Pre-Qualification Document

Prequalification of Firms for the Supply of Drugs and Medicines (Bulk Purchase) to PIMS and SSP funded Patients for the Period of 02 Years (Goods)

National

Single Stage-Two Envelope

August 03, 2026

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

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INVITATION FOR PRE-QUALIFICATION

PROCUREMENT OF GOODS

1. The **Purchase Department (Pakistan Institute of Medical Sciences (PIMS))** has reserved Funds for the procurement planned for FY **2026-27**. The **Purchase Department (Pakistan Institute of Medical Sciences (PIMS))** intends to apply part of the proceeds of this Fund to cover eligible payments under the contract for the "**Prequalification of Firms for the Supply of Drugs and Medicines (Bulk Purchase) to PIMS and SSP funded Patients for the Period of 02 Years**" with the reference of **"P56051"**

2. The **Purchase Department (Pakistan Institute of Medical Sciences (PIMS))** intends to pre-qualify suppliers for Invitation to Bid(s), and sign the contract agreement(s) with the selected bidder(s) subsequent to bidding process.

3. The objective of the intended pre-qualification is the on-demand supply of " **Prequalification of Firms for the Supply of Drugs and Medicines (Bulk Purchase) to PIMS and SSP funded Patients for the Period of 02 Years** " through subsequent signing of contract with successful bidders, and the purpose of this Pre-qualification Notice is to provide the very basic information to enable the potential applicants to decide whether or not to respond to this Pre-qualification Notice.

4. Only the pre-qualified applicants shall be entitled to participate in the procurement proceedings, and it is expected that the Invitation to Bids will be made to the Pre-qualified Applicants.

5. Pre-qualification process is open for all **National** Applicants subject to fulfilling the eligibility requirements mentioned in the respective Pre-qualification Documents. Interested Applicants may obtain further information from the Purchase Department (Pakistan Institute of Medical Sciences (PIMS)) through **EPADS v2.0** during office hours. A complete set of Pre-qualification Documents may be accessed by interested Applicants through **EPADS v2.0** at **https://epads.gov.pk/opportunities/federal/procurements/56051.**

6. The application, prepared in accordance with the instructions in the Pre-qualification Documents, must be submitted through **EPADS v2.0** on or before **Saturday, August 29, 2026 11:00 AM**. E-bids will be opened using **EPADS v2.0** on the same day at **Saturday, August 29, 2026 11:30 AM**. Manual submission of Bids shall not be entertained. Those vendors who have not yet registered on the new version of **EPADS v2.0,** may register themselves on **https://vendors.epads.gov.pk/**. A tutorial to explain the registration process is available at **https://www.youtube.com/watch?v=MNW6T38v7tc**

In terms of Rule 48 of Public Procurement Rules, 2004 Grievance Redressal Committee (GRC) is notified for the subject procurement and notification copy is available on the procuring agency's website and on Authority's website at (**www.ppra.org.pk**).

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

A. General

B. Contents of the Prequalification Documents

Sections of Prequalifica¬tion Documents

1. **Scope of Application**

1.1. In connection with the "**Invitation for Prequalification**", the Procuring Agency, as defined in Section II (Prequalification Data Sheet abbreviated as PDS), issues this set of Prequalification Documents (PD) to prospective applicants (also hereinafter referred as Applicants) interested in submitting applications (also hereinafter referred as Applications) to determine the capacity and capability of the Applicant(s) for supply of Goods and Related Services incidental thereto as specified in Section VII (Schedule of Requirements).

2. **Source of Funds**

2.1. Source of funds is same as referred in Invitation for Pre-qualification.

3. **Fraud and Corruption**

3.1. The Procuring Agency requires that the Applicants /Bidders/ Suppliers/Contractors under Government financed contracts, observe the highest standard of ethics during the procurement and execution of such agreements and contracts.

3.2. The Applicants/Bidders shall permit and shall cause their agents (whether declared or not), sub-contractors, sub-consultants, service providers, suppliers, and their personnel, to permit the Procuring Agency to inspect all accounts, records and other documents relating to any, Application/Bid submission, Secondary Procurement process, and to have them audited by auditors appointed by the Procuring Agency.

3.3. Any communications between the Applicant and the Procuring Agency related to matters of alleged corrupt and fraudulent practices must be made in writing or in electronic forms that provide record of the content of

communication.

3.4. Procuring Agency will reject an application or bid or proposal, if it is established that the Applicant or the Bidder or Prosper was engaged in corrupt and fraudulent practices in competing for the contract.

3.5. Procuring Agency will also declare the Applicant as blacklisted in accordance with rules and predefined standard mechanism.

4. **Eligible Applicants**

4.1. An Applicant may be a private entity, a state-owned enterprise or institution subject to ITB 4.6, or any combination of such entities in the form of a joint venture (JV) under an existing JV agreement or with the intent to enter into such an agreement supported by a letter of intent.
In case of single (private or state-owned entity), it shall be liable for execution of all the provisions of the Contract Agreement.
In the case of a joint venture, all members shall be jointly and severally liable for the execution of all the provisions of the Contract Agreement (if signed b/w the Procuring Agency and the JV), in accordance with the Contract conditions that apply.
The JV shall nominate a Representative who shall have the authority to conduct all business for and on behalf of any and all the members of the JV during the Prequalification process, Bidding process (in the event the prequalified JV submits a Bid) and during the period of contract agreement and contract execution (in the event the JV is awarded the Contract). Unless specified in the PDS, there is no limit on the number of members in a JV.

4.2. An Applicant may apply for Prequalification both individually, and as part of a joint venture, or participate as a subcontractor. If prequalified as a JV only, it will not be permitted to bid for the same contract as an individual entity. Bids submitted in violation of this provision will be rejected.

4.3. An Applicant and any of its affiliates (that directly or indirectly control, are controlled by or are under common control with that entity) may submit its Application for Prequalification either individually, as joint venture or as a sub-contractor among them for the same contract. However, if prequalified only one prequalified Applicant will be allowed to bid for the same contract.

All Bids submitted in violation of this provision will be rejected.

4.4. Applicants shall be considered to have a conflict of interest, if they participated as a consultant in the preparation of the design or technical specifications or have been hired or proposed to be hired by the Procuring Agency for execution of subsequent Contract Agreement. In addition, Applicants may be considered to have a conflict of interest if they have a close business or family relationship with such professional staff of the Procuring Agency (or a recipient of a part of the funds) who:

4.4.1. are directly or indirectly involved in the preparation of the Prequalification Documents or Bidding Documents or specifications of Contract and/or the Prequalification or Bid evaluation process of such Contract; or

4.4.2. would be involved in the implementation or supervision of such Agreement t, unless the conflict stemming from such relationship has been resolved throughout the Procurement Process, Bidding.

4.5. An Applicant that has been declared debarred or blacklisted shall be ineligible to be prequalified to bid for such period of time and for such type of procurement for which he has been declared debarred or blacklisted. The list of debarred firms and individuals is available at PPRA's website.

4.6. An Applicant shall provide such documentary evidence for determining the eligibility of the Applicant to the reasonable satisfaction of the Procuring Agency.

5. **Eligibility (in terms of Nationality)**

5.1. Applicants may be ineligible if they are nationals of ineligible countries as indicated in Section V.

B. Contents of the Prequalification Documents

6. **Sections of Prequalification Documents**

6.1. This set of Prequalification Documents consists of Parts 1 and 2 which comprise all the sections indicated below, and which should be read in conjunction with any Addendum issued in accordance with ITA 8.

PART 1 Prequalification Procedures PART 2 Supply Requirements

6.2. Section I - Instructions to Applicants (ITA)

6.3. Section II - Prequalification Data Sheet (PDS)

6.4. Section III - Qualification Criteria and Requirements

6.5. Section IV - Application Forms

6.6. Section V - Eligible Countries

6.7. Section VI - Fraud and Corruption

6.8. Section VII – Schedule of Requirements

6.9. The Procuring Agency accepts no responsibility for the completeness of the Prequalification documents, responses to requests for clarification, the minutes of the pre-Application meeting (if any), or Addenda to the Prequalification documents in accordance with ITA 8. In case of any discrepancies, documents issued directly through ePADS shall prevail.

6.10. The Applicant is expected to examine all instructions, forms, and terms in the Prequalification Documents and to furnish with its Application all information or documentation as is required by the Prequalification Documents.

7. **Clarification of Pre-qualification Documents and Pre-Application Meeting**

7.1. An Applicant requiring any clarification of the Pre-qualification Documents shall contact the Procuring Agency in writing through ePADS as. The Procuring Agency will respond in writing through ePADS to any request for clarification provided that such request is received no later than three (03) days prior to the deadline for submission of the Applications. The Procuring Agency shall forward a copy of its response to all prospective

Applicants through ePADS who have obtained the Prequalification Documents from ePADS , including a description of the inquiry but without identifying its source. If so indicated in the PDS, the Procuring Agency shall also promptly publish its response through ePADS . Should the Procuring Agency deem it necessary to amend the Prequalification Documents as a result of a clarification, it shall do so in accordance with the provisions of ITA 16.2.

7.2. If indicated in the PDS, the Applicant's designated representative is invited at the Applicant's cost to attend a pre-Application meeting through online platform / **EPADS v2.0** as per date and time mentioned in the PDS. During this Pre-Application meeting, prospective Applicants may request clarification of the schedule of requirement, the qualification criteria or any other aspects of the Pre-qualification Documents.

7.3. Minutes of the Pre-Application meeting, if applicable, including the text of the questions asked by Applicants, including those during the meeting (without identifying the source) and the responses given, together with any responses prepared after the meeting will be transmitted promptly through ePADS to all prospective Applicants who have obtained the Pre-qualification Documents. Any modification to the Pre-qualification Documents that may become necessary as a result of the pre-Application meeting shall be made by the Procuring Agency exclusively through the use of an Addendum pursuant to ITA 8 and through **EPADS v2.0**. Non-attendance at the pre-Application meeting will not be a cause for disqualification of an Applicant.

8. **Amendment of Prequalifica¬tion Documents**

8.1. At any time prior to the deadline for submission of Applications, the Procuring Agency may amend the Prequalification Documents by issuing an Addendum through **EPADS v2.0**

8.2. Any Addendum issued shall be part of the Prequalification Document and shall be communicated in writing through ePADS to all Applicants who have obtained the Prequalification Documents from the Procuring Agency. The Procuring Agency shall promptly publish the Addendum at the Procuring Agency's web page and **EPADS v2.0**.

Provided that an Applicant who had already submitted their Applications

prior to the issuance of any such addendum shall have the right to withdraw his already submitted Application and submit the revised Application prior to the original or extended Application submission deadline through **EPADS v2.0**.

8.3. To give Applicants reasonable time to take an Addendum into account in preparing their Applications, the Procuring Agency may at its discretion, extend the deadline for the submission of Applications in accordance with ITA 16.2:
Provided that the Procuring Agency shall extend the deadline for submission of Applications, if such an addendum is issued within last three (03) days of the Application submission deadline.

C. Preparation of Applications

9. **Cost of Applications**

9.1. The Applicant shall bear all costs associated with the preparation and submission of its Application. The Procuring Agency will in no case be responsible or liable for those costs, regardless of the conduct or outcome of the Pre-qualification process.

10. **Language of Application**

10.1. The Application as well as all correspondence and documents relating to the Pre-qualification exchanged by the Applicant and the Procuring Agency, shall be written in the language specified in the PDS. Supporting documents and printed literature that are part of the Application may be in another language, provided they are accompanied by an accurate translation of the relevant passages in the language specified in the PDS, in which case, for purposes of interpretation of the Application, the translation shall govern.

11. **Documents Comprising the Application**

11.1. The Application shall comprise the following:

11.1.1. **Application Submission Letter**, in accordance with ITA 12.1;

11.1.2. **Eligibility**: documentary evidence establishing the Applicant's eligibility, in accordance with ITA 13.1;

11.1.3. **Qualifications**: documentary evidence establishing the Applicant's qualifications, in accordance with ITA 14; and

11.1.4. any other document required as specified in the PDS.

11.2. **Application Submission Letter**

11.2.1. The Applicant shall complete an Application Submission Letter as provided in Section IV (Application Forms). This Form must be completed without any alteration to its format.

11.3. **Documents Establishing the Eligibility of the Applicant**

11.3.1. To establish its eligibility in accordance with ITA 4, the Applicant shall complete the eligibility declarations in the Application Submission Letter and Form ELI-1.1 (eligibility), included in Section IV (Application Forms).

11.4. **Documents Establishing the Qualifications of the Applicant**

11.4.1. To establish its qualifications to perform the contract(s) in accordance with Section III (**Qualification Criteria and Requirements**), the Applicant shall provide the information requested in the corresponding Information Sheets included in **Section IV (Application Forms)**.

11.4.2. Wherever an Application Form requires an Applicant to state a monetary amount, Applicants should indicate the Pak Rupee equivalent using the rate of exchange determined as follows:

11.4.2.1. for turnover or financial data required for each year - Exchange rate prevailing on the last day of the respective calendar year (in which the amounts for that year is to be converted).

11.4.2.2. value of single contract - Exchange rate prevailing on the date of the contract.

11.4.3. Exchange rates shall be taken from the publicly available source identified in the PDS. Any error in determining the exchange rates in the Application may be corrected by the Procuring Agency.

11.4.4. The documentary evidence of the Applicant’s qualifications to conclude a contract Agreement, shall establish to the Procuring Agency’s satisfaction:

11.4.4.1. that, if required in the BDS, an Applicant that does not manufacture or produce the Goods it offers to supply shall submit the Manufacturer’s Authorization using the form included in Section IV A (Bidding Forms) to demonstrate that it has been duly authorized by the manufacturer or producer of the Goods to supply these Goods in the Procuring Agency’s Country;

11.4.4.2. that, if required in the BDS, in case of an Applicant not doing business within Islamic Republic of Pakistan (or the country where the procurement is being made), the Applicant is, or will be, (if awarded the call off contract) represented by an Agent in the country, equipped and able to carry out the Supplier’s maintenance, repair, and spare parts stocking obligations in respect of the Goods.

D. Submission of Applications

15. **Submission of the Applications through EPADS v2.0**

15.1. The Bidder shall prepare and submit Bid with due diligence after carefully reading all the terms and condition before submission through ePADS in accordance with the procedures specified in the PDS.

15.2. In case the Applicant is a JV, the Application shall submit an authorized representative of the JV on behalf of the JV and so as to be legally binding on all the members as evidenced by a power of attorney signed by their legally

authorized signatories.

16. **Deadline for Submission of Applications**

16.1. Applicants shall be submitted through ePADS no later than the deadline indicated in the PDS.

16.2. If required in accordance with the provisions of ITA 8.3, the Procuring Agency will extend the deadline for the submission of Applications, in which case all rights and obligations of the Procuring Agency and the Applicants subject to the previous deadline shall thereafter be subject to the deadline as extended.

16.3. The deadline will be extended in the same manner as that of original Invitation for Prequalification (or the advertisement) through **EPADS v2.0**.

17. **Opening of Applications**

17.1. The Procuring Agency shall open all Applications on the date and time specified in the PDS through **EPADS v2.0**. Late Applications shall be treated in accordance with ITA 16.1.

E. Procedures for Evaluation of Applications

18. **Confidentiality**

18.1. Information relating to the Applications, their evaluation and results of the Prequalification shall not be disclosed to Applicants or any other persons not officially concerned with the Prequalification process until the notification of Prequalification results is made to all Applicants in accordance with ITA 26 through **EPADS v2.0**.

18.2. From the deadline for submission of Applications to the time of notification of the results of the Prequalification in accordance with ITA 26, any Applicant that wishes to contact the Procuring Agency on any matter related to the Prequalification process may do so only in writing through **EPADS v2.0**

19. **Clarification of Applications**

19.1. To assist in the evaluation of Applications, the Procuring Agency may, ask an Applicant for a clarification (including missing documents) of its Application, to be submitted within a stated reasonable period of time. Any request for clarification from the Procuring Agency and all clarifications from the Applicant shall be in writing through **EPADS v2.0**

19.2. If an Applicant does not provide clarifications and/or documents requested by the date and time set in the Procuring Agency's request for clarification, its Application shall be evaluated based on the information and documents available at the time of evaluation of the Application.

20. **Responsiveness of Applications**

20.1. The Procuring Agency may reject any Application which is not responsive to the requirements of the Prequalification Documents. In case the information furnished by the Applicant is incomplete or otherwise requires clarification as per ITA 19.1, and the Applicant fails to provide satisfactory clarification and/or missing information within prescribed time, it may result in disqualification of the Applicant.

21. **Margin of Preference**

21.1. Unless otherwise specified in the PDS, a margin of preference shall not apply in the Bidding process resulting from this Pre-qualification.

22. **Sub-contractors**

22.1. Subcontractors' qualification and experience will not be considered for evaluation of the Applicant. The Applicant on its own (without taking into account the qualification and experience of the Subcontractor) should meet the qualification criteria.

F. Evaluation of Applications and Prequalification of Applicants

23. **Evaluation of Applications**

23.1. The Procuring Agency shall use the factors, methods, criteria, and requirements defined in Section III, Qualification Criteria and Requirements, to evaluate the qualifications of the Applicants, and no other methods, criteria, or requirements shall be used. The Procuring Agency reserves the right to waive minor deviations from the qualification criteria if they do not materially affect the technical capability and financial resources of an Applicant to perform the contract, however subject to the provisions of ITA 25.

23.2. Subcontractors proposed by the Applicant shall be fully qualified for their parts of the Scope of Supply of the Goods and Allied Services.

23.3. In case of multiple contracts, Applicants should indicate in their Applications the individual contract or combination of contracts in which they are interested. The Procuring Agency shall prequalify each Applicant for the maximum combination of contracts for which the Applicant has thereby indicated its interest and for which the Applicant meets the appropriate aggregate requirements. The Qualification Criteria and Requirements are mentioned in Section III.

Only the qualifications of the Applicant shall be considered. The qualifications of other related entities such as the Applicant's subsidiaries, parent entities, affiliates, subcontractors or any other firm(s) different from the Applicant shall not be taken into consideration in determining the qualifications of the Applicant.

24. **Procuring Agency's Right to Accept or Reject Applications**

24.1. The Procuring Agency reserves the right to accept or reject all the Applications, and to annul the Prequalification process at any time, without thereby incurring any liability to the Applicants. However, the procuring agency shall record its reasons and justifications on **EPADS v2.0**, duly approved by the Principal Accounting Officer or Head of Organization.

25. **Pre-qualification of Applicants**

25.1. All Applicants whose Applications substantially meet or exceed the specified qualification requirements will be prequalified by the Procuring

Agency.

25.2. An Applicant may be “conditionally prequalified,” that is, qualified subject to the Applicant submitting or correcting certain specified nonmaterial documents or deficiencies to the satisfaction of the Procuring Agency.

25.3. Applicants that are conditionally prequalified will be so informed along with the statement of the condition(s) which must be met to the satisfaction of the Procuring Agency before or at the time of submitting their Bids.

26. **Notification of Prequalification**

26.1. The Procuring Agency shall notify all Applicants in writing through **EPADS v2.0** indicating the names of those Applicants who have been prequalified or conditionally prequalified. In addition, those Applicants who have been disqualified will be informed separately through **EPADS v2.0**.

26.2. The procuring agency shall communicate to those suppliers or contractors who have not been pre-qualified the reasons for not pre-qualifying them through **EPADS v2.0**

27. **Request for Bids**

27.1. Promptly after the notification of the results of the Prequalification, the Procuring Agency will invite the Bids from all the Applicants that have been prequalified through **EPADS v2.0**.

28. **Changes in Qualifications of Applicants**

28.1. Any change in the structure or formation of an Applicant after being prequalified in accordance with ITA 25 and invited to bid (including, in the case of a JV, any change in the structure or formation of any member thereto) shall be subject to the written approval of the Procuring Agency prior to the deadline for submission of Bids. Such approval shall be denied if:

28.1.1. a prequalified Applicant proposes to associate with a disqualified Applicant or in case of a disqualified joint venture, any of its members;

28.1.2. as a consequence of the change, the Applicant no longer substantially meets the qualification criteria set forth in Section III, Qualification Criteria and Requirements; or

28.1.3. in the opinion of the Procuring Agency, the change may result in a substantial reduction in competition.

28.2. Any such change should be submitted to the Procuring Agency before the date of “Invitation to Bids”.

29. **Redressal of Grievances**

29.1. Procuring agency shall constitute a Grievance Redressal Committee (GRC) and proceed in accordance with the procedure and mechanism defined under Rule-48 of Public Procurement Rules, 2004.

29.2. The GRC shall not have any of the members of Procurement Evaluation Committee. The committee must have one subject specialist depending on the nature of the procurement.

30. **Mechanism of Blacklisting**

30.1. The procuring agency shall initiate blacklisting or debarment proceedings against any bidder, supplier or contractor in accordance with the mechanism prescribed under Rule-19 of Public Procurement Rules, 2004 read with “Mechanism for Blacklisting Regulations, 2024”.

Pre-qualification Data Sheet

Prequalification Data Sheet (PDS)

The following specific data for the Prequalification of Applicants shall complement, supplement, or amend the provisions in the Instructions to Applicants (ITA). Whenever there is a conflict, the provisions herein shall prevail over those in ITA.

PDS Clause No
ITA No

Amendments of, and Supplements to, Clauses in the Instructions to Applicants

A. General

PDS Clause No 1
ITA No 1.1

Identification Number of the Invitation for Prequalification: **P56051**
The Procuring Agency is: **Purchase Department (Pakistan Institute of Medical Sciences (PIMS))**
List of Contracts*:*
See section items and Lots

PDS Clause No 2
ITA No 2.1

The name of Procuring Agency is: **Purchase Department (Pakistan Institute of Medical Sciences (PIMS))**
The name of Project / Procurement is: **Prequalification of Firms for the Supply of Drugs and Medicines (Bulk Purchase) to PIMS and SSP funded Patients for the Period of 02 Years**

PDS Clause No 3
ITA No 4.2

Maximum number of members in a Joint Venture (JV): **Nil**

PDS Clause No 4
ITA No 4.5

A list of debarred firms and individuals is available on PPRA website: **https://ppra.gov.pk**

B. Contents of the Prequalification Document

PDS Clause No 5
ITA No 7.1

For clarification, the Applicant shall seek clarifications through: **EPADS v2.0**

PDS Clause No 6
ITA No 7.1 & 8.2

Information related to Prequalification shall be published on: **EPADS v2.0**

PDS Clause No 7
ITA No 7.2

Pre-Application Meeting: **Clarification Date: Thursday, August 20, 2026**

C. Preparation of Applications

PDS Clause No 8
ITA No 10.1

This Prequalification Document has been issued in the language: **English**

PDS Clause No 9
ITA No 11.1(d)

Additional documents to be submitted through EPADS v2.0:
No

PDS Clause No 10
ITA No 14.2

Source for determining exchange rates: **Not Applicable**

D. Submission of Applications

PDS Clause No 11
ITA No 16.1

Deadline for Application Submission:
Day: ***Saturday***
Date: ***Saturday, August 29, 2026***
Time: ***11:00 AM***

PDS Clause No 12
ITA No 17.1

Opening of Applications shall be conducted through: **EPADS v2.0**
Day: ***Saturday***
Date: ***Saturday, August 29, 2026***
Time: ***11:30 AM***
Virtual participation link: **https://vendors.epads.gov.pk/**

E. Procedures for Evaluation of Applications

PDS Clause No 13
ITA No 21.1

Margin of Domestic Preference: **Not Applicable**
(Applicable only if authorized in Procurement Plan)

PDS Clause No 14
ITA No 29.1

Prequalification-related complaints / grievances shall be submitted in writing through: **EPADS v2.0**
A complaint may challenge:

- The terms of the Prequalification Documents
- The Procuring Agency’s decision not to prequalify an Applicant

Eligibility & Qualification Criteria

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

Eligibility Criteria	Document
Name, Designation & Specimen signature of authorized / focal person with cell / what’s app No. and E mail addresses on official Firm’s letter pad.	Yes
The firm shall submit original receipt of tender fee of Rs. 2000/- with prequalification application.	Yes
The firm undertakes that currently it is not blacklisted / debarred by any Govt. / Semi government / Autonomous dept. /hospital and provide undertaking in this regard on legally notarized e-stamp paper of rupees 100/-	Yes
The firm shall provide valid Drugs Manufacturing License/Drug Distributor License/ Drug Sale License for Sole Agents of Foreign Principal issued / import license by DRAP accordingly	Yes

Valid GMP Certificate or Valid Satisfactory GMP Inspection Report issued by DRAP (for local manufacturers/distributors). Only those manufacturing sections and pharmaceutical categories that are declared Satisfactory in the GMP Inspection Report and/or are specifically mentioned in the GMP Certificate shall be considered for prequalification. For Sole Agents of Foreign Principles, a Valid COPP (Certificate of Pharmaceutical Product).	Yes
Facility is having functional and validated, Heating Ventilation & Air Conditioning System (HVAC). Firm shall provide undertaking in this regard on legally notarized e-stamp paper of rupees 100.	Yes
The firm shall submit undertaking on Rs.100 e-stamp paper that the firm complies with all the regulatory requirement of the country.	Yes
The firm shall submit undertaking on Rs.100 e-stamp paper that none of its supplied batch of any drap registered section to any hospital has been declared Spurious/Adulterated since last two years till the closing date of Prequalification Document submission.	Yes
The firm shall submit undertaking on Rs.100 e-stamp paper that tested samples of the quoted item declared Substandard/Misbranded (if any) by any DTLs (Not over 5%) since last two years till the closing date of documents submission.	Yes
The applicant shall submit an affidavit on Rs.100/- on e-stamp paper (Notarized) stating the applicant is submitted all documents and accept all the terms and conditions of the prequalification tender Documents.	Yes
The applicant shall submit an affidavit on Rs.100/- on e-stamp paper (Notarized) stating that the applicant will not skip tender proceedings in future if pre-qualified.	Yes

Evaluation Criteria

Quality Based Selection (QBS)

Technical Marks	100
Passing Marks	**70**
Technical Evaluation Criteria	
Annual Financial turnover for any of single financial year (i.e. during the last three financial years) / single calendar year (i.e. during the last three calendar years) (Attach audited financial statements) (Qualitative)(Doc Required) Minimum Annual Financial turnover for any of single financial year (i.e. during the last three financial years) / single calendar year (i.e. during the last three calendar years) more than 30 million (Attach audited financial statements) **(40)** Minimum Annual Financial turnover for any of single financial year (i.e. during the last three financial years) / single calendar year (i.e. during the last three calendar years) within 20 million to 30 million (Attach audited financial statements) **(30)** Minimum Annual Financial turnover for any of single financial year (i.e. during the last three financial years) / single calendar year (i.e. during the last three calendar years) within 10 million to 20 million (Attach audited financial statements) **(20)** Minimum Annual Financial turnover for any of single financial year (i.e. during the last three financial years) / single calendar year (i.e. during the last three calendar years) within 5 million to 10 million (Attach audited financial statements) **(10)**	40

The firm shall provide Experience letter for the supply of Drug Medicine to teaching government, Semi government / Autonomous dept. /hospital (Attach documentary evidence), i.e., experience certificate and copy of award letter etc. (Qualitative)(Doc Required) More than 04 years **(40)** More than 03 years to 04 years **(30)** More than 02 years to 03 years **(20)** More than 01 to 02 years **(10)**	40
Number of licensed and qualified Pharmacist in Regular employment of the firm (Qualitative)(Doc Required) If more than 02 Pharmacist **(20)** 01 to 02 Pharmacist **(10)**	20

PUBLIC PROCUREMENT REGULATORY AUTHORITY
(PPRA)
EPADS

Annexure

Annexure - 1 Request for Prequalification Documents

<u>Annex-1-(On firm’s Original Letter Head)</u>
Technical Submission (Vendor)

Document Required

See Form Under Additional Forms and Documents: **Annexure - 1 Request for Prequalification Documents** (page number: 32)

GENERAL FIRM’S INFORMATION

Technical Submission (Vendor)

Document Required

See Form Under Additional Forms and Documents: **GENERAL FIRM’S INFORMATION** (page number: 33)

Application Submission Form

Technical Submission (Vendor)

Document Required

See Form Under Additional Forms and Documents: **Application Submission Form** (page number: 39)

Instructions to Applicants (ITA)

Information (Read-Only)

See Form Under Additional Forms and Documents: **Instructions to Applicants (ITA)** (page number: 40)

Schedule

Information (Read-Only)

See Form Under Additional Forms and Documents: **Schedule** (page number: 41)

Procurement Forms

PUBLIC PROCUREMENT REGULATORY AUTHORITY
(PPRA)
EPADS

PUBLIC PROCUREMENT REGULATORY AUTHORITY
(PPRA)
EPADS

Additional Forms and Documents

Ref.No/ Dated: _____

The Executive Director,
PIMS, Islamabad

Subject: ***Request Application for Pre-qualification Documents for the Supply of Drugs and Medicines (Bulk Purchase) to PIMS & SSP funded Patients for Period of 02 Years.***

Dear Sir,

With reference to your advertisement regarding Pre-qualification of firms for the Supply of Drugs and Medicines (Bulk Purchase) to PIMS & SSP funded Patients for Period of 02 years advertised on _____in the _____ Newspaper, PPRA Website as well as on EPADS 2.0 of PPRA, it is requested to provide the Pre-qualification Documents.

(Tick Appropriate Box)

1. Local Manufacturers ***(Drug Medicine)*** ☐

2. Sole Agents ***(Drug Medicine)*** ☐

3. Authorized Distributors ***(Drug Medicine)*** ☐

M/s _____ hereby authorizes Mr./Ms. _____

Designation _____ CNIC No. _____

Official Email_____ Mobile No. _____

*Firm's NTN:*_____ *Firm's STN:*_____

Authorized By

Name_____ Signature _____

Designation _____ Contact No. _____

Stamp _____

GENERAL FIRM'S INFORMATION
(Manufacturer of Drug Medicine)

I.Company Profile.

1. Name of company:

Year established:

Form of company : [ ] Individual
[ ] Partnership
[ ] Corporation
[ ] Other (specify)

2. Legal status: _____

Trade registers number: _____

NTN & Sales Tax number (If applicable): _____

Mfg. License Number: (attach valid copy): _____

3. Address of Company: _____

Telephone: _____ Tele/Fax: _____

E-mail: _____

4. **Employees of the Manufacturer**

Sr No.	Category	Quantity
1.	Management	
2.	R &D	
3.	Sales	
4.	Administrative	
5.	Production and quality control	
6.	Others (specify)	
	Total	

Please attach the company organizational chart

II. QUALITY DEPARTMENT

1. Do you maintain your own quality control laboratory?

[ ] YES [ ] NO (if NO please provide details of alternate arrangements)

2. Number of specialized personnel working in your quality control, quality assurance and microbiological laboratory/ies (excluding administrative personnel). Provide their academic and professional details on a separate sheet.

Pharmacists: _____
Chemists: _____
Others : _____

3. List of Equipment installed in quality control, quality assurance and microbiological laboratory/ies for quality assurance as per BP/USP.

_____

_____

4. Are these equipment calibrated & validated.

[ ] YES [ ] NO

5. Are all raw materials completely tested prior to use or is a Certificate of Analysis accepted?

[ ] YES [ ] NO [ ] Certificate of Analysis

6. Are control samples of each batch retained?

[ ] YES [ ] NO

7. Name and title of the authorized person (s) responsible for batch release:
Name:_____

Title: _____

Experience in pharmaceuticals: _____ years

8. Name and qualification of the head of the Quality Control department:

Name: _____

Qualification: _____

Experience in pharmaceuticals: _____ Years

9. **Describe your storage facilities:**

_____

Arbitration History (if any): _____

Authorized Sole agent for Foreign Principal's Qualification (Drug Medicine)

I. Company Profile.

1. Name of company: _____

Year established: _____

Form of company : [ ] Individual
[ ] Partnership
[ ] Corporation
[ ] Other (specify)

Legal status : _____

Trade registers number : _____

NTN & Sales Tax number:

Valid sole agency agreement (attach valid copy)

2. Address : _____

Telephone : _____ Tele fax: _____

E-mail & Web : _____

Please attach the company organizational chart

3. Type of activity carried out by the company (tick the appropriate category/ies)

[ ] Manufacturer
[ ] Branded products
[ ] Generic products
[ ] Medical supplies
[ ] Other products (specify below)

(In case Sole Agent is authorized for more than one Principal then detail/s of each Principal).

4. Employees of Sole Agent:

Sr No.	Category	Quantity
1.	Management	
2.	R &D	
3.	Sales	
4.	Administrative	
5.	Production and quality control	
6.	Others (specify)	
	Total	

6. Capital value of the company (specify currency)

(a) Authorized capital: _____

(b) Paid up capital: _____

(c) Administration: _____

7. Annual sales turnover in the previous one year. Mention Private Sector and Public Sector sales separately (in Pak Rupees) (In Million)

Annual turnover	Open market sales	Public Sector Sale	Year

8. Arbitration History (if any): _____

9. Describe your storage facilities:

_____

Authorized Distributors for Qualification (Drug Medicine)

I. Company Profile.

1. Name of company: _____

Year established: _____

Form of company : [ ] Individual
[ ] Partnership
[ ] Corporation
[ ] Other (specify)

Legal status : _____

Trade registers number : _____

NTN & Sales Tax number:

Valid Distribution agreement with the Principal (attach valid copy)

2. Address : _____
Telephone : _____ Tele fax: _____
E-mail & Web : _____

3. Type of activity carried out by the Principal (tick the appropriate category/ies)

[ ] Manufacturer
[ ] Branded products
[ ] Generic products
[ ] Medical supplies
[ ] Other products (specify below)

(In case Distributor is authorized for more than one Principal then add detail of each authorized firm)

4. Employees of the Distributor:

Sr No.	Category	Quantity
1.	Management	
2.	Sales	
3.	Administrative	
4.	Distribution	
5.	Others (specify)	
	Total	

5. Capital value of the Distribution (In PKR) _____

6. Annual sales turnover in the previous Three years. Mention Private Sector and Public Sector sales separately (in Pak Rupees) (In Million)

Year	Open market sales	Public Sector Sale	Annual turnover

7. Stock Storage Capacity mention area in Sqft. _____

8. SOP's regarding Cold chain Maintenance: (Attach Copy) _____

9. Arbitration History (if any):_____

Date:_____

To

The Executive Director,
PIMS, Islamabad.

I/we, the undersigned, apply to be prequalified for the referenced Pre-qualification and declare that:

(a) I/we have examined and have no reservations to the Pre-qualification Documents issued in accordance with Instructions to Applicants (ITA).

(b) I/we, have nationalities from eligible countries, in accordance with instruction to applicant .

(c) I for any part of the application resulting from this Pre-qualification, do not have any conflict of interest;

(d) I/we for any part of the contract resulting from this Pre-qualification, have not been declared disqualified/blacklisted by any of the public organization of the Procuring Agency's country

(e) I/we understand that you may cancel the Pre-qualification process at anytime; the Pre-qualification does not bound the procuring agency to call for the bids from the prequalified firms.

(f) All information, statements and descriptions contained in the Application (online through EPADs 2.0 of PPRA) are in all respect true, correct and complete to the best of our knowledge and belief and there is no difference in information provided online and submitted in hard copy.

Signature: _____
(*signature of an authorized representative of the Applicant)*

Name: _____
(*full name of person signing the application)*

In the Capacity of: _____
(capacity of person signing the application)

Duly authorized to sign the application for and on behalf of:

Applicant's Name: _____

Address: _____

Date: ____/____ / 2026

Instructions to Applicants (ITA)		
1	**Scope of Application**	
1.1		In connection with the Invitation for Pre-qualification "as per PPRA rules 2004" the Pakistan Institute of Medical Sciences (PIMS), issues this Pre-qualification Document (PQD) to applicants interested to pre-qualifying Pharmaceutical Manufacturing Units, Authorized Distributors, and Sole Agents of Foreign Principals for Drug Medicine against the list of items/sections contained in the Pre-qualification Documents. This Pre-qualification will be concluded for PIMS.
1.2		Pre-qualification will be carried only for the items which comes under the definition of drugs under Drugs Act 1976/DRAP Act.
1.3		Department may physically verify firm's claim/s regarding submitted documents, in case any discrepancy found at any stage.
2	**Eligible Applicants**	
2.1		(a) An Applicant can be a private entity registered with FBR having NTN & STRN Registration.
2.2		(b) If Government of Pakistan prohibits commercial relations with any Country, the firms dealing with such countries are ineligible to apply.
2.3		(c) A firm declared disqualified/blacklisted/debarred by any of the public sector organization in Pakistan shall be ineligible for Pre-qualification
2.4		(d) The prequalified firms are required to participate in RFP/bidding process announced by procuring agency i.e. PIMS. In case of failure to participate, procuring agency may disqualify respective firm from Pre-qualification for the period of 02 Years and that firm will also not be entertaining for next 02 years.
3	**Tender Fee**	
3.1		The firm will deposit tender fee of Rs. 2,000/- to the Accounts Branch, of PIMS, Islamabad. ***(Upload tender receipt copy on EPADs 2.0).***
4	**Documents Comprising the Application**	
4.1		The application shall comprise the following:
		a. Application Submission Form, in accordance with Information to Applicants (ITA);
		b. Documentary evidence establishing the Applicant's eligibility to pre-qualifying, in accordance with ITA & Pre-qualification Criteria;
		c. Any other document required as specified in the Pre-qualification Documents.
		d. A Statement should be on letter head containing all information, statements and description contained in the Application (online) are in all respect true, correct and complete to the best of our knowledge and belief and there is no difference in information provided online.
5	**Signing of the Application**	Applicant shall prepare and submit the application for Pre-qualification as described in ITA & Pre-qualification Documents. The application shall be typed and shall be signed by a person duly authorized to sign on behalf of the Applicant.
6	**Notification of Pre-qualification**	Once the Pre-qualification has completed it shall be notified and communicated to all applicants in writing as well as through EPADS of PPRA & PIMS website.
7	**Contract Period**	The Pre-qualification contract shall be valid for **the period of 02 Years.**

S. No	Title	Total Estimated Quantity
1	Diclofenac Sodium 75mg/3ml Injection	345000
2	Diclofenac Sodium 50mg Tablet	1150000
3	Acetyl Salicylic Acid 300mg Tablet(Soluble)	230000
4	Ibuprofen 100mg/5ml Suspension	184000
5	Ibuprofen 400mg Tablet	924600
6	Ketorolac tromethamine 30mg/ml Injection	594550
7	Mefenamic Acid 500mg Tablet	115000
8	Nalbuphine HCl 10mg/ml Injection	157550
9	Paracetamol 120mg/5ml (60ml) Suspension	190900
10	Paracetamol +Orphenadrine 450+35mg Tablet	1150000
11	Paracetamol 1gm/100ml Infusion	690000
12	Paracetamol 500mg Tablet	5784500
13	Tramadol HCL 100mg/2ml Injection	140300
14	Tramadol HCl 50mg Capsule	46000
15	Amikacin 100mg Injection	64400
16	Amikacin 500mg Injection	69000
17	Amoxicillin +Clavulanic acid 0.6gm Injection	23000
18	Amoxicillin +Clavulanic acid 1.2gm Injection	57500
19	Amoxicillin + Clavulanic acid 500mg+125mg tablet	506000
20	Amoxicillin + Clavulanic acid 250mg+62.5mg Suspension	73600
21	Amoxicillin 500mg Capsule	920000
22	Amoxicillin + Salbactum 1.5g Injection	11500
23	Amoxicillin as Trihydrate 250mg/5ml Syrup	69000
24	Ampicillin 500mg Injection	80500
25	Azithromycin 500mg Injection	13800
26	Azithromycin 200mg Suspension	46000
27	Azithromycin 250mg Capsule	345000
28	Moxifloxacin 400mg Injection	23000
29	Benzathine Penicillin 1.2MU Injection	32200
30	Benzyl Pencillin 1000000 IU Injection	32200
31	Cefixime 100 mg/5ml Suspension	36800
32	Cefixime 400mg Capsule	46000
33	Cefotaxime 500mg Injection	23000
34	Ceftazidime 1gm Injection	29900
35	Ceftriaxone 1gm injection	939550
36	Cefuroxime Sodium 750mg injection	80500
37	Cefepime 1gm with with Water for injection	23000
38	Cephradine 500mg with Water for injection	80500
39	Cephradine 500mg Capsule	368000
40	Cephradine 250mg/5ml Suspension	23000
41	Ciprofloxacin 200mg /100ml Injection	80500
42	Ciprofloxacin 500mg Tablet	345000
43	Gentamycin Sulphate 80mg Injection	55200
44	Imipenem + Cilastatin Sodium 500mg Injection	41400
45	Levofloxacin 500mg/100ml Injection	37950
46	Levofloxacin 500mg Tablet	345000
47	Meropenem 500mg Injection	23000
48	Meropenem 1g Injection	211600
49	Linezolid 600mg Infusion	46230
50	Piperacillin+Tazobactam 2.25 gm with Water for injection	20700
51	Piperacillin+Tazobactamwith 4.5gm with Water for injection	134550
52	Piperacillin+Tazobactamwith 4.5gm (Lyophilized)	88550

53	Salbactam + Cefoperazone injection 1gm Injection	101200
54	Salbactam + Cefoperazone 2gm Injection	241500
55	Ticarcillin clavulanic acid 3.2g injection	34500
56	Vancomycin HCl 1gm Injection	129950
57	Colistimethate Sodium 2 million i.u Injection	44850
58	Tigecycline 50mg injection	18400
59	Tobramycin 20mg Injection	11500
60	Metronidazole 500mg/ 100ml Infusion	460000
61	Metronidazole 400mg Tablet	1334000
62	Metronidazole 200mg/5ml (60ml) Suspension	23000
63	Mebendazole 100mg/5ml Suspension	34500
64	Clotrimazole (vaginal) 100mg Tablet	4600
65	Nystatin 30 ml Drops 100000 unita/ml Drop	13800
66	Fluconazole 150 mg cap	11500
67	Fluconazole 200mg injection	6900
68	Voriconazole 200 mg suspension	6900
69	Voriconazole 100 mg Injection	4600
70	Amphotericin 50mg injection	1150
71	Artemether +Lumefantrine 20mg+120mg Tablet	23000
72	Artemether 80mg Injection	460
73	Phenylephrine HCl 10mg Injection	23230
74	Isosorbide Dinitrate 10mg Injection	69000
75	Sodium nitroprusside 50mg Injection	1150
76	Dobutamine HCl 250mg Injection	34500
77	Dopamine HCl 200mg Injection	23000
78	Metoprolol tartorate 1mg Injection	6900
79	Labetalol 50mg Injection	34500
80	Methyldopa 250mg Tablet	11500
81	Amlodipine 5mg Tablet	575000
82	Propranolol 10mg tab	69000
83	Atenolol 50mg Tablet	115000
84	Atorvastatin 10mg Tablet	1150000
85	Captopril 25mg Tablet	230000
86	Carvedilol 6.25mg Tablet	345000
87	Apixaban 2.5mg Tablet	230000
88	Glyceryl Trinitrate 0.5mg S/L Tablet	46000
89	Glyceryl Trinitrate 2.6mg Tablet	460000
90	Glyceryl Trinitrate 10ml Injection	2300
91	Lisinopril 10mg Tablet	230000
92	Losartan Potassium 50mg Tablet	345000
93	Losartan Potassium + Hydrochlorthiazide 50mg + 12.5mg Tablet	345000
94	Amlodopine besylate + Valsartan 5mg + 80mg Tab	1150000
95	Amlodopine besylate + Valsartan 5mg + 160mg Tab	460000
96	Amlodopine besylate + Valsartan 10mg + 160mg Tab	230000
97	Metoprolol tartorate 50mg Tablet	345000
98	Metoprolol Tartrate 25mg Tablet	230000
99	Rosuvastatin 10mg Tablet	1150000
100	Nebivolol 2.5mg Tablet	230000
101	Nebivolol 5mg Tablet	230000
102	Bisoprolol (Fumarate) 2.5mg Tablet	1150000
103	Bisoprolol (Fumarate) 5mg Tablet	690000
104	Ivabradine 5mg Tablet	230000
105	Amiodarone 150mg Injection	46000
106	Nifedipine R 20mg Tablet	23000

107	Lanoxin 50mcg Syrup	1150
108	Lanoxin 100mcg Injection	1150
109	Tenecteplase Injection	460
110	Enoxaparin 40mg Injection	69000
111	Enoxaparin 60mg Injection	46000
112	Heparin Sodium 5000IU/ml Injection	29670
113	Streptokinase 1.5MIU Injection	4600
114	Phytomenadion B.P 2mg/ml Injection	161000
115	Phytomenadion B.P 10mg/ml Injection	115000
116	Tranexamic Acid 500mg Capsule	34500
117	Tranexamic Acid 500mg Injection	233680
118	Aspirin (Enteric coated) 75mg Tablet	1150000
119	Clopidogrel 75mg Tablet	1150000
120	Loprin+Clopidogrel 75+75mg Tablet	1150000
121	Aspirin 300mg Tablet	115000
122	Spironolactone 40mg Tablet	161000
123	Frusemide 20mg/2ml Injection	382950
124	Mannitol 20% 500ml Infusion	52900
125	Betamethasone ointment 1% 30g Cream	11500
126	Clobetasol propionate 0.05% cream	23000
127	Mupirocin 2% 5g ointment	23000
128	Fluorometholone, Sodium Cromoglycate 4.0%	2300
129	Carboxymethylcellulose 10ml Eye Drop	2300
130	Polymyxin B Sulphate 10000units + Bacitracine Zinc 500units Eye ointment	65550
131	Xylometazoline 15ml Nasal Drop	34500
132	Gentamycin 0.3% Ear Drop	6900
133	Dexamethasone +Tobramycin 0.1%+0.3% Eye Drop	6900
134	Fusidic Acid 2% 15gm Cream	41400
135	Permethrine 5% Lotion	46000
136	Polymyxin + Bacitracin +Neomycin + Lidocain USP 10gm Ointment	143750
137	Polymyxin B Sulphate BP 10000 units/gm, Bacitracin Zinc 500 units/gm.	63250
138	Silver Sulphazine Cream 1% 250gm jar Cream	11500
139	Protamin Sulphate 10mg Injection	23000
140	Naloxone HCl 0.4mg/ml Injection	15065
141	Parlidoxime methysulphate 200mg injection	23000
142	Flumazenil 0.1mg/ml Injection	230
143	Octreotide acetate 0.1mg/ml Injection	46000
144	Octreotide acetate 0.5mg/ml Injection	46000
145	Drotaverine HCL 40mg Injection	391000
146	Omeprazole 40mg Injection	972900
147	Pantoprazole 40mg Injection	460000
148	Pantoprazole 40mg Tablet	1150000
149	Omeprazole 20mg Capsule	5060000
150	Omeprazole 40mg Capsule	1150000
151	Famotidine 40mg Tablet	920000
152	Domperidone 10mg Tablet	575000
153	Domperidone 1mg/ml Syp	11500
154	Zinc Sulphate 60ml Syrup	23000
155	Lactulose 3.35gm/5ml Syrup (120ml)	34500
156	Erythropoietin vial 2000iu/ml Injection	9200
157	Erythropoietin vial 4000iu/ml Injection	34500
158	Iron Sucrose 20mg/ml Injection	41400
159	Iron Polymaltose Complex eq to Elemental Iron + Folic Acid 120ml Syrup	25300
160	Calcium carbonate + Vit-D Tablet	1150000

161	Calcium Gluconate 10mg Injection	139380
162	Calcium Phosphate 210mg/5ml +Vit D3 350 units/5ml Syp (120ml)	46000
163	Ferrous Sulphate (Blister Pack) Tablet	460000
164	Mecobalamine 500mcg Tablet	575000
165	Multivitamin (Blister Pack) Tablet	575000
166	Ferrous Sulphate Syrup	23000
167	Pyridoxin hydrochlorid 100mg Tablet	11500
168	Vitamin B Complex Syrup	34500
169	Sodium Valproate 500mg Injection	46000
170	Sodium Valproate 500mg Tablet	460000
171	Carbamazepine 200mg Tablet	184000
172	Sodium Valprate 250mg/5ml (120ml) Syrup	23000
173	Carbamazepine 100mg/5ml Syrup	11500
174	Phenytoin Sodium 250mg Injection	11500
175	Levetiracetam 500mg Injection	163300
176	Levetiracetam 500mg Tablet	230000
177	Citicoline 250mg/ml injection	23000
178	Lacosamid 200mg Injection	11500
179	Acetazolamide 250mg tablets	11500
180	Sevoflurane 250ml INH	2760
181	Isoflurane Liquid 250ml	3220
182	Bupivacaine HCl 0.5% 10ml Injection	31280
183	Spinal 0.5% Bupivacaine HCL in dextrose 4ml Injection	92460
184	Glycopyrrolate 0.2mg/ml Injection	12420
185	Glycopyrrolate+Neostigmine Methylsulphate 0.5mg+2.5mg/ml Injection	40020
186	Ketamine HCl 50mg/ml Injection	70150
187	Lignocaine HCl 2% 10ml Injection	147200
188	Lignocaine HCl 2% +Adrenaline 10ml Injection	100280
189	Propofol MCT and LCT 10mg/ml Injection	85100
190	Midazolam HCl 5mg Injection	172500
191	Lignocain 4% 50ml Solution	23000
192	Lignocain 2% 15gm Gel	143750
193	Albumin human 20% 100ml Infusion	3450
194	Albumin human 20% 50ml Infusion	1150
195	Parentral Amino Acid 5% 500ml Infusion	13800
196	Polygeline 3.5% 500ml Infusion	16100
197	Succinylated Fluid Gelatin 4% 500ml Infusion	16790
198	Long chain Fat emulsion Soyabean oil + Medium chain triglyceride 250ml	16100
199	Parentral Amino Acid 10% 500ml Infusion	23000
200	Potassium Chloride 7.45% 20ml Injection	140300
201	Infusion 4.3% Glucose with 0.18% Sodium Chloride 500ml Infusion Without IV SET	73600
202	Infusion 0.45% Sodium Chloride with 5% Glucose 500ml Infusion Without IV SET	73600
203	Infusion Dextrose 5% With 0.9% Sodium Chloride 1000ml Infusion without iv set	27600
204	Infusion 10% Dextrose Water 1000ml Infusion without IVSET	25300
205	Dextrose Water 25% 1000ml Infusion	46000
206	Dextrose Water 25% 20ml Infusion	220800
207	Infusion Dextrose water 5% 1000ml Infusion without IVSET	34500
208	Infusion Dextrose water 5% 500ml Infusionwithout IVSET	23000
209	Reduced osmolarity Oral Rehydration Salt Sachet	69000
210	Ringer lactate 1000ml Infusion without iv set	351900
211	Ringer lactate 500ml Infusion without iv set	34500
212	Sodium Bicarbonate 8.4% 20ml Infusion	16100
213	Normal 0.9% saline 100ml Infusion without iv set	1667500

214	Normal 0.9% saline 1000ml Infusion without iv set	851000
215	Normal 0.9% saline 500ml Infusion without iv set	92000
216	Atracurium Besylate 30mg/3ml Injection	34500
217	Atracurium Besylate 50mg/5ml Injection	194810
218	Dexmedetomidine 200mcg/2ml Injection	51175
219	Suxamethonium Chloride 100mg Injection	18400
220	Salbutamol sulphate 2mg/5ml 60ml Syrup	23000
221	Terbutaline 0.3mg/5ml 60ml Syrup	34500
222	Ipratropium Bromide 0.25mg/ml (20ml) Nebules	57960
223	Salbutamol Nebulization Solution	23000
224	Beclomethasone Dipropionate 0.8mcg /2ml Nebuliser suspension	46000
225	Thyroxine sodium anhydrous 50mcg Tablet	230000
226	Pheniramine Maleate 25mg/ml Injection	600300
227	Loratidine 10mg Tablet	345000
228	Fexofenadine 120mg Tablet	690000
229	Chlorphenramine Maleate 2mg/5ml Syp	115000
230	Dimenhydrinate 50mg/ml Injection	345000
231	Dimenhydrinate 12.5mg/4ml Syrup	11500
232	Montelukast 10mg Tablet	690000
233	Montelukast 5mg Tablet	230000
234	Cetirizine 10mg Tablet	2300000
235	Cetirizine 5mg/5ml 60ml Syrup	184000
236	Ondansetron 8mg Injection	487600
237	Ondansetron 4mg/5ml Syrup	23000
238	Dexamethasone 4mg/ml Injection	598000
239	Hydrocortisone 100mg Injection	296700
240	Hydrocortisone 250mg Injection	46000
241	Prednisolone 5mg Tablet	345000
242	Methylprednisolone 500mg Injection	6900
243	Hydrogen Peroxide Solution 6% 450ml Solution	5635
244	NPH Insulin (Human) 100IU/ml 10ml Injection	2645
245	Regular Insulin (Human) 100IU/ml 10ml Injection	4945
246	Insulin Regular 30% + Isophane Insulin 70% (Human) 100IU/ml 10ml Injection	11960
247	Glimepiride 2mg Tablet	230000
248	Metformin HCl 500mg Tablet	690000
249	Sitagliptin + Metformin HCl 50/500 Tablet	690000
250	Sitagliptin + Metformin HCl 50/1000 Tablet	690000
251	Tetanus Toxoid Vaccine 0.5ml (Single Dose) Injection	218960
252	Anti-D Immunoglobulin 300mcg Injection	690
253	Anti snake Venom (Lyophlised) Dried indian cobra + Dried Comman Krait + Dried Russell's Viper + Dried Saw scaled Viper Injection	6900
254	I/M Rabies Vaccine for human use (WHO approved) Single dose	13800
255	Norepinephrine bitartrate 1mg/ml Injection	119600
256	Procyclidine 5mg Tablet	575000
257	Adrenaline HCl 1mg/ml Injection	166750
258	Atropine 1mg/ml Injection	140300
259	Oxytocin 5iu/ml Injection	345000
260	Gadopentetate Dimeglumine (Contrast Media for MRI) Injection	7130
261	Non Ionic contrast Media for CT 100ml Injection	7130
262	Non Ionic contrast Media for CT 50ml Injection	2530
263	Diazepam 10mg/2ml Injection	92230
264	Alprazolam 0.5mg Tablet	345000
265	Ecitalopram 10mg Tablet	575000
266	Risperidone 2mg Tablet	575000

267	Haloperidol 5mg/2ml Injection	115000
268	Tamsulocin 0.4mg Capsule	690000
269	Acyclovir 500mg/2ml Injection	36800
270	Lung Surfactent Bovine Lipid extract 3ml suspension	1840
271	Lung Surfactent Bovine Lipid extract 5ml suspension	690
272	Poractant alfa 1.5ml intratracheal suspension	1150
273	Poractant alfa 3ml intratracheal suspension	1150
274	Methotrexate 2.5mg Tablet	69000
275	Methotrexate 10mg Tablet	69000
276	Methotrexate 250mg injection	2300
277	Povidone Iodine Scrub 450ml	21850
278	Povidone Iodine Solution 10% 450ml	37950
279	I/V Set Blister Pack (Length 120cm or More)	3806500
280	Paracetamol 250mg/5ml Syrup	115000
281	Ibuprofen 100mg/5ml Suspension	46000
282	Linezolid 100mg/5ml Syrup	2300
283	Water for Injection 5ml	713000
284	Caffeine Citrate 20mg/ml Injection	1150
285	Magnesium Sulphate 500mg Injection	127190
286	Misoprostol 200mcg Tablet	69000
287	Thiocolchicoside 4 mg Tablet	115000
288	Famciclovir 250mg Tablet	69000
289	Diphtheria Anti toxin Injection	2300
290	Promethazine HCL 25mg Injection	115000
291	liquid paraffin 450ML	13800
292	liquid glycerin 450ML	6900
293	Ketotifen 0.2mg/ml syrup	69000
294	Ketotifen 1mg Tablet	115000
295	Daprodoustat 4mg	4600
296	Monobasic & Diabasic Sodium Phosphate Rectal Solution iabasic S	24150
297	Baby infant formula (NICU & PICU) 400gm Powder	11500
298	Baby milk low birth weight formula (NICU & PICU) 400gm Powder	11500
299	Balanced isotonic Electrolyte Solution, Lactate free 1000ml	4600
300	Ferric Carboxy maltose 500mg/10ml Injection	46000
301	Factor VIII Injection	1150
302	Deferasirox 500 mg	46000
303	Deferasirox 250 mg	46000
304	Deferasirox 180 mg	46000
305	Deferasirox 360 mg	46000
306	Isophane Insulin 70% (Human) 100IU/ml 10ml Injection	11500
307	Aztreonam 1g Injection	46000
308	Aztreonam 2g Injection	46000
309	Daptomycin 500mg injection	2300
310	Caspofungin 50mg injection	2300
311	Gencyclovir 500mg Injection	2300
312	Voriconazole 200mg Injection	2300
313	Ceftazidime + Avibactum 2.5g Injection	23000
314	Valganciclovir 450 mg Injection	2300
315	Minocycline 100 mg Injection	2300
316	Minocycline 100 mg Tablet	23000
317	l-ornithine l-aspartate Injection	115000
318	Terlipressin 1mg injection	23000
319	Citacholine 250mg injection	46000
320	Ceftaroline 600mg Injection	11500

321	Glargine 100 unit/ml Pen	23000
322	Acetylcysteine 200mg water soluble granules	69000
323	Tacrolimus 1mg capsules	46000
324	Tacrolimus 1mg/ml Injection	11500
325	lornoxicam 8mg injection	23000
326	Cefazolin 1g injection	23000
327	Acyclovir 400mg Tablet	230000
328	Nifedipine 30 mg Tablet	23000
329	Tropicamide ophthalmic solution 1%	23000
330	Moxifloxacin 0.5% Eye Drop	46000
331	Ciprofloxacin 0.3% ophthalmic solution	46000
332	Spironolactone 25 mg tablet	230000
333	Amphotericin inj 50mg	11500
334	Apixaban 5 mg tablet	1150000
335	Dapagliflozin propanediol 10 mg tablet	46000
336	Nimodipine 30mg tablets	46000
337	Alfacalcidol 1 mcg Tablet	460000
338	Clindamycin 600mg injection	11500
339	Sodium bicarbonate 300mg tablet	1150000
340	Cream of magnisium with liq parrafin 120ml Syp	115000
341	Carbocisteine 250mg /5m Syp	11500
342	Acefylline piperazine and diphenhydramin hcl 120ml Syp	115000
343	Miconazole 2% Oral gel	69000
344	Ferric carboxymaltose 50mg injection	46000
345	Filgrastrim 300mcg injection	11500
346	Artesunate 30mg injection	11500
347	Gabapentin 100mg capsules	115000
348	Gabapentin 50mg capsules	115000
349	Febuxostat 40mg tablet	46000
350	Itopridehydrochloride 50mg tablet	46000
351	levopride 25 tablet	230000
352	Hydralazine 25mg tablet	46000
353	Azathioprine 50 mg tablet	46000
354	Pottacium chloride Oral solution USP Syp	46000
355	Monobasic sodium phosphate enema	115000
356	Magnesium sulphate injection 2ml Injection	115000
357	Fostoin Sodium 150mg/2ml injection	11500
358	Artesunate 60mg injection	11500
359	Fluconazole 5mg/ml Syrup	34500
360	Sildenafil citrate 50mg Syrup	11500
361	Ibutilide Injection	230
362	Sotalol 150mg/10ml Injection	230
363	Flecainide 150mg/15ml Injection	230
364	Disopyramide 100mg/5ml Injection	230
365	Sotalol 150mg/10ml Injection	230
366	Propafenone 70mg/20ml Injection	230
367	Isoprenaline 1mg/5ml Injection	1150
368	Vitamin B1+B6+B12 Injection	115000
369	Minocycline hydrochloride 100 mg capsule	69000
370	Doxyccucline HCL 100 mg capsule	69000
371	Linezolid 600mg Tablet	115000
372	Neomercazole 5mg Tablet	230000
373	Naproxen 500 mg tablet	230000
374	5% Dextrose and Electrolyte 500ml Inf	69000

375	Gabapentin tablets 100mg Capsule	115000
376	Cyclosporin Syrup	1150
377	Cyclosporin 25mg Tablet	23000
378	Ursodeoxycholic acid Syrup	23000
379	cefixime 400 mg Capsule	115000
380	Calcium Acetate Tablet	115000
381	Oxythazine + Almunium + Magnisium 120ml SUSP	230000
382	Betahistine dihydrochloride 16 mg Tablet	115000
383	Sucralfate Oral Susp	11500
384	Valganciclovir 450 mg Injection	1150
385	Vasopressin 20units/ml injection	1150
386	Vitamin D3 Injection	46000
387	Vitamin D3 Capsule	460000
388	Cefuroxime Sodium 250mg injection	115000
389	salbutamol inhaler 100mcg/dose	11500
390	Salmeterol + fluticasone 25/125 mcg Inhaler	11500
391	beclometasone dipropionate 100/6 Inhaler	11500
392	Eltrombopag 50mg Tablet	2300
393	Dextrose 5% With 0.9% Sodium Chloride 500ml Inf	46000
394	lobetasol propionate cream 0 05 %	46000
395	Paroxetine hcl 20mg tablet	46000
396	Tizanidine 2mg Tablet	230000
397	Esomeprazole 40mg Cap	1150000
398	Activated Charcoal Sachet	23000
399	Albendazole 400 mg Tablet	115000
400	Amitriptyline 10 mg Tablet	23000
401	Baclofen 10 mg Tablet	23000
402	Bisacodyl 5 mg Tablet	23000
403	Bosentan 62.5 mg Tablet	23000
404	Cholestyramine Sachet	23000
405	Cinacalcet 30 mg Tablet	23000
406	Clarithromycin 500 mg Tablet	23000
407	Clobazam 10 mg Tablet	23000
408	Clozapine 25 mg Tablet	23000
409	Co-trimoxazole 160/800 mg Tablet	23000
410	Dabigatran 150 mg Capsule	23000
411	Duloxetine 30 mg Capsule	23000
412	Erythromycin 500 mg Tablet	23000
413	Finerenone 10 mg Tablet	23000
414	Fludrocortisone 100 mcg Tablet	23000
415	Influenza Vaccine	2300
416	Itraconazole 100 mg Capsule	23000
417	Ivermectin 6 mg Tablet	23000
418	Lamotrigine 25 mg Tablet	23000
419	Loperamide 2 mg Capsule	23000
420	Metoclopramide 10 mg Tablet	23000
421	Midodrine 5 mg Tablet	23000
422	Nitrofurantoin 100 mg Capsule	23000
423	Olanzapine 5 mg Tablet	23000
424	Oxcarbazepine 300 mg Tablet	23000
425	Pneumococcal Vaccine	4600
426	Prasugrel 10 mg Tablet	46000
427	Pregabalin 150 mg Capsule	46000
428	Pregabalin 50 mg Capsule	46000

429	Propylthiouracil 50 mg Tablet	46000
430	Quetiapine 100 mg Tablet	46000
431	Rifaximin 550 mg Tablet	46000
432	Rivaroxaban 20 mg Tablet	46000
433	Sacubitril/Valsartan 100 mg Tablet	46000
434	Sildenafil 50 mg Tablet	46000
435	Terbinafine 250 mg Tablet	46000
436	Ticagrelor 90 mg Tablet	46000
437	Tinidazole 500 mg Tablet	46000
438	Topiramate 100 mg Tablet	46000
439	Topiramate 25 mg Tablet	46000
440	Topiramate 50 mg Tablet	46000
441	Warfarin 5 mg Tablet	46000
442	4-Factor Prothrombin Complex Concentrate	23
443	Tetanus Immunoglobulin	230
444	Rabies Immunoglobulin	1150
445	Phentolamine 10mg injection	1150
446	Calcitonin 100iu Injection	1150
447	Ertapenem 1g injection	1150
448	Pamidronate 30mg Injection	1150
449	Ceftolozane/Tazobactam 1.5g Injection	1150
450	Meropenem/Vaborbactam 2g Injection	1150
451	Posaconazole IV 300mg	1150
454	IVIG 5g Vial	1150
455	Ringer Acetate 1000ml	1150
456	Tirofiban 50ml Injection	1150
457	Cisatracurium 10mg Injection	1150
458	Etomidate 20mg Injection	1150
459	Milrinone 10mg Injection	1150
460	Procainamide 1g Injection	1150
461	Rocuronium 50mg Injection	1150
462	Sugammadex 200mg Injection	1150
463	Vecuronium 10mg Injection	1150
464	Desmopressin Injection	1150
465	Quinine 300mg Injection	2000
466	Aminophylline 250mg injection	20000
467	Teicoplanin 400mg Injection	5000
468	Fosfomycin 1g IV	1150
469	Deferoxamine 500mg Injection	1150
470	Budesonide 0.5mg /2ml Respules	1150
471	Granisetron 3mg Injection	1150
472	Metoclopramide 10mg Injection	50000
473	Verapamil 5mg Injection	2000
474	Lorazepam 4mg Injection	2000