottle of 15ml) (Pack in carton with leaflet)	2000	66.98	133960	2	2679.2
357	Eye Drop Tobramycin 0.3% (Bottle of 5ml) (Pack in carton with leaflet)	4000	34.78	139120	1	2782.4
359	Eye Drop Tropicamide 1% (Bottle of 15ml or less) (Pack in carton with leaflet)	1000	72.67	72670	1	1453.4
360	Eye Oint Polymyxin Beta sulphate 10,000 units/gm, Bacitracin 500 units/gm 6grm (Pack in carton with leaflet)	10000	19.87	198700	1	3974
361	Nasal Spray Xylometazolin 1:1000 Bottle of 20ml or Less (Pack in carton with leaflet)	3000	34.33	102990	1	2059.8
365	Tab. Ibuprofen 400mg (Blister Pack) (Pack in carton with leaflet)	100000	2.7	270000	1	5400
366	Syp. Domperidone 1mg/ml 120ml (Pack in carton with leaflet)	20000	80	1600000	2	32000
367	Tab. Sofosbuvir 400mg + Velpatasvir 100mg (Pack in carton with leaflet)	201600	303	61084800	2	1221696
368	Tab. Entacavir 0.5mg (Pack in carton with leaflet)	20000	136	2720000	2	54400
369	Tab. Tenofovir Alafenamide Fumarate 25mg	24000	94	2256000	2	45120
370	Hepatitis B Vaccine with Disposable Syringes 1ml	4800	279	1339200	2	26784
372	Tissue Plasminogen Activata (TPA)	120	80500	9660000	3	193200
4	Inj Acyclovir Sodium 500mg with water for injection (Vial) (Pack in carton with leaflet)	20000	250	5000000	2	100000
5	Inj Acyclovir Sodium 250mg with water for injection (Vial) (Pack in carton with leaflet)	12000	410	4920000	2	98400
41	Inj Bupivacaine HCl 0.5% (Amp of 10ml) (Pack in carton with leaflet)	30000	19	570000	1	11400
54	Inj Amikacin Sulphate 500mg(vial) (Pack in carton with leaflet)	50000	47.9	2395000	2	47900
63	Inj Ceftriaxone 1G IV,IM(Vial) with water for injection (Pack in carton with leaflet)	400000	66.5	26600000	2	532000
75	Inj. Cefuroxime Sodium 750mg, Vial with water for injection (Pack in carton with leaflet)	6000	59	354000	1	7080
100	Tab. Terbinafine 125mg (Blister Pack) (Pack in carton with leaflet)	5000	18	90000	1	1800

105	Susp/Syp. Cefixime 100mg/5ml Bottle of 30ml (Pack with teaspoon in carton with leaflet)	12000	80	960000	1	19200
115	inj Dacarbazine 200mg(Vial) (Pack in carton with leaflet)	180	425	76500	1	1530
119	Inj Doxorubicin 10mg (Vial) (Pack in carton with leaflet)	6000	403	2418000	2	48360
137	Inj Topotecan 4mg (Vial) (Pack in carton with leaflet)	60	7000	420000	2	8400
140	Inj Zolidronic Acid 4mg (Vial) (Pack in carton with leaflet)	2400	1674	4017600	2	80352
141	Tab/Cap. Lenvatinib 4mg (Blister Pack) (Pack in carton with leaflet)	6000	1100	6600000	1	132000
144	Tab/Cap. Palbociclib 125mg (Blister Pack) (Pack in carton with leaflet)	13000	12000	156000000	2	3120000
155	Tab. Sunitinib 12.5mg(Blister Pack) (Pack in carton)	21600	500	10800000	1	216000
161	Inj Lung Surfactant(Bovine lipid Extract) (Vial of 3ml) (Pack in carton with leaflet)	500	12750	6375000	2	127500
182	Inj. Dexamethasone 4 mg/ml (amp of 1 ml) (pack in carton with leaflet)	240000	8.77	2104800	2	42096
187	Inj. Dobutamine 250mg (Amp of 5ml) (Pack in carton with leaflet)	24000	330	7920000	2	158400
216	Inj. Phenytoin Sodium 50mg/ml (Ampoule of 5ml) (Packed in carton with leaflet)	40000	175.47	7018800	2	140376
222	Inj. Tramadol HCl 100mg/2ml (Amp of 2ml) (Pack in carton with leaflet)	60000	7.9	474000	1	9480
234	Cap/Tab. Tranexamic acid 500mg (Blister Pack) (Pack in carton with leaflet)	10000	15.55	155500	1	3110
240	Tab. Amlodipine 5mg (Blister Pack) (Pack in carton with leaflet)	50000	1.72	86000	1	1720
254	Tab. Carvedilol 6.25mg (Blister Pack) (Pack in carton with leaflet)	12000	1.56	18720	1	374.4
255	Tab. Cetirizine 10mg (Blister Pack) (Pack in carton with leaflet)	120000	1.24	148800	1	2976
257	Tab. Clopidogrel 75mg (Blister Pack) (Pack in carton with leaflet)	240000	3.4	816000	2	16320
258	Tab. Clopidogrel 75mg + Aspirin 75mg (Blister Pack) (Pack in carton with leaflet)	80000	7.5	600000	2	12000
262	Tab. Diclofenac Sodium 50mg (Blister Pack) (Pack in carton with leaflet)	200000	0.79	158000	1	3160
268	Tab. Escitalopram 10mg (Blister Pack) (Pack in carton with leaflet)	10000	3.1	31000	1	620
275	Tab. Glimipride 4mg (Blister Pack) (Pack in carton with leaflet)	40000	2.75	110000	1	2200
281	Tab. Lisinopril 5mg (Blister Pack) (Pack in carton with leaflet)	60000	2.8	168000	2	3360
282	Tab. Losartan Potassium 50mg (Blister Pack) (Pack in carton with leaflet)	15000	3.8	57000	1	1140

291	Tab. Naproxen 500mg (Blister Pack) (Pack in carton with leaflet)	100000	8.9	890000	1	17800
292	Tab. Paracetamol 500mg (Blister Pack) (Pack in carton with leaflet)	900000	1.41	1269000	2	25380
293	Tab. Paracetamol 650mg + Orphenadrin Citrate 50mg (Blister Pack) (Pack in carton with leaflet)	150000	4.44	666000	1	13320
314	Syp. Cetirizine 1mg/ml (Bottle of 60ml) (Pack in carton with leaflet)	12000	19.7	236400	1	4728
329	Clotrimazole 1% + Hydrocortisone 1% Skin Cream (10gm) (Pack in carton with leaflet)	1200	55	66000	1	1320
331	Fusidic Acid 20mg + Hydrocortisone 10mg / gm Cream (15gm) (Pack in carton with leaflet)	9600	195	1872000	1	37440
332	Fusidic Acid Cream 15gm (Pack in carton with leaflet)	9600	78	748800	1	14976
340	Ipratropium Bromide Nebulizing Solution 250mcg/ml (20ml) (Pack in carton with leaflet)	24000	116.82	2803680	2	56073.6
364	Tab. Amlodipine 5mg + Valsartan 80mg (Blister Pack) (Pack in carton with leaflet)	20000	20	400000	1	8000
371	Paracetamol Suppository 250mg (Pack in carton)	1000	184.9	184900	1	3698