

**ALL INDIA INSTITUTE OF MEDICAL SCIENCES,
ANSARI NAGAR, NEW DELHI-110029, INDIA
1ST FLOOR, STORE SECTION (HOSPITAL),
NEAR BLOOD BANK (Main)**

TENDER ENQUIRY DOCUMENT



Advertised Tender Enquiry No. : 39/H/Drugs/2025-26

Rate Contract items : Purchase of Drugs/Medicines/I.V. Fluids

Period of Rate Contract : 02 Years Rate Contract Basis

SECTION-I



ALL INDIA INSTITUTE OF MEDICAL SCIENCES ANSARI NAGAR, NEW DELHI-110 029

NOTICE INVITING TENDERS (NIT)

Advertised Tender Enquiry No. : **39/H/Drugs/2025-26**

On behalf of Director, AIIMS, Ansari Nagar, New Delhi-110 029, online bids are invited in two bid system (Techno-Commercial Bid and Financial Bid) from eligible and qualified firms/manufacturer for supply of following Goods for conclusion of Rate Contract for a period of 02 Years: -

S. No.	Brief Description of Goods	Amount of Bid Security/EMD (INR)
1.	Purchase of Drugs/Medicines/I.V. Fluids on 02 years rate contract basis	Rs.50,000/-

CRITICAL DATE SHEET

Published Date & Time	03.02.2026 at 04:00 PM
Bid Document Download/Sale Start Date	03.02.2026 at 04:00 PM
Bid Submission Start Date & Time	13.02.2026 at 04:00 PM
Bid Submission End Date & Time	27.02.2026 at 04:00 PM
Bid Opening Date & Time	28.02.2026 at 04:00 PM

Instructions:

1. Bids shall be submitted online only at CPPP website:
<https://eprocure.gov.in/eprocure/app>.
2. The Bidder shall download the Tender Enquiry Document directly from the websites <https://eprocure.gov.in/eprocure/app> and shall not tamper/modify it including downloaded Price Bid template in any manner. In case if the same is found to be tampered/modified in any manner, Tender/Bid will be summarily rejected and EMD would be forfeited.
3. The complete bidding process is online. Bidders should be in possession of valid Digital Signature Certificate (DSC) of class III for online submission of bids. Prior to bidding DSC need to be registered on the website mentioned above.
4. Bidders are advised to follow the instructions provided in the “Instructions for Online Bid Submission” in Para No. 11 of GIB of Tender Enquiry Document.
5. Bidders are advised to visit this website regularly to keep them updated, for any changes / modifications in the Tender Enquiry Document.
6. Intending bidder are advised to visit CPPP website <https://eprocure.gov.in/eprocure/app> regularly till closing date of submission of bid, for any corrigendum.
7. The documents to be submitted in their bid may be scanned with 100 dpi with black and white option which helps in fast uploading.
8. The EMD / Bid Security of **Rs. 50000/-** shall be deposited through Bank Guarantee / Demand Draft / FDR drawn in favor of the AIIMS MAIN GRANT. The original Earnest Money / Bid Security must be submitted to *Stores Officer (Hospital), Hospital Stores, 1st Floor, M.S. Office, Near Blood Bank, AIIMS, New Delhi-110 029* till “Bid Submission End Date & Time” as mentioned in “Critical Date Sheet” failing which the bid shall be summarily rejected.

SECTION - II
GENERAL INSTRUCTIONS TO BIDDERS (GIB)

A. PREAMBLE

1. Definitions and Abbreviations

1.1 The following definitions and abbreviations, which have been used in these documents shall have the meanings as indicated below:

1.2 Definitions:

- i) "Purchaser" means the organization i.e. AIIMS/Center/Hospital/Department/Sections purchasing goods as incorporated in the Tender Enquiry Document.
- ii) "Bid" means Quotation / Tender received from a Firm / Tenderer / Bidder.
- iii) "Bidder" means Tenderer/ the Individual or Firm submitting Bids / Quotation / Tender
- iv) "Supplier" means the individual or the firm supplying the goods as incorporated in the Rate Contract/Purchase Order.
- v) "Goods" means all articles, material, commodity, livestock, furniture, fixtures, raw material, spares, instruments, machinery, equipment, vehicles, medicines, assemblies, sub-assemblies, accessories, intangible products like software, technology transfer, licenses, patents or other intellectual properties purchased or otherwise acquired for the use of Government but excludes books, publications, periodicals, etc. for a library. The term 'goods' also includes works and services which are incidental or consequential to the supply of such goods, such as, transportation, insurance, installation, commissioning, training and maintenance.
- vi) "Services" means services allied and incidental to the supply of goods, such as transportation, installation, commissioning, provision of technical assistance, training, after sales service, maintenance service and other such obligations of the supplier covered under the Rate Contract.
- vii) "Bid Security" (BS) means Earnest Money Deposit / monetary or financial guarantee to be furnished by a bidder along with its tender.
- viii) "Contract" means Rate Contract/Purchase Order which means the written agreement entered into between the purchaser and the supplier, together with all the documents mentioned therein and including all attachments, annexure etc. therein.
- ix) "Performance Security" means monetary or financial guarantee to be furnished by the successful bidder for due performance of the Rate Contract/Purchase Order placed on it. Performance Security is also known as Security Deposit.
- x) "Consignee" means the Center/Hospital/Department/Sections /person to whom the goods are required to be delivered as specified in the Purchase Order.
- xi) "Specification" also called Technical Specifications means the document/standard that prescribes the requirement with which goods has to conform.
- xii) "Inspection" means activities such as measuring, examining, testing, gauging one or more characteristics of the product and comparing the same with the specified requirement mentioned in the Rate Contract/Purchase Order to determine conformity.
- xiii) "Day" means calendar day.

1.3. Abbreviations:

- i) "ATE" means Advertised Tender Enquiry
- ii) "NIT" means Notice Inviting Tenders.
- iii) "GIB" means General Instructions to Bidders
- iv) "SIB" means Special Instructions to Bidders
- v) "GCC" means General Conditions of Contract
- vi) "SCC" means Special Conditions of Contract
- vii) "DP" means Delivery Period
- viii) "BG" means Bank Guarantee
- ix) "GST" means Goods & Service Tax
- x) "RC" means Rate Contract

2. Introduction

2.1 The AIIMS is the premier multi-disciplinary super specialty health sciences institution of India. It was established in 1956 by an Act of Parliament. AIIMS has a trinity of mission, which is medical education, research and patient care. It has around 2400 indoor beds with over 2.5 lakhs admissions per annum and an annual out-patient attendance of around 40,00,000 patients. The All India Institute of Medical Sciences (AIIMS) is catering Drugs/Medicines/I.V. Fluids to all E.H.S. patients, all essential drugs and I.V. Fluids to indoor patients. The list of Drugs/Medicines/I.V. to all E.H.S. patients, all essential drugs and I.V. Fluids to indoor patients. The list of Drugs/Medicines/I.V. Fluids required by AIIMS, is enclosed herewith for your information/reference (enclosed at Annexure-A).

2.2 This tender is for the purpose for executing rate-contract for supply of medicines at whole of the AIIMS (including all centres viz. CT & NS centre, Dr. BRA IRCH, NDDTC Ghaziabad, Rural Health Centre Ballabhgarh, JPNATC, DR. RPC and Main Hospital).

2.3 The Purchaser has issued these Tender Documents for purchase of goods as mentioned in Section – VI – "Schedule of Requirements", which also indicates, *inter alia*, the required delivery schedule, terms and place of delivery.

2.4 This section (Section II - "General Instructions to Bidders") provides the relevant information as well as instructions to assist the prospective bidders in preparation and submission of bids. It also includes the mode and procedure to be adopted by the bidder for receipt and opening as well as scrutiny and evaluation of bids and subsequent placement of Rate Contract/Purchase Order.

2.5 The bidder shall also read the Special Instructions to Bidders (SIB) related to this purchase, as contained in Section III of these documents and follow the same accordingly. Whenever there is a conflict between the GIB and the SIB, the provisions contained in the SIB shall prevail over those in the GIB.

2.6 Before formulating the bid and submitting the same to the purchaser, the bidder should read and examine all the terms, conditions, instructions, etc. contained in the Tender Document. Failure to provide and/or comply with the required information, instructions etc. incorporated in these Tender Documents may result in rejection of its Bid.

2.7 The rates quoted, approved and accepted by the Director, AIIMS shall be valid for **two years** from the date of signing of the agreement deed (**extendable up-to one year on mutual agreement, if required**).

2.8 **The tenders are to be submitted by the manufacturers/sole importer only. Tenders quoted by suppliers on behalf of manufacturers will not be entertained even if they are authorized by the manufacturers.** However, manufacturers can give authority letter to the supplier / distributor / stockiest for the purpose of making supplies, raising bills, collecting payment etc. only after selection in the tender. **In such cases, the manufacturer has to accept**

responsibility for any lapse on the part of the distributor/supplier and an undertaking to this effect from the manufacturer will have to be submitted. Failure to submit such an undertaking will lead to rejection of authorization and manufacturer will have to supply drugs directly. This authorization should be valid for the entire duration of the contract. **No change in the authorized supplier/distributor will be allowed during the rate contract period. Different distributors of a manufacturer for different Centers/Hospital will not be allowed. Sub authorization further to any other agent for delivery of the goods or for raising bills/collecting payment etc. will not be accepted.**

3. Availability of Funds

3.1 Expenditure to be incurred for the proposed purchase will be met from the funds available with the purchaser/consignee.

4. Language of Bid

4.1 The bid submitted by the bidder and all subsequent correspondence and documents relating to the bid exchanged between the bidder and the purchaser, shall be written in the English language. However, the language of any printed literature furnished by the bidder in connection with its bid may be written in any other language provided the same is accompanied by an English translation and, for purposes of interpretation of the bid, the English translation shall prevail.

5. Bid Expense

5.1 The bidder shall bear all costs and expenditure incurred and/or to be incurred by it in connection with its bid including preparation, uploading of its bid and for subsequent processing the same. The purchaser will, in no case be responsible or liable for any such cost, expenditure etc regardless of the conduct or outcome of the Tender process.

B. TENDER ENQUIRY DOCUMENT

6. Content of Tender Enquiry Document

6.1 In addition to Section I – “Notice Inviting Tender” (NIT), the Tender Enquiry Document includes:

- Section II – General Instructions to Bidders (GIB)
- Section III – Special Instructions to Bidders (SIB)
- Section IV – General Conditions of Contract (GCC)
- Section V – Special Conditions of Contract (SCC)
- Section VI – Schedule of Requirements
- Section VII – Specifications
- Section VIII – Qualification Criteria
- Section IX – Tender Acceptance Form
- Section X – Price Schedules (BoQs)
- Section XI – Bank Guarantee Form for Bid Security
- Section XII – Bank Guarantee Form for Performance Security
- Section XIII – Rate Contract Forms
- Section XIV – Performa of Consignee Receipt Certificate
- Section XV – Performa of Final Consignee Acceptance Certificate
- Section XVI – List of items quoted
- Section XVII – Performa to be filled by the tenderer
- Section XVIII – Manufacturing & Marketing Certificate
- Section XIX – Production Capacity Assessment Certificate

6.2 The relevant details of the required goods, the terms, conditions and procedure for Tender, bid evaluation, placement of Rate Contract/Purchase Order, the applicable contract terms and, also, the standard formats to be used for this purpose are incorporated in the above-mentioned documents. The interested bidders are expected to examine all such details etc to proceed further.

7. **Corrigendum to Tender Enquiry Document**

7.1 At any time prior to the deadline for submission of bids, the purchaser may, for any reason deemed fit by it, modify the Tender Enquiry Document by issuing suitable Corrigendum to it

7.2 Corrigendum will be notified through <https://eprocure.gov.in/eprocure/app> only.

7.3 In order to provide reasonable time to the prospective bidders to take necessary action in preparing their bids as per the amendment, the purchaser may, at its discretion extend the deadline appropriately for the submission of bids and other allied time frames, which are linked with that deadline.

8. **Clarification of Tender Enquiry Document**

8.1 A bidder requiring any clarification or elucidation on any issue of the Tender Enquiry Document may take up the same with the purchaser through CPP Portal only. The purchaser will respond through CPP Portal to such request provided the same is uploaded within the time schedule mentioned in "Critical Date Sheet".

C. PREPARATION OF BIDS

9. **Documents Comprising the Bid**

9.1 The **Two Bid System**, i.e. "Techno – Commercial Bid" and "Price Bid" prepared by the bidder shall comprise the following:

A) **Techno – Commercial Bid (Un-priced Bid)**

i) Scanned copy of "EMD/Bid Security" furnished in accordance with GIB alternatively, documentary evidence as per GIT for claiming exemption from payment of EMD/Bid security to be uploaded. THE EMD/BID SECURITY DEPOSITED AGAINST OTHER TENDERS CANNOT BE ADJUSTED OR CONSIDERED FOR THIS TENDER. NO INTEREST IS PAYABLE ON EMD/BID SECURITY. EMD/Bid Security of the approved firms, who fulfills pre-qualification requirements, would be retained till the firm is registered at AIIMS for the supply of Drugs/Medicines items.

FIRM WHICH HAD BEEN DECLARED ELIGIBLE ON THE BASIS OF PATENT/NICHE MOLECULE SHALL NOT BE EXEMPT UNDER THIS CLAUSE AND SHALL HAVE TO SUBMIT ALL DOCUMENTS AS PER THE REQUIREMENT OF THIS TENDER

- ii)** Scanned copy of "List of Items Quoted" as per **SECTION – XVI** of Tender Enquiry Document.
- iii)** Scanned copy of "Tender Acceptance Form" as per **Section IX** to be uploaded.
- iv)** Scanned Copy of GST Registration Certificate.
- v)** The Scanned Copies of following documents, wherever applicable may be uploaded under "Other Important Documents":

- a)** Scanned copy of Documentary evidence, as necessary in terms of clauses of GIB establishing that the bidder is eligible to submit the bid and, also, qualified to perform the Rate Contract if its bid is accepted to be uploaded.
- b)** Scanned copy of Power of Attorney in favor of signatory of Tender/Bid to be uploaded.
- c)** Scanned copy of Documents and relevant details to establish in accordance with GIB that the goods to be supplied by the bidder conform to the requirement of the Tender Enquiry Document to be uploaded.
- d)** Scanned copy of Documents confirming to Sole Proprietorship/ Partnership/Private Limited Firm in the country of origin as the case may be to be uploaded.
- vi)** Scanned Copy of undertakings and Other Documents as per TED.

Note:

1. It is the responsibility of bidder to go through the Tender Enquiry Document to ensure uploading all required documents in addition to above, if any

B) Price Bid:

Price Schedule(s) as per BoQ format filled up with all the details including Make, Model etc. of the goods offered to be uploaded.

Schedule of price bid in the form of BoQ_XXXX.xls:

The below mentioned (Section X) price bid format is provided as BoQ_XXXX.xls along with this Tender Enquiry Document at <https://eprocure.gov.in/eprocure/app>. Bidders are advised to download this BoQ_XXXX.xls as it is and quote their offer/rates in the permitted column and upload the same in the commercial bid. Bidder shall not tamper/modify downloaded price bid template in any manner. In case if the same is found to be tempered/modified in any manner, tender will be completely rejected and tenderer is liable to be banned from doing business with AIIMS New Delhi.

- 9.2 The authorized signatory of the bidder must digitally sign the bid. Individuals digitally signing the bid or other documents connected with a Rate Contract must specify whether he signs as:
 - i) A 'Sole Proprietor' of the firm or constituted attorney of such Sole Proprietor.
 - ii) In case of partnership firm he must have authority to quote & to refer to arbitration dispute concerning the business of the partnership either by virtue of the partnership agreement or a power of attorney;
 - iii) Constituted attorney of the firm if it is a company.

Note:

- 1) In case of (ii) above, a copy of the partnership agreement duly registered with "Registrar of Firm's" or general power of attorney, in either, case, attested by a Notary Public should be uploaded, or affidavit on stamped paper of all the partners admitting execution of the partnership agreement or the general power of attorney should be uploaded.
- 2) In case of the partnership firms, where no authority to refer disputes concerning the business of the partnership has been conferred on any partner, the bid and all other related documents must be signed by every partner of the firm and uploaded.
- 3) Person digitally signing the Tender Acceptance Form or any documents forming part of the contract on behalf of another shall be deemed to warrantee that he has authority to bind such other persons and if, on enquiry, it appears that the persons so signing had no authority to do so, the purchaser may, without prejudice to other civil and criminal remedies, liable

for rejection of bid or cancel of contract and hold the signatory liable for all cost and damages.

9.3 A bid, which does not fulfill any of the above requirements and/or gives evasive information/reply against any such requirement, shall be liable to be ignored and rejected.

9.4 Bid sent by fax/email shall be ignored.

10. Bid Currencies

10.1 The bidder supplying indigenous goods or already imported goods shall quote only in Indian Rupees (INR).

10.2 Bids, where prices are quoted in any other way shall be treated as non -responsive and rejected.

11. Bid Prices

11.1 The Bidder shall indicate in the Price Schedule provided in BoQ all the specified components of prices shown therein including the unit prices on Free Delivery At Site basis, applicable GST, HSN Code, it proposes to supply against the requirement. The Bidders shall indicate MRP in the relevant column against each item of BoQ. The details about make & model, if applicable, may also be indicated. All the columns shown in the Price Schedule should be filled up as required.

11.2 In no case the quoted rates should be more than MRP at the time of submission of quotation. If subsequently during the currency of Rate Contract there is decrease in MRP, the bidder shall inform the purchaser promptly along with revised reduced rates on pro-rata basis. In case, if bidder quotes more than MRP and/or does not inform purchaser about reduction in MRP, it will be viewed seriously and appropriate administrative action will be taken including de-barring the firm.

11.3 If there is more than one schedule in the "Schedule of Requirements", the bidder has the option to submit its bid for any one or more schedules. However, while quoting for a schedule, the bidder shall quote for the complete requirement of goods as specified in that particular schedule.

11.4 The need for indication of all such price components by the bidders, as required in this clause is for the purpose of comparison of the bids by the purchaser and will in no way restrict the purchaser's right to award the Rate Contract on the selected bidder on any of the terms offered.

11.5 In case of controlled drugs by the Government (Under DPCO), the quotation must be sent subject to the controlled rates and other conditions and supplier will be paid at the controlled price or rates offered by the supplier whichever is less. Controlled drugs must be clearly mentioned as such in the bidders' quotations.

12. Firm Price

12.1 Prices quoted by the bidder shall remain firm and fixed during the currency of the Rate Contract and not subject to variation on any account. Purchase Orders will be placed by Centers/Hospital/Departments/Store Sections against this Rate Contract till the currency period of Rate Contract.

12.2 Statuary variation in GST will be applicable.

13. Alternative Models/Brands/Quality

13.1 Alternative Models/Brands/Quality are not permitted. The Bidders are required to quote Models/Brands/Quality of best quality meeting tender specifications. Wherever, a bidder quotes alternative Models/ Brands/ Quality, there bid will not be considered for that item.

14. Documents Establishing Bidder's Eligibility and Qualifications

14.1 The bidder shall furnish, as part of its bid, relevant details and documents establishing its eligibility to quote and its qualifications to perform the Rate

Contract if its bid is accepted. The “Qualification Criteria” have been given in Section VIII.

14.2 Quotations shall be strictly according to the required specifications, and in the case of formulations, detailed formula along with the connected Ltrature, Drug licenses etc. should be furnished. The name of the manufacturer and the brand name should also be stated.

15. Documents establishing good's Conformity to Tender Enquiry Document.

15.1 The bidder shall upload in its bid the required as well as the relevant documents like technical data, Ltrature, drawings etc. to establish that the goods offered in the bid fully conform to the goods specified by the purchaser in the Tender Enquiry Document. For this purpose, the bidder shall also upload a clause-by-clause commentary on the technical specifications and other technical details incorporated by the purchaser in the Tender Enquiry Document to establish technical responsiveness of the goods offered in its bid.

15.2 In case there is any variation and/or deviation between the goods prescribed by the purchaser and that offered by the bidder, the bidder shall list out the same in a chart form without ambiguity and provide the same along with its bid.

15.3 If a bidder furnishes wrong and/or misguiding data, statement(s) etc. about technical acceptability of the goods offered by it, its bid will be liable to be ignored and rejected in addition to other remedies available to the purchaser in this regard.

16. Bid Security (BS) /EMD

16.1 Pursuant to the bidder shall furnish along with its bid, Bid Security for amount as shown in the Notice Inviting Tenders (NIT).

16.2 The original Earnest Money/Bid Security must be delivered to address as given in NIT till bid opening date and time as mentioned in “Critical Date Sheet” failing which the bid shall be summarily rejected. The scanned copy of original Bid Security/EMD may be uploaded along with the bid.

16.3 The bidders who are currently registered with MSME for the goods as per Tender document specification shall be eligible for exemption from Bid Security as defined in MSE Procurement Policy issued by the department of MSME. In case the bidder falls in this category, the bidder shall upload relevant certificate of registration for the subject goods issued by department of MSME.

16.4 The Bid Security shall be denominated in Indian Rupees. The Bid Security shall be furnished in one of the following forms:

- Account Payee Demand Draft/ Banker's cheque
- Fixed Deposit Receipt
- Bank Guarantee

16.5 The demand draft or banker's cheque shall be drawn on any commercial bank in India, in favour of as indicated in the NIT payable at New Delhi. In case of Bank Guarantee, the same is to be provided from any commercial bank in India or country of the bidder as per the format specified under Section XI in these documents.

- 16.6 The Bid Security shall be valid for a period of forty-five (45) days beyond the validity period of the bid. As validity period of Bid is 270 days, the Bid Security shall be valid for 315 days from Techno – Commercial Bid opening date.
- 16.7 The Bid Security of successful bidder will be returned without any interest, after receipt of performance security from that bidder.
- 16.8 Bid Security is required to protect the purchaser's right against the risk of the Bidder's conduct, which would warrant the forfeiture of the Bid Security. Bid Security of a bidder will be forfeited, if the bidder withdraws or amends its bids or impairs or derogates from the bid in any respect within the period of validity of its bid or if it comes to the notice that the information/documents furnished in its bid is incorrect, false, misleading or forged without prejudice to other rights of the purchaser. The Bid Security of the successful bidder will be forfeited without prejudice to other rights of Purchaser if it fails to furnish the required performance security within the specified period.

17. Bid Validity

- 17.1 The bid shall remain valid for acceptance for a period of 270 days (Two hundred and Seventy days) after the date of bid opening prescribed in the Tender Document. Any bid valid for a shorter period shall be treated as unresponsive and rejected.
- 17.2 In exceptional cases, the bidder may be requested by the purchaser to extend the validity of their bids up to a specified period. Such request(s) and responses thereto shall be conveyed by mail/fax/email. The bidders, who agree to extend the bid validity, are to extend the same without any change or modification of their original bid and they are also to extend the validity period of the Bid Security accordingly. A bidder, who may not agree to extend its bid validity after the expiry of the original validity period, their bid will not be considered further and the Bid Security furnished by them shall be returned.
- 17.3 In case the day up to which the bids are to remain valid falls on/ subsequently declared a holiday or closed day for the purchaser, the bid validity shall automatically be extended up to the next working day.

18. Instructions for Online Bid Submission and Registration on CPP Portal:

- 18.1 The bidders shall submit their online bids as per the instruction given for online bid process. The bidders are required to submit soft copies of their bids electronically on the CPP Portal, using valid Digital Signature Certificates. The instructions given below are meant to assist the bidders in registering on the CPP Portal, prepare their bids in accordance with the requirements and submitting their bids online on the CPP Portal. More information useful for submitting online bids on the CPP Portal may be obtained at: <https://eprocure.gov.in/eprocure/app>.
- 18.2. Registration on CPP Portal:
 - i) Bidders are required to enrol on the e-Procurement module of the Central Public Procurement Portal (URL: <https://eprocure.gov.in/eprocure/app>) by clicking on the link "Online bidder Enrolment" on the CPP Portal which is free of charge.
 - ii) As part of the enrolment process, the bidders will be required to choose a unique username and assign a password for their accounts.
 - iii) Bidders are advised to register their valid email address and mobile numbers as part of the registration process. These would be used for any communication from the CPP Portal.

- iv) Upon enrolment, the bidders will be required to register their valid Digital Signature Certificate (Class II or Class III Certificates with signing key usage) issued by any Certifying Authority recognized by CCA India (e.g. Sify/nCode /eMudhra etc.), with their profile.
- v) Only one valid DSC should be registered by a bidder. Please note that the bidders are responsible to ensure that they do not lend their DSC's to others which may lead to misuse.
- vi) Bidder then logs in to the site through the secured log-in by entering their user ID / password and the password of the DSC / e-Token.

18.3. Searching for Tender Enquiry Document on CPP Portal:

- i) There are various search options built in the CPP Portal, to facilitate bidders to search active tenders by several parameters. These parameters could include Tender ID, Organization Name, Location, Date, Value, etc. There is also an option of advanced search for tenders, wherein the bidders may combine a number of search parameters such as Organization Name, Form of Contract, Location, Date, Other keywords etc. to search for a tender published on the CPP Portal.
- ii) Once the bidders have selected the tenders they are interested in, they may download the required documents / tender schedules. These tenders can be moved to the respective 'My Tenders' folder. This would enable the CPP Portal to intimate the bidders through SMS / e-mail in case there is any corrigendum issued to the tender document.
- iii) The bidder should make a note of the unique Tender ID assigned to each tender, in case they want to obtain any clarification / help from the Helpdesk.

18.4. Preparation of Bids for uploading on CPP Portal

- i) Bidder should take into account any corrigendum published on the tender document before submitting their bids.
- ii) Please go through the tender advertisement and the Tender Enquiry Document carefully to understand the documents required to be submitted as part of the bid. Please note the number of covers in which the bid documents have to be submitted, the number of documents - including the names and content of each of the document that need to be submitted. Any deviations from these may lead to rejection of the bid.
- iii) Bidder, in advance, should get ready the documents/BoQ to be uploaded as indicated in the Tender Enquiry Document and generally, they can be in PDF / XLS / RAR / DWF/JPG formats. Scanned documents to be uploaded may be scanned with 100 dpi with black and white option which helps in reducing size of the scanned document and resulting in fast uploading. It is the responsibility of the bidder to ensure that uploaded scanned documents are legible.
- iv) To avoid the time and effort required in uploading the same set of standard documents which are required to be submitted as a part of every bid, a provision of uploading such standard documents has been provided to the bidders. Bidders can use "My Space" or "Other Important Documents" area available to them to upload such documents. These documents may be directly submitted from the "My Space" area while submitting a bid, and need not be uploaded again and again. This will lead to a reduction in the time required for bid submission process.

19. Submission of Bids for uploading on CPP Portal

19.1 Bidder should log into the site well in advance for bid submission so that they can upload the bid in time i.e. on or before the bid submission time. Bidder will be responsible for any delay due to other issues.

- 19.2 The bidder has to digitally sign and upload the required bid documents one by one as indicated in the Tender Enquiry document.
- 19.3 Bidder has to select the payment option as “offline” to pay the Bid Security/ EMD as applicable and enter details of the instrument.
- 19.4 Bidder should prepare the Bid Security/EMD as per the instructions specified in the Tender Enquiry Document. The original should be posted/couriered/given in person to the concerned official, latest by the last date of bid submission or as specified in the Tender Enquiry Document. The details of the DD/any other accepted instrument, physically sent, should tally with the details available in the scanned copy and the data entered during bid submission time. Otherwise the uploaded bid will be rejected.
- 19.5 Bidders are requested to note that they should necessarily submit their financial bids in the format provided and no other format is acceptable. If the price bid has been given as a standard BoQ format with the tender document, then the same is to be downloaded and to be filled by all the bidders. Bidders are required to download the BoQ file, open it and complete the white coloured (unprotected) cells with their respective financial quotes and other details (such as name of the bidder). No other cells should be changed. Once the details have been completed, the bidder should save it and submit it online, without changing the filename. If the BoQ file is found to be modified by the bidder, the bid will be rejected.
- 19.6 The server time (which is displayed on the bidders’ dashboard) will be considered as the standard time for referencing the deadlines for submission of the bids by the bidders, opening of bids etc. The bidders should follow this time during bid submission.
- 19.7 All the documents being submitted by the bidders would be encrypted using PKI encryption techniques to ensure the secrecy of the data. The data entered cannot be viewed by unauthorized persons until the time of bid opening. The confidentiality of the bids is maintained using the secured Socket Layer 128 bit encryption technology. Data storage encryption of sensitive fields is done. Any bid document that is uploaded to the server is subjected to symmetric encryption using a system generated symmetric key. Further this key is subjected to asymmetric encryption using buyers/bid openers’ public keys. Overall, the uploaded tender documents become readable only after the tender opening by the authorized bid openers.
- 19.8 The uploaded Tender/Bid shall become readable only after the tender opening by the authorized bid openers.
- 19.9 Upon the successful and timely submission of bids (i.e. after Clicking “Freeze Bid Submission” in the portal), the portal will give a successful bid submission message & a bid summary will be displayed with the bid no. and the date & time of submission of the bid with all other relevant details.
- 19.10 The bid summary has to be printed and kept as an acknowledgement of the submission of the bid. This acknowledgement may be used as an entry pass for any bid opening meetings.
- 19.11 Assistance to Bidders for uploading CPP Portal:
 - i) Any queries relating to the Tender Enquiry Document and the terms and conditions contained therein should be addressed to the Tender Inviting Authority for a tender or the relevant contact person indicated in the NIT.

- ii) Any queries relating to the process of online bid submission or queries relating to CPP Portal in general may be directed to the 24x7 CPP Portal Helpdesk

E. BID OPENING

20. Opening of Bids

- 20.1 E- Bids will be opened after due time and date and the bidders may check the status etc. on CPP Portal.
- 20.2 No change/alteration on plea of clerical or typographical error in rates or other terms in the tender will be permitted under any circumstances.
- 20.3 Withdrawal of the complete tender can be allowed but in such cases, the earnest money shall be forfeited in full.
- 20.4 Partial withdrawal (in respect of one or more items quoted) will not be allowed under any circumstances.

F. SCRUTINY AND EVALUATION OF BIDS

21. Basic Principle

- 21.1 Bids will be evaluated on the basis of the terms & conditions already incorporated in the Tender Enquiry Document, based on which bids have been received and the terms, conditions etc. mentioned by the bidders in their bids. No new condition will be brought in while scrutinizing and evaluating the bids.

22. Scrutiny of Bids

- 22.1 The Purchaser will examine the Bids to determine whether they are complete, whether any computational errors have been made, whether required Bid Securities have been furnished, whether the documents have been properly signed stamped and whether the Bids are generally in order.

- 22.2 The Purchaser's determination of a Bid's responsiveness is to be based on the contents of the Bid itself without recourse to extrinsic evidence.

- 22.3 The Bids will be scrutinized to determine whether they are complete and meet the essential and important requirements, conditions etc. as prescribed in the Tender Enquiry Document.

22.4 PHARMACOPOEIAL SPECIFICATION:

Pharmacopoeia' specifications i.e. IP/BP/USP should be clearly mentioned against each drug/constituent of the drug quoted as per the provisions of Drug and Cosmetics Act, 1945.

- 22.5 In the absence of submission of the following, a bid shall be declared non-responsive during the evaluation and will be ignored;

- i) Tender Acceptance Form as per Section IX (signed & stamped) not uploaded.
- ii) Bid validity is shorter than the required period.
- iii) Required Bid Security (Amount, validity etc.)/exemption documents have not been uploaded as per stipulated provisions.
- iv) Bidder has not agreed to give the required Performance Security of required amount in an acceptable form for due performance of the contract.
- v) Bidder has not agreed to other essential condition(s) specially incorporated in the Tender document like terms of payment, liquidated damages clause, shelf life clause, warranty clause, dispute resolution mechanism, and applicable law.
- vi) Poor/unsatisfactory past performance.

- vii) Bidders who stand de-registered/banned/blacklisted by any Central Govt. /State Govt. Ministries/AIIMS, New Delhi.
- viii) Bidder has not agreed to currency of Rate Contract period.
- ix) Bidder has not agreed for the delivery terms and delivery period.

22.6 INSPECTION OF FIRM'S PREMISES:

The Director or his nominee reserves the right for inspection of the pharmaceutical firms participating in the tenders, by officers appointed by the Director. They can carry out inspection for assessing the capacity/capability/eligibility of the firm to make supplies on the basis of rate-contract and to ensure that good manufacturing practices are being followed by the manufacturer. The decision of the Director shall be final in this regard.

23. Minor Infirmitiy/Irregularity/Non-Conformity

23.1 If during the evaluation, the purchaser finds any minor informality and/or irregularity and/or non-conformity in a bid, the purchaser will convey its observation on such 'minor' issues, which has not price implication, to the bidders by registered/speed post/ e-mail/fax etc. asking the bidder to respond by a specified date. If the bidder does not reply by the specified date or gives evasive reply without clarifying the point at issue in clear terms, that bid will be liable to be ignored.

24. Qualification Criteria

24.1 Bids of the bidder, who have not uploaded required documents or do not meet the required Qualification Criteria prescribed in Section VIII, will be treated as non - responsive and will not be considered further.

25. Item-wise Evaluation

25.1 In case the Schedule of Requirements contains multiple items, the responsive bids will be evaluated and compared separately for each item.

26. Comparison of Bids

26.1. The comparison of the responsive Bids shall be carried out on Free Delivery at consignee site basis.

27. Purchase Preference for Evaluation

27.1 The Purchaser reserves the right to give the price preference to small-scale sectors etc. and purchase preference to central public sector undertakings as per the instruction in vogue while evaluating, comparing and ranking the responsive Bids.

28. Bidder's capability to perform the Rate Contract

28.1 The purchaser, through the above process of bid scrutiny and bid evaluation will determine to its satisfaction whether the bidder, whose bid has been determined as the lowest evaluated responsive bid is eligible, qualified and capable in all respects to perform the Rate Contract satisfactorily.

28.2 The above-mentioned determination will, inter alia, take into account the bidder satisfying all the requirements of the purchaser as incorporated in the Tender Enquiry Document. Such determination will be based upon scrutiny and examination of all relevant data and details submitted by the bidder in its bid as well as such other allied information as deemed appropriate by the purchaser.

29. Contacting the Purchaser

29.1 From the time of submission of bid to the time of awarding the Rate Contract, if a bidder needs to contact the purchaser for any reason relating to NIT/Tender Enquiry Document and / or its bid, it should do so only through CPP portal.

29.2 In case a bidder attempts to influence the purchaser in the purchaser's decision on scrutiny, comparison & evaluation of bids and awarding the contract, the bid

of the bidder shall be liable for rejection in addition to appropriate administrative actions being taken against that bidder, as deemed fit by the purchaser.

G. AWARD OF RATE CONTRACT

30. Purchaser's Right to accept any bid and to reject any or all bids.

30.1 The purchaser reserves the right to accept in part or in full any bid or reject any or more bid(s) without assigning any reason or to cancel the Tender process and reject all bids at any time prior to award of Rate Contract , without incurring any liability, whatsoever to the affected bidder(s).

31. Award Criteria

31.1 Subject to the above, the Rate Contract will be awarded to the lowest evaluated responsive bidder decided by the purchaser. In cases where advance samples have been called in "Special Instructions to Bidders" in Section III,

32. Purchase Orders to be placed during currency of Rate Contract

32.1 Purchase Orders will be placed from time to time by the Centers/Hospitals/Department/ Store Sections of AIIMS during the currency of Rate Contract, as per actual requirement, in which the exact quantities required on each occasion together with the date of delivery shall be specified in the purchase order.

33. Notification of Award

33.1 Before expiry of the bid validity period, the purchaser will notify the successful bidder (s) in writing, by registered / speed post or by fax/ email (to be confirmed by registered / speed post) that its bid for Goods, which have been selected by the purchaser, has been accepted, also briefly indicating there in the essential details like description, specification and quantity of the goods and corresponding prices accepted. The successful bidder must furnish to the purchaser the required Performance Security within thirty days from the date of dispatch of this notification, failing which the Bid Security will be forfeited and the award will be cancelled. Relevant details about the Performance Security have been provided in clause 3 of GCC under Section IV.

33.2 The Notification of Award shall constitute the conclusion of the Rate Contract.

34. Issue of Rate Contract

34.1 Promptly after notification of award, the Purchaser will mail the Rate Contract form (as per Section XIII) duly completed and signed, in duplicate, to the successful bidder by registered / speed post.

34.2 Within twenty one days from the date of the Rate Contract, the successful bidder shall return the original copy of the Rate Contract, duly signed and dated, to the Purchaser/ by registered / speed post/courier.

35. Non-receipt of Performance Security by the Purchaser

35.1 Failure of the successful bidder in providing Performance Security and / or returning Rate Contract copy duly signed in terms of GIB clauses above shall make the bidder liable for forfeiture of its Bid Security and, also, for further actions by the Purchaser it as per the clause 12-Termination of default of GCC under Section IV.

36. Return of Bid Security/EMD

36.1 The Bid Security/EMD of the successful bidder and the unsuccessful bidder will be returned to them without any interest, whatsoever, in terms of Clause 19 of GIB.

37. Publication of Bid Result

37.1 The name and address of the successful bidder(s) receiving the Rate Contract(s) will be mentioned in the CPP Portal.

H. CORRUPT OR FRAUDULENT PRACTICES

38. Corrupt or Fraudulent Practices

38.1 It is required by all concerned namely the Bidder /Suppliers/ Purchaser/Consignee/End User etc. to observe the highest standard of ethics during the procurement and execution of such Rate Contract/Purchase Orders. In pursuance of this policy, the Purchaser: -

- a) defines, for the purposes of this provision, the terms set forth below as follows:
 - i) "corrupt practice" means the offering, giving, receiving or soliciting of anything of value to influence the action of a public official in the procurement process or in Rate Contract/Purchase Orders execution; and
 - ii) "fraudulent practice" means a misrepresentation of facts in order to influence a procurement process or the execution of a Rate Contract/Purchase Orders to the detriment of the Purchaser, and includes collusive practice among bidders (prior to or after Bid submission) designed to establish Bid prices at artificial non-competitive levels and to deprive the Purchaser of the benefits of free and open competition;
- b) Will reject a proposal for award if it determines that the Bidder recommended for award has engaged in corrupt or fraudulent practices in competing for the Rate Contract/Purchase Orders in question;
- c) Will declare a firm ineligible, either indefinitely or for a stated period of time, to be awarded a Rate Contract/Purchase Orders by the purchaser if it at any time determines that the firm has engaged in corrupt or fraudulent practices in competing for, or in executing the Rate Contract/Purchase Orders.

SECTION – III
SPECIAL INSTRUCTIONS TO BIDDERS (SIB)

The following Special Instructions to Bidders will apply for this purchase. These special instructions will modify/substitute/supplement the corresponding General Instructions to Bidders (GIB) incorporated in Section II. The corresponding GIB clause numbers have also been indicated in the text below:

In case of any conflict between the provision in the GIB and that in the SIB, the provision contained in the SIB shall prevail.

Sl. No.	GIB Clause No.	Topic	SIB Provision
1.	1 - 38		No Change

1. If required, the bidder will submit the samples for each item in original packing, duly labeled (Printed) and sealed having date of manufacturing, date of Expiry, manufactured by with batch No. Stores Officer (H) within 10 days. If the bidder fails to submit the sample within given time, the bid will be summarily rejected and no correspondence will be entertained in this regard.

SECTION - IV
GENERAL CONDITIONS OF CONTRACT (GCC)

1. Application

1.1 The General Conditions of Contract incorporated in this section shall be applicable for this purchase to the extent the same are not superseded by the Special Conditions of Contract prescribed under Section V, Schedule of Requirements under Section VI and Technical Specification under Section VII of this document.

2. Patent Rights

2.1 The supplier shall, at all times, indemnify and keep indemnified the purchaser, free of cost, against all claims which may arise in respect of goods to be provided by the supplier under the Rate Contract/Purchase Orders for infringement of any intellectual property rights or any other right protected by patent, registration of designs or trademarks. In the event of any such claim in respect of alleged breach of patent, registered designs, trademarks etc. being made against the purchaser, the purchaser shall notify the supplier of the same and the supplier shall, at his own expenses take care of the same for settlement without any liability to the purchaser.

3. Performance Security

3.1 Within Thirty (30) days from date of the issue of Notification of Award by the Purchaser, the supplier shall furnish Performance Security to the Purchaser for an amount equal to three percent (3%) of the Total Estimated Quantity of the items for which Rate Contract is being awarded.

3.2 The Performance Security shall be denominated in Indian Rupees in any of the following forms:

- i) Account Payee Demand Draft
- ii) Fixed Deposit Receipt drawn from any Scheduled bank in India
- iii) Bank Guarantee issued by a Scheduled bank in India, in the prescribed form as provided in Section XII of this document

3.3 In the event of any failure /default of the supplier with or without any quantifiable loss to the government, the amount of the Performance Security is liable to be forfeited equivalent to the amount of Supply Order. The needful will be done to cover any failure/default of the supplier with or without any quantifiable loss to the Government.

3.4 In the event of any extension of currency of Rate Contract, the supplier shall, within fifteen (15) days of issue of the amendment, furnish the corresponding amendment to the Performance Security (as necessary), rendering the same valid in all respects in terms of the Rate Contract , as amended.

3.5 Subject to above, the Purchaser will release the Performance Security without any interest to the supplier on completion of the supplier's all contractual obligations including the warranty obligations (if applicable).

4. Technical Specifications

4.1 The Goods to be provided by the supplier under this Rate Contract shall conform to the 'Technical Specification' under Sections VII of this document.

5. Inspection, Testing and Quality Control

5.1 The purchaser has contractual right to inspect, test and, if necessary, reject the goods to confirm their conformity to the Rate Contract specifications and other quality control details incorporated in the Rate Contract.

5.2 If during such inspections and tests the contracted goods fail to conform to the required specifications and standards, the purchaser may reject them and the supplier shall either replace the rejected goods or make all alterations necessary to meet the specifications and standards, as required, free of cost to the purchaser and re-submit the same to the purchaser for conducting the inspections and tests again. No payment shall be made for rejected material and In case rejected goods are not removed, these will be disposed off in a manner as deemed fit by the authorities at the risk and responsibility of the suppliers without any further notice.

5.3 Regular and random testing of drugs will be undertaken by AIIMS from any NABL accredited /Govt. approved laboratories (Annexure attached) at the time of supply and at any time during the shelf life or whenever any defect is noticed. The Director AIIMS shall be at liberty to undertake regular and random testing of the drugs supplied by the pharmaceutical firm/ bidder at regular interval to maintain and ensure the quality of drugs.

5.4 The report of the NABL accredited/Govt. approved laboratory shall be accepted by the pharmaceutical firm. In case the same is disputed by the pharmaceutical firm, the report of the approved Central Drug Testing Laboratory as approved by CDSCO (Appellate Authority) only will be accepted as final. However, the same should be submitted within three months, from the date of communication of the disputed test report to the pharmaceutical firm. For this, the pharmaceutical firm should approach the concerned Drug Control Authorities for getting the drugs tested, as per procedure.

5.5 If any drug sample fails the test or is found to be of substandard quality, action as below will be initiated:

- (a) If any store/stores supplied against the contract are found to be not of standard quality as per specifications on analysis and/or on inspection by competent authority, the Institute will destroy the entire consignment against the particular invoice, irrespective of fact that part of the supplied stores may have been consumed. The institute shall not be liable to make any payments in lieu of inferior items.
- (b) If the firm fails to make fresh supplies in lieu of substandard quality of drug, it is liable to be debarred for three years in respect of all the items in the rate-contract of this Institute and EMD/Performance security shall be forfeited.
- (c) If the product is found to be not of standard quality, the cost of testing done by the Institute will be recovered from the supplier.
- (d) In case, the supplies are found to be of inferior quality on three occasions, the firm shall be liable for debarment for subsequent tender of Drugs and EMD/Performance security shall be forfeited.
- (e) A copy of the test report will be sent to the DCGI for necessary action at their end.

(f) If any drugs supplied against this Rate Contract are found to be not of standard quality on inspection by Competent Authority, the pharmaceutical firm will be liable to replace the entire quantity within 15 days otherwise risk purchase will be charged from the company and the cost of testing will be recovered from the supplier.

5.6 Goods accepted by the purchaser/consignee in inspection in terms of the Rate Contract/Purchase Orders shall in no way dilute purchaser's/consignee's right to reject the same later, if found deficient in terms of the warranty clause, if applicable.

Quality Control

- I. The stores offered should comply with the provisions of the Drugs and Cosmetics Act, 1940 and the Rules made there under as and Drug Price Control order.
- II. While quoting against items with ISI Mark, it should be ensured that ISI code number is indicated on quotation and at the time of making the supplies, the pharmaceutical firm should ensure that the items supplied has ISI Mark as well as Code Number, as is the statutory requirement of the Bureau of Indian Standards. The attested copy of the valid ISI Marking license issued by Bureau of Indian Standards should be enclosed along with the quotation.

6. Terms of Delivery

6.1 Goods shall be delivered by the supplier on "Free Delivery At Site" basis and delivered as per Delivery Period specified in the Purchase Order placed against Rate Contract. Please note that the time shall be the essence of the contract.

6.2 The goods are to be supplied by F.O.R. destination and all the transit loss/expenses whatsoever, will be borne by the supplier/firm.

7. Warranty

7.1 The supplier warrants comprehensively that the goods supplied under the Rate Contract is new, unused and incorporate all recent improvements in design and materials unless prescribed otherwise by the purchaser in the Rate Contract. The supplier further warrants that the goods supplied under the Rate Contract/Purchase Orders shall have no defect arising from design, materials or workmanship or from any act or omission of the supplier that may develop under normal use of the supplied goods under the conditions prevailing in India.

7.2 The warranty period (if applicable as stated in Schedule of Requirement in Section-VI or Technical Specification in Section- VII) shall include all spares, labor and preventive maintenance from the date of completion of the satisfactory installation and acceptance till warranty period.

8. Prices

8.1 Prices quoted by the bidder shall remain firm and fixed during the currency of the Rate Contract and not subject to variation on any account. Purchase Orders will be placed by Centers/Hospital/Departments/Store Sections against this Rate Contract till the currency period of Rate Contract.

8.2 Statuary variation in GST will be applicable during currency of the contract, during the original Delivery Period of Purchase Order after

submitting supporting documents (Government notifications) issued by concern department.

8.3 **Rate Revision:** Successful bidders shall not be entitled to any rate-revision of price for any reason except Govt. levies which become applicable after finalization of rate contract along with adequate documentary proof thereof.

9. Payment Terms

9.1 100% payment would be made on receipt of goods in good condition and acceptance, upon the submission of the following documents:

- i) Original copies of supplier's invoice showing Rate Contract/Purchase Orders number, goods description, quantity, packing list, unit price and total amount;
- ii) "Consignee Receipt Certificate" as per Section XIV of Tender document in original
- iii) "Final Consignee Acceptance Certificate" as per Section XV of goods to be issued by the End User subject to recoveries, if any, either on account of non-rectification of defects/deficiencies not attended by the Supplier or otherwise.

9.2 Any dues or payments that have arisen to the Institution from the supplier for which no specific time-limit has been laid down in the terms & conditions, shall be payable by the supplier within such time limit as may be prescribed in the various letters/orders addressed to the contractors. On failure to do so the supplier shall be liable to be debarred for not paying dues or payment etc. to the hospital for a period as decided by the Director or his nominee.

9.3 Conditions of advance payments or payment against delivery shall not be accepted.

10. Delivery

10.1 The supplier shall deliver the goods under the Rate Contract within the time schedule specified by the Purchaser Order as per in the Schedule of Requirements and as incorporated in the Rate Contract. The time for and the date of delivery of the goods stipulated in the Purchase Order shall be deemed to be of the essence of the contract and the delivery must be completed no later than the date (s) as specified in the Purchase Order.

10.2 Supply orders placed against the contract, on or just before last date of the tenure of contract will have to be accepted /honored by the supplier.

10.3 **No guarantee can be given as to the minimum quantity which will be demanded against this contract, but the supplier will supply such quantity as may be ordered by the Stores Officer during the tenure of the contract.**

10.4 Subject to the provision under Force Majeure clause of GCC, any unexcused delay by the supplier in maintaining its contractual obligations towards delivery of goods shall render the supplier liable to any or all of the following sanctions:

- i) Imposition of liquidated damages,
- ii) Forfeiture of its Performance Security and
- iii) Termination of the Rate Contract/Purchase Orders for default.

10.5 If at any time during the currency of the Rate Contract, the supplier encounters conditions hindering timely delivery of the goods, the supplier shall promptly inform the Purchaser in writing but not later than 10 days from the date of issue of the Purchase Order about the same and its likely duration and make a request to the Purchaser for extension of the delivery schedule accordingly. In case no communication is received within 10 days from the date of issue of Purchase Order, it will be presumed that supplier has accepted the Purchase Order in all regards. On receiving the supplier's communication, the Purchaser shall examine the situation as soon as possible and, at its discretion, may agree to extend the delivery schedule, with or without liquidated damages for completion of supplier's contractual obligations by issuing an amendment to the Purchase Order.

10.6 When the period of delivery is extended due to unexcused delay by the supplier, the amendment letter extending the delivery period shall, inter alia contain the following conditions:

- i) The Purchaser shall recover from the supplier, under the provisions of the Force Majeure clause of the General Conditions of Contract, Liquidated Damages on the goods, which the Supplier has failed to deliver within the delivery period stipulated in the Purchase Order.
- ii) That no increase in price on account of any ground, whatsoever, including any stipulation in the Rate Contract for increase in price on any other ground and, also including statutory increase in or fresh imposition of GST levied in respect of the goods specified in the Purchase Order, which takes place after the date of delivery stipulated in the Purchase Order shall be admissible on such of the said goods as are delivered and performed after the date of the delivery stipulated in the Purchase Order.
- iii) But nevertheless, the Purchaser shall be entitled to the benefit of any decrease in price on account of reduction in GST which takes place after the expiry of the date of delivery stipulated in the Purchase Order.

10.7 The supplier shall not dispatch the goods after expiry of the delivery period. The supplier is required to apply to the Purchaser for extension of delivery period and obtain the same before dispatch. In case the supplier dispatches the goods without obtaining an extension, it would be doing so at its own risk and no claim for payment for such supply and / or any other expense related to such supply shall lie against the purchaser.

10.8 Passing of Property

- (i) The property in the goods shall not pass to the purchaser unless and until the goods have been delivered to the consignee in accordance with the contract.
- (ii) Where there is a contract for sale of specific goods and the supplier is bound to do something to the goods for the purpose of putting them into a deliverable state the property does not pass until such thing is done.
- (iii) Unless otherwise agreed, the goods remain at the supplier's risk until the property therein is transferred to the purchaser.

10.9 The delivery period should not exceed 45 (forty five) days for all supplies but in emergency the delivery period may be reduced up to 15 days and firm is bound to supply the items within DOD (Date of delivery) period. Bidders are hereby directed to quote the rates of only those drugs/medicines for which they can ensure supply within 45 days of issue of supply-order along with Test Report either on Form 39 from Govt. approved analytical testing laboratory or from in house Test Lab (approved by NABL (National Accreditation Board for Testing and Calibration Laboratories or GLP (Good Lab Practice) accredited Lab. without which the supply will not be accepted. It will be the responsibility of the vendor to provide the certificate of NABL/GLP accredited of the laboratory from which the test report is given. In case the total value of supply order of drugs is less than Rs.-10,000/- in house Lab Test Report will be accepted. However, AIIMS reserves the right to get the supplies tested again from a Govt. /NABL accredited laboratory. In case of failure to either supply the goods within DOD (Date of delivery) period or if goods are not accompanied with lab. test report, they may be debarred, after three defaults, from participating in the next tender for a period of three years and their EMD/ Bid Security/Performance Security Money may be forfeited and risk purchase clause will be invoked. However, in case of imported drugs, In house Test Report of the manufacturing Company will be accepted.

10.10 Supply time: Timing 2.00 P.M to 4.00 P.M (from Monday to Friday) & 11.00 A.M to 12.00 Noon (on Saturday).

10.11 Before making the supply, approved rate contract holder should ensure that all labels of cartons, ampoules, vials, bottles, jars, tubes etc. should be embossed, imprinted, stamped with letters, other requirements like "AIIMS SUPPLY NOT FOR SALE" stamp with permanent ink on each item/strip up to primary level. The supply Challan should be accompanied by test report from NABL accredited lab/Govt. Approved Lab. While delivering the supplies, the firm will ensure that quantities are as per challan, quality of material is as per Rate contract specifications etc. All the items which are stamped with "AIIMS SUPPLY NOT FOR SALE" mark, including rejected stores, cannot be sold to the public by the bidder.

10.12 The supplier shall arrange to effect free replacement of any quantity which may deteriorate in potency, strength approaching expiry or expired etc. before the date of expiry marked on the labels.

10.13 If the supplied item is not utilized before expiry date the supplier should undertake to replace with fresh stock of items as and when required.

10.14 MARKING: Each packing shall be marked with nomenclature of the drug and shall be labeled in accordance with the requirement of the Drugs and Cosmetics Act, 1940 and the rules made there under.

10.15 **PACKING:**

- 1) Tendering firms must quote for the packing specified against each item in the schedule annexed to the rate-enquiry, as any other packing may not be accepted.
- 2) Where no pack is specified, bidders may quote for standard pack which is available in the market.
- 3) Loose supplies / damaged packing / tampered or damaged labeled supplies shall not be accepted under any circumstances.
- 4) Rates should be quoted for strip packing only except where mentioned.
- 5) Supplies to be made in the box of Standard packing. However tablets/capsules in loose pack (tin/bottle) shall not be accepted.

- 6) Liquid orals to be supplied only in glass / plastic bottles conforming to IP/BP/USP/Drugs & Cosmetics Act, 1940.
- 7) Large volume parenteral to be quoted and supplied only in glass/plastic bottles / poly packs conforming to I.P. /BP/USP/ Drug & Cosmetic Act, 1940.
- 8) It should be ensured that only first use packaging material of uniform size including bottles and vials, is used for making supplies on the basis of rate-contract.
- 9) All primary packing containers should be strictly conforming to the specification included in the relevant pharmacopoeia.
- 10)Packing should be able to prevent damage or deterioration during transit.
- 11)All containers i.e. bottles, cartons, tubes etc. are required to be secure with pilferage-proof seals to ensure genuineness of the products packed and the correctness of the contents.MRP should not be written/embossed/should be defaced with indelible ink on any labels otherwise it will disqualified for that supply.

10.16 The supply offered should comply with the provisions of the Drugs and Cosmetics Act, 1940 and the Rules made there under as amended up to date and Drug Price Control order.

11. Liquidated Damages

11.1 PENALTY FOR NON-SUPPLY/LATE SUPPLY

- i) Subject to Force Majeure clause of the General Conditions of Contract, if the supplier fails to deliver any or all of the goods within the time frame(s) incorporated in the Purchase Order, the Purchaser shall, without prejudice to other rights and remedies available to the Purchaser under the Rate Contract, deduct from the Purchase Order, as liquidated damages, a sum equivalent to 0.5% per week of delay or part thereof on delayed supply of goods, until actual delivery or performance subject to a maximum of 10% of the Purchase Order price. Once the maximum is reached Purchaser may consider termination of the Purchase Order as per GCC.
- ii) If supplier fails to execute the supply order three times during the period of rate contract, it shall be debarred for the next three years with effect from the last failure and forfeiting of Performance Security for that drug

11.2 In case of default institute will have the right to procure the ordered item from open market /another party at their own risk and expenses under risk purchase clause.

11.3 The approved rate contract holders should supply all their ordered items within DOD period as per supply order terms and these terms should be strictly adhered to. **In case they fail to supply the item within DOD period, the reminder letter would not be issued in any circumstances and penalty will be imposed.** The item would be arranged either through local purchase or from open market under Risk Purchase Clause without any information in this regard. The difference amount shall be recovered from the pending dues of the firm. **In the eventuality of such instances being repeated, administrative action shall be initiated as per AIIMS procedure which may lead to debarring of the firm for subsequent tenders (up to 3 years).**

11.4 It is hereby also informed that in case any administrative action (imposing of liquidated damages, warning letter, risk purchase, short supply etc.) is taken by the AIIMS during the rate contract period against any approved

vendor, it would be reflected during finalization of the next rate contract as "Past performance" of that firm.

11.5 The Director or his nominee reserves the right to invite at his sole discretion, separate quotations to effect purchase outside this contract in the event of any urgent demand arising in hospital, where no stock is held or otherwise.

12. Termination for Default

12.1 The Purchaser without prejudice to any other contractual rights and remedies available to it the Purchaser, may, by written notice of default sent to the supplier, terminate the Rate Contract and/or Purchase Order in whole or in part, if the supplier fails to deliver any or all of the goods or fails to perform any other contractual obligation(s) within the time period specified in the Purchase Order, or within any extension thereof granted by the Purchaser.

12.2 The Performance Security in such cases will be forfeited equivalent to the amount of Purchase Order.

12.3 Unless otherwise instructed by the Purchaser, the supplier shall continue to perform the Rate Contract/Purchase Orders to the extent not terminated.

13. Termination for Insolvency

13.1 If the supplier becomes bankrupt or otherwise insolvent, the purchaser reserves the right to terminate the Rate Contract/Purchase Orders at any time, by serving written notice to the supplier without any compensation, whatsoever, to the supplier, subject to further condition that such termination will not prejudice or affect the rights and remedies which have accrued and / or will accrue thereafter to the Purchaser.

14. Force Majeure

14.1 Notwithstanding the provisions contained in above clauses of GCC, the supplier shall not be liable for imposition of any such sanction so long the delay and/or failure of the supplier in fulfilling its obligations under the Rate Contract/Purchase Orders is the result of an event of Force Majeure.

14.2 For purposes of this clause, Force Majeure means an event beyond the control of the supplier and not involving the supplier's fault or negligence and which is not foreseeable and not brought about at the instance of the party claiming to be affected by such event and which has caused the non – performance or delay in performance. Such events may include, but are not restricted to, wars or revolutions, hostility, acts of public enemy, civil commotion, sabotage, fires, floods, explosions, epidemics, quarantine restrictions, strikes excluding by its employees, lockouts excluding by its management and freight embargoes.

14.3 If a Force Majeure situation arises, the supplier shall promptly notify the Purchaser in writing of such conditions and the cause thereof within twenty one days of occurrence of such event. Unless otherwise directed by the Purchaser in writing, the supplier shall continue to perform its obligations under the Rate Contract/Purchase Orders as far as reasonably practical, and shall seek all reasonable alternative means for performance not prevented by the Force Majeure event.

- 14.4 If the performance in whole or in part or any obligation under this Rate Contract/Purchase Orders is prevented or delayed by any reason of Force Majeure for a period exceeding sixty days, either party may at its option terminate the Rate Contract/Purchase Orders without any financial repercussion on either side.
- 14.5 In case due to a Force Majeure event the Purchaser is unable to fulfill its contractual commitment and responsibility, the Purchaser will notify the supplier accordingly and subsequent actions taken on similar lines described in above sub-paragraphs.

15. Termination for Convenience

- 15.1 The Purchaser reserves the right to terminate the Rate Contract, in whole or in part for its Purchaser's convenience, by serving written notice on the supplier of 30 days at any time during the currency of the Rate Contract.
- 15.2 The Supplier reserves the right to terminate the Rate Contract, in whole or in part for its Purchaser's convenience, by serving written notice by the supplier of 90 days at any time during the currency of the Rate Contract.

16. Resolution of Disputes

- 16.1 If dispute or difference of any kind shall arise between the Purchaser/Consignee and the supplier in connection with or relating to the Rate Contract/Purchase Orders, the parties shall make every effort to resolve the same amicably by mutual consultations.
- 16.2 If the parties fail to resolve their dispute or difference by such mutual consultation within twenty-one days of its occurrence, then, unless otherwise provided in the SCC, either the Purchaser/Consignee or the supplier may give notice to the other party of its intention to commence arbitration, as hereinafter provided the applicable arbitration procedure will be as per the Arbitration and Conciliation Act, 1996 of India.
- 16.3 In the case of a dispute or difference arising between the Purchaser and a domestic Supplier relating to any matter arising out of or connected with the Rate Contract/Purchase Orders, such dispute or difference shall be referred to the sole arbitration to be appointed by the Director, AIIMS. The award of the arbitrator shall be final and binding on the parties to the Rate Contract/Purchase Orders subject to the provision that the Arbitrator shall give reasoned award in case the value of claim in reference exceeds Rupees One lakhs (Rs. 1,00,000/-)
- 16.4 **Venue of Arbitration:** The venue of arbitration shall be the place from where the Rate Contract/Purchase Orders has been issued, i.e., New Delhi, India.
- 16.5 **Jurisdiction of the court** will be from the place where the Tender Document has been issued, i.e., New Delhi, India.
- 16.6 **Applicable Law:** The Rate Contract/Purchase Orders shall be governed by and interpreted in accordance with the laws of India for the time being in force.

17 Withholding and Lien in respect of sums claimed

- 17.1 Whenever any claim for payment arises under the Rate Contract/Purchase Orders against the supplier the purchaser shall be entitled to withhold and also have a lien to retain such sum from the security deposit or sum of money arising out of under any other Rate Contract/Purchase Orders made by the supplier with the purchaser, pending finalization or adjudication of any such claim.
- 17.2 It is an agreed term of the Rate Contract/Purchase Orders that the sum of money so withheld or retained under the lien referred to above, by the purchaser, will be kept withheld or retained till the claim arising about of or under the Rate Contract/Purchase Orders is determined by the Arbitrator or by the competent court as the case may be and the supplier will have no claim for interest or damages whatsoever on any account in respect of such withholding or retention.

SECTION – V
SPECIAL CONDITIONS OF CONTRACT (SCC)

The following Special Conditions of Contract (SCC) will apply for this purchase. The corresponding clauses of General Conditions of Contract (GCC) relating to the SCC stipulations have also been incorporated below.

These Special Conditions will modify/substitute/supplement the corresponding (GCC) clauses.

Whenever there is any conflict between the provision in the GCC and that in the SCC, the provision contained in the SCC shall prevail.

The warranty conditions, Shelf life, if applicable, will be as mentioned in the Schedule of Requirement as per section VI of the Tender Enquiry Document.

- 1. The quantity shown in the tender can be increased or decreased to any extent depending upon the actual requirement.**

SECTION – VI
SCHEDULE OF REQUIREMENTS
As per “Annexure A”- List of Drugs/Molecules

Terms of Delivery:

Free Delivery at Consignee's Site(s)

1. Delivery Period:

- 1.1 The Delivery Period is maximum 45 days from date of issue of Purchase Order against the Rate Contract. In case of exigency, a shorter Delivery Period can be given and if, it is not acceptable to Supplier, it may be intimated to the Purchase Officer within seven days from the date of issue of the Purchase Order, otherwise it will be assumed that the Purchase Order has been accepted. The date of delivery will be the date by when it is to be delivered at consignee site.
- 1.2 The purchaser will not pay separately for transit insurance and the contractor will be responsible for delivery of items covered by the supply-order in good condition at the specified destination and for this purpose, freight, insurance, octroi etc., if any will have to be borne by the supplier. The consignee will, as soon as possible, but not later than 07 days of the date of arrival of stores at destination, notify the supplier/ bidder, of any loss or damage to the stores that may have occurred in the transit.

2. Shelf-Life:

- a) Short- life items (which have a life-period of eighteen months or less), should not have passed 5/6th of their total shelf life at the time of supply.
- b) In respect of items not covered by clause (i) above, stores should not be older than one year from the date of manufacturing at the time of supply.
- c) For all those drugs, which are required to be stored under controlled temperature / cold chain, have to be supplied under controlled temperature/cold chain.
- d) If the supplied item is not utilized before expiry date the supplier should undertake to replace with fresh stock of items as and when required.
- e) The supplier shall arrange to effect free replacement of any quantity which may deteriorate in potency, strength approaching expiry or expired etc. before the date of expiry marked on the labels.
- f) For Drugs having shelf life of Two years or less: As on the date of delivery, Drugs should not be older than one fourth (1/4) of its shelf life from the date of manufacture.
- g) For Drugs having shelf life more than Two years: As on the date of delivery, Drugs should not be older than one sixth (1/6) of its shelf life from the date of manufacture.
- h) For Imported Drugs: As on the date of delivery, Drugs should have a minimum 50% of valid shelf life from the date of manufacture.

However, the consignee may relax this criteria in case of exigencies with reasons duly recorded and shall be responsible for use of that stores within its given shelf life, with a suitable undertaking from the supplier, the terms of which shall be decided by the consignee as per the requirement of the stores and usage pattern. The Consignee should ensure that there should not be any loss to the Corporation.

3. The supply offered should comply with the provisions of the Drugs and Cosmetics Act, 1940 and the Rules made there under as amended up to date and Drug Price Control order.

(ii) While making quotations against re-packing and chemical items, it must be ensured that ISI code number is indicated on quotation and at the time of making the supplies, the firm should ensure that the item supplied has ISI mark as well as code number, as is the statutory requirement of the Bureau of Indian Standards. The attested copy of the valid ISI marking license issued by Bureau of Indian Standards should be enclosed along with quotation.

If a molecule is being repacked all the requirements of 28 (i) to (xviii) must be fulfilled for the repacked molecule.

For delayed delivery, liquidated damages will get applied as per GCC.

SECTION – VII
SPECIFICATION
As per “Annexure A”- List of Drugs/Molecules

Section – VIII
QUALIFICATION CRITERIA

1. Scanned copy of **Manufacturing & Market standing/ experience certificate** of minimum “**Three Years**” of the molecule quoted by them duly certified by centre/ State Drug Controller in the Performa Section- XVIII. The certificate should have been issued recently i.e. not more than one year old from the date of the opening of the tender.
2. WHO GMP/GMP Certificate Scanned Copy of Valid WHO-GMP certificate/ Valid Schedule ‘M’ certificate issued by Centre/ State Drug Controller and should not have been issued more than five years old.
3. In case of imported drugs (i.e. not manufactured in India), **COPP (Certificate of Pharmaceutical Products)/ import license** and copy of the import registration of that particular molecule quoted in the tender indicating the list of products should be submitted as per WHO norms and ‘3-years’ Marketing experience certificate issued by the Drug Controller.
4. Scanned copy of **valid manufacturing license** issued by Centre/State Drug Controller indicating the list of products should be submitted. Public Sector Undertakings with at least “3-years” market standing having manufacturing license issued by Centre/ State Drug Controller.
5. Scanned copy of **valid narcotic license** issued by Central/State Excise Commissioner should be submitted by the bidder.
6. In case of newly introduced drugs/molecules, the manufacturer can be eligible provided the firm submits a certificate from the DCGI, in this regard. In such cases, the firm has to submit an MMC of the molecule concerned from the date of issue of Certificate by the DCGI of the new drug to that firm. In such case MMC of 03 years is not cleared/ completed, it will be relaxed accordingly. Also, in case of imported Drug/Formulations Form-45 (Permission Certificate) issued by DCGI will also be accepted.

In case of the newly off-patent molecules wherein MMC of 03 years is not cleared/ completed, it will be relaxed in accordance with the time from which the molecule has been declared off-patent.

In case, renewal of MMC is pending, the application for renewal will be read in continuation with the last MMC provided there is no break between the two. However, the renewed MMC will have to be provided by the firm before completion of technical evaluation; failing which the bid will summarily be rejected.

7. Firms which have **US-FDA** approval for export/selling of specified drugs in USA, may submit copies of approval documents from FDA in support of their claim.

8. Manufacturing firm should upload the scanned copy of performance certificate of 02 years for supply of drugs/medicines/iv fluids within last 05 financial years i.e. **2020-21, 2021-22, 2022-23, 2023-24 and 2024-25** from any Govt. Hospital/PSUs./reputed hospital/Institutions/International buyer on the purchaser letter head where the bidders is supplying these items in reference to this tender.
9. **Production-Capacity assessment certificate:** The manufacturing firm should enclose the certificate issued by the Chartered Accountant/ concerned State Drug Controller indicating actual production detail of a particular molecule batch wise for the items quoted and at least one analysis batch report per year for any two of the last three years for each molecule quoted (i.e. minimum of two reports of at least **2-different years** of the last three financial years (**2022-23, 2023-24 and 2024-25**) in the enclosed Performa at **Section-XIX**.
10. Tender shall be rejected if the Copy of GST Registration Certificate is not furnished. Firm shall furnish a certificate on their letter head stating that up to date returns have been filed and there are no dues with the concerned department. Firm will also submit Scanned copies of last 01 (one) year's returns submitted to the concerned department.
11. Turnover Clause:
 - (a) Participating pharmaceutical Firms will have to submit audited financial statement by registered Chartered Accountant for last three preceding financial years (i.e. **2022-23, 2023-24 and 2024-25**) in support of the annual turnover.
 - (b) Group turnover (other than drugs and their formulations) will not be considered for determining the eligibility and such tenders will be rejected summarily.
 - (c) The manufacturing firm quoting for the items mention below have to Submit the documents of annual turnover of the company audited by a Chartered Accountant of the pharmaceutical products during any three consecutive financial years (Financial year **2020-21, 2021-22, 2022-23 2023-24 and 2024-25**) :
 - i) Narcotic drugs, Enemas should have minimum annual turnover of Rs. 1.5 Crores.
 - ii) Niche products/Patented Products/MSE have minimum annual turnover of Rs. 1.5 Crores.
 - iii) Cream/Ointment, lotion, eye/ear drops, mouth wash/Gargles , Contrast media, I.V fluids(large volume parenterals) should have a minimum annual turnover of Rs. 30.00 Crores.
 - iv) Tablets, Capsules, Injections should have a minimum annual turnover of at least Rs. 150.00 Crores
12. If a firm is the sole manufacturer of the product, the same can be treated as a Proprietary drug, provided the firm submits a certificate to this effect from the competent authority in India.

13. Scanned copy of **Non-conviction certificate** issued by the Centre/State Drug Controller to the effect that the manufacturer has not been convicted under the Drugs and Cosmetics Act, 1940 and rules there under during the last three years in respect of any of the drugs for which prices have been quoted by the firm. In case the DCGI does not mention the name of the molecules in their certificates, **a relevant undertaking will be provided with list of drug/molecules along with non-conviction certificate, by the vendor in addition to the above mentioned certificate.** Non-Conviction Certificate must have been issued *by the Drug Controller of the concerned State* within preceding one year from the date of the publication of the tender.

14. In case of imported products the financial turnover of overseas manufacturing firm (Principal firm) will be considered.

15. The contractor should also give a guarantee as follows, in case of biological and other products having a particular life-period to provide safe-guard against loss on account of deterioration within their stated period of potency.

"The seller hereby declares that the goods/store/articles sold to the buyer under this contract shall be of the best quality and shall be strictly in accordance with the specification and particulars mentioned in the description clauses hereof and the seller hereby guarantees that the said goods/stores/articles would continue to confirm to their description and quality for a period of one year from the date of delivery of the said goods/stores/articles or such portion thereof as may be discovered not to conform to the description and quality. Such rejection of the goods/articles/ stores will be at the seller's risk and all the provisions herein contained relating to rejection of goods etc., or such portion thereof if rejected by the purchaser shall be applicable. Otherwise the contractor/seller shall pay to the purchaser such damages as may arise by reason of the breach of conditions herein contained. Nothing herein contained shall prejudice any other right of the purchase in that behalf under this contract or otherwise".

16. Certificate on self-attested non-judicial stamp paper of Rs.10/- stating that there is no vigilance/ CBI case pending against the firm/supplier and the firm has not been blacklisted/debarred on the date of submission of the bid by any Central Govt./State Govt. department/hospital/PSUs etc. Bidder should also provide information regarding blacklisting/debarring of the firm in last three years (**2022-23, 2023-24 and 2024-25**) by any Government or Private organization/Hospital. In case of any false information provided or concealed the information by any bidder, the bidder shall be debarred for two years and EMD/Bid Security/Performance Security submitted by the firm shall be forfeited.

17. The firms should give an undertaking to the effect that they will be legally bound to supply the medicines/drugs, for which they have quoted the rates in the tender during the validity of the contract. In case, they fail to execute any supply-order placed to them within 45 days from the date of placement of purchase order, they will be liable for action against them as per tender terms.
18. Scanned copy of Information as per the format enclosed (**Section-XVII**) should be submitted with the tender. Furnishing of false information will make the bidder ineligible and the firm will stand blacklisted.
19. Scanned copy of List of Items quoted as per **Section- XVI**.
 - a) Participating Pharmaceutical firm should submit a notarized undertaking on an affidavit of Rs. 100/- (Rupees One Hundred only) stating that:
 - i. They will comply with all the statutes & legislation regarding manufacturing, import, sale and supply of drugs in India and in particular the following Acts/Enactments viz., the Drugs and Cosmetics Act, 1940, The Drugs and Cosmetics Rules, 1945 (as amended), The Legal Metrology Act, 2009, The Drugs (Control) Act, 1950, The Indian Statistical Institute Act, 1959, GST Act.
 - ii. To supply drugs of standard quality as prescribed under the provisions of Drug and Cosmetic Act, 1940 (as amended). The bidder shall also undertake not to supply items / drugs “not of standard”, “Grossly sub-standard” and “Spurious and adulterated drugs” as per the guidelines issued by the Drug Controller of India from time to time.
 - b) The participating pharmaceutical firm should submit an affidavit of Rs. 100/- (Rupees One Hundred only) duly signed by the Notary (Annexure T) asunder:-
 - i. “The pharmaceutical firm hereby declare that the drugs/items sold to the AIIMS under this contract shall be of best quality and workmanship and shall be strictly in accordance with the specifications and particulars contained/mentioned in the description clauses hereof and the pharmaceutical firm/bidder hereby guarantees that the said drugs/items would continue to conforms to their description/ specification and the provisions of law as stated in the contract and that notwithstanding the fact that the purchaser (inspector) may have inspected and/or approved the said drugs/items. If the same be discovered not to conform to the description and quality aforesaid or have deteriorated, the decision of the AIIMS in that behalf will be final and conclusive. AIIMS will be entitled to reject said drugs/items or such portion thereof as may be discovered not to conform to the said description and quality in the manner as prescribed. Such rejection of the drugs/items will be at the seller's risk and all the provisions herein contained relating to rejection of drugs/items etc. or such portion thereof if is rejected by the purchaser. Nothing herein contained shall prejudice any other right of AIIMS in that behalf under this contract or otherwise”.

- ii. The Bidder submits stating that the drugs, which are being quoted, are not banned under Section 26 (A) of Drugs & Cosmetics Act. Or any other provision of law prevailing in India.
- iii. It is declared that the firm / company/ corporation and any of its director / proprietors/ partners/ Authorised signatories are not convicted/ or a criminal case filed against or pending in any court of India by any department of Govt. under prevention of Corruption Act or for cheating/ defrauding Govt/ embezzlement of Govt fund or any criminal conspiracy in the said matter.
- iv. The Bidder submits an undertaking that it is not submitting bid for any drug/ combination of drugs which is not approved by DCGI".
- v. Company/Authorised Signatory has to submit an affidavit giving address of manufacturing unit.

20. For the drugs which are being imported, the Participating Pharmaceutical firm will submit valid import license issued by Drug Controller General of India and valid marketing license issued by concerned Licensing Authority (Form 10 & Form 41). That Firm will be eligible if one batch of new drug has been imported at the time of bidding.

21. In case of patented drugs, Participating Pharmaceutical firm will submit valid certificate to this effect from the Licensing Authority else bidder's claim will not be considered.

22. The firm / company/ corporation should not be convicted/ or a criminal case filed against or pending in any court of India by any department of Govt. under prevention of Corruption Act or for cheating/ defrauding Govt./ embezzlement of Govt. fund or any criminal conspiracy in the said matter.

23. For the drugs quoted in the tender enquiry, Participating Pharmaceutical firm will have to submit the samples on demand. If bidder fails to submit the samples within the period specified, the tender will be rejected

Section – IX
TENDER ACCEPTANCE FORM

To _____

The Director,
All India Institute of Medical Sciences
Ansari Nagar, New Delhi-110 029 India.

Ref. Your ATE No. _____ due for opening on

_____ *insert date*

We, the undersigned have examined the above mentioned Tender Enquiry Document, including amendment/corrigendum (*if any*), the receipt of which is hereby confirmed. We now offer to supply and deliver in conformity with your above referred document for the sum as shown in the Price Schedules (BoQ) uploaded herewith and made part of this bid. If our bid is accepted, we undertake to supply the items for which Rate Contract has been concluded, in accordance with the delivery schedule specified in the Schedule of Requirements.

We further confirm that, if our bid is accepted, we shall provide you with a Performance Security of required amount in an acceptable form in terms of “General Conditions Contract”, Section - IV read with modification, if any “Special Conditions of Contract”, in Section - V, for due performance of the Rate Contract/Purchase Orders.

We agree to keep our bid valid for acceptance as required in the “General Instruction to Bidders”, read with modification, if any in “Special Instructions to Bidders”, Section – III or for subsequently extended period, if any, agreed to by us. We also accordingly confirm to abide by this bid up to the aforesaid period and this bid may be accepted any time before the expiry of the aforesaid period. We further confirm that, until a formal Rate Contract is executed, this bid read with your written acceptance thereof within the aforesaid period shall constitute a binding contract between us.

We further understand that you are not bound to accept the lowest or any bid you may receive against your above-referred advertised tender enquiry.

We confirm that we do not stand deregistered/banned/blacklisted by Central Govt./State Govt. Ministries/AIIMS, New Delhi.

We confirm that we fully agree to the terms and conditions specified in above mentioned Tender Enquiry Document, including amendment/ corrigendum if any.

We hereby certify that if at any time, information furnished by us is proved to be false or incorrect, we are liable for any action as deemed fit by the purchaser in addition to forfeiture of the Bid Security/Performance Security.”

Name: _____
Business Address _____

Place: _____

Date: _____

SECTION – X
PRICE SCHEDULE

BoQ may be uploaded as per instructions given in Tender Enquiry Document.

SECTION - XI
BANK GUARANTEE FORM FOR BID SECURITY

Whereas _____ (Name and address of the Bidder)
(hereinafter called the "Bidders")

has submitted its Bid dated _____ for the supply of _____
(hereinafter called the "Bid")

against the purchaser's ATE No. _____

Know all persons by these presents that we _____

having our registered office at _____
(Hereinafter called the "Bank")

are bound to AIIMS MAIN GRANT, New Delhi
(hereinafter called the "Purchaser")

in the sum of _____ for which payment will and truly to be made to the said Purchaser, the Bank binds itself, its successors and assigns by these presents. Sealed with the Common Seal of the said Bank this

_____ day of _____ 20 _____.

The conditions of this obligation are:

- 1) If the Bidder withdraws or amends, impairs or derogates from the bid in any respect within the period of validity of this Bid.
- 2) If the Bidder having been notified of the acceptance of his Bid by the Purchaser during the period of its validity:-
 - a. If the bidder fails or refuses to furnish the performance security for the due performance of the Rate Contract/Purchase Orders or
 - b. If the bidder fails or refuses to accept/execute the Rate Contract/Purchase Orders or
 - c. If it comes to notice at any time, that the information/documents furnished in its Bid are false or incorrect or misleading or forged

We undertake to pay the Purchaser up to the above amount upon receipt of its first written demand, without the Purchaser having to substantiate its demand, provided that in its demand the Purchaser will note that the amount claimed by it is due to it owing to the occurrence of one or more the three conditions, specifying the occurred condition(s).

This guarantee will remain in force up to _____ (insert date of additional forty-five days after Bid validity) and any demand in respect thereof should reach the Bank not later than the above date.

.....
(Signature with date of the authorized officer of the Bank)

.....
(Name and designation of the Officer)

.....
(Seal, name & address of the Bank and address of the Branch)

SECTION - XII
BANK GUARANTEE FORM FOR PERFORMANCE SECURITY

WHEREAS _____ (Name and address of the Supplier) (Hereinafter called "the Supplier")

has undertaken, in pursuance of Rate Contract No._____

dated _____ valid from _____ to _____ for supply

_____ (*insert description of goods*)
(Hereinafter called "the Contract"),

to AIIMS MAIN GRANT, New Delhi-110 029
(Hereinafter called "the Purchaser")

AND WHEREAS it has been stipulated by you in the said contract that the supplier shall furnish you with a bank guarantee by a scheduled commercial bank recognized by you for the sum specified therein as security for compliance with its obligations in accordance with the contract;

AND WHEREAS we have agreed to give the supplier such a bank guarantee;

NOW THEREFORE we hereby affirm that we are guarantors and responsible to you, on behalf of the supplier, up to a total of

_____ (*insert Amount of the Performance Security in words and figures*), and we undertake to pay you, upon your first written demand declaring the supplier to be in default under the contract and without cavil or argument, any sum or sums within the limits of (amount of guarantee) as aforesaid, without your needing to prove or to show grounds or reasons for your demand or the sum specified therein.

We hereby waive the necessity of your demanding the said debt from the supplier before presenting us with the demand.

We further agree that no change or addition to or other modification of the terms of the contract to be performed there under or of any of the contract documents which may be made between you and the supplier shall in any way release us from any liability under this guarantee and we hereby waive notice of any such change, addition or modification.

This guarantee will remain in force up to _____ (*insert last date of currency of Rate Contract plus Warranty Period (if applicable) plus additional Ninety days*) and any demand in respect thereof should reach the Bank not later than the above date.

.....
(Signature with date of the authorized officer of the Bank)

.....
Name and designation of the officer

.....
Seal, name & address of the Bank and address of the Branch

SECTION – XIII
RATE CONTRACT FORM FOR GOODS
(To be executed on Non-Judicial Stamp Paper worth of Rs. 100/-)

ALL INDIA INSTITUTE OF MEDICAL SCIENCES
(Insert Name of concerned Centre/Hospital/Department/Section)
ANSARI NAGAR, NEW DELHI-110 029

Rate Contract No._____ dated_____

To _____
(insert name of Supplier with address)

This is in continuation to this office's Notification of Award No.: _____ dated _____

1. Name & address of the Supplier: _____
2. Advertised Tender Enquiry No. of Tender Documents: _____
and subsequent Amendment No.: _____, dated: _____
(if any), issued by the Purchaser
3. Supplier's Bid No.: _____ dated: _____ and subsequent communication(s) No.: _____ dated: _____ (if any), exchanged between the supplier and the purchaser in connection with this Tender Document.
4. In addition to this Rate Contract Form, the following documents etc, which are included in the Tender Enquiry Documents mentioned under paragraphs 2 and 3 above, shall also be deemed to form and be read and construed as integral part of this Rate Contract:

- i) General Conditions of Contract;
- ii) Special Conditions of Contract;
- iii) Schedule of Requirements;
- iv) Technical Specifications;
- v) Tender Acceptance Form uploaded by the supplier;
- vi) Price Schedule(s)/BoQ uploaded by the supplier in its Bid;
- vii) Manufacturers' Authorization Form (if applicable);
- viii) Purchaser's Notification of Award

Note: The words and expressions used in this Rate Contract shall have the same meanings as are respectively assigned to them in the conditions of Rate Contract referred to above. Further, the definitions and abbreviations incorporated under clause 1 of Section II – "General Instructions to Bidders" of the Tender Enquiry Document shall also apply to this Rate Contract.

5. Some terms, conditions, stipulations etc. out of the above-referred documents are reproduced below for ready reference:

- i) Brief particulars of the goods which shall be supplied by the supplier against Rate Contract are as under:

Item	Brief Description	Unit	Unit	GST Rate	Total Unit
------	-------------------	------	------	----------	------------

No.	of Goods		Price (in INR)	(in %age)	Price with GST (in INR)

ii) Terms of Delivery: Free Delivery At Site
 iii) Delivery schedule: 45 Days from the Date of Issue of Purchase Order
 iv) Performance Security of Rs._____ valid upto _____ to be furnished by _____

6. Currency of Rate Contract from: _____ to: _____
 7. Shelf Life: At the time of supply, the supplier will supply fresh stock, and the remaining shelf life should be more than 5/6 of shelf life.
 8. The supplier shall arrange to effect free replacement of any quantity which may deteriorate in potency, strength etc. before the date of expiry marked on the labels.
 9. Payment terms: As per General Conditions of Contract
 10. The Supplier will supply the goods as per Rate Contract against Purchase Orders issued by various Centers/Hospital/Section/Departments/Store Sections of AIIMS, New Delhi.

Signature, name and designation of the Purchaser authorized official for and on behalf of Director, AIIMS, may be called as First Party

Received and accepted this Rate Contract

Signature, name and address of the supplier's executive duly authorized to sign on behalf of the supplier, may be called as Second Party

for and on behalf of _____
(Insert Name and address of the supplier)

(Seal of the Supplier)

Date: _____

Place: _____

SECTION – XIV

CONSIGNEE RECEIPT CERTIFICATE

(To be given by consignee's authorized representative)

The following store(s) has/have been received in good condition:

- 1) Rate Contract No. &date : _____
- 2) Purchase Order No. &date : _____
- 3) Supplier's Name : _____
- 4) Consignee's Name & Address: _____
- 5) Name of the item supplied : _____
- 6) Quantity Supplied : _____
- 7) Date of Receipt by the Consignee : _____

Signature of Consignee with date: _____

Name and designation of Consignee: _____

Seal of the Consignee: _____

SECTION – XV
FINAL CONSIGNEE ACCEPTANCE CERTIFICATE
(To be given by consignee's authorized representative)

1 This is to certify that the goods as detailed below have been received in good conditions along with all the standard and special accessories in accordance with the Rate Contract/Purchase Order and the same has been installed and accepted.

- 1) Rate Contract No. & date : _____
- 2) Purchase Order No. & date : _____
- 3) Supplier's Name: _____
- 4) Consignee's Name & Address: _____
- 5) Name of the item Supplied : _____
- 6) Quantity Supplied : _____
- 7) Date of Receipt by the Consignee : _____
- 8) Quantity Accepted : _____
- 9) Date of Acceptance by the Consignee : _____
- 10) The supplier has fulfilled its contractual obligations including installation (if applicable) satisfactorily

OR

The supplier has failed to fulfill its contractual obligations with regard to the following:

- i)
- ii)
- iii)
- iv)
- 11) The amount of recovery on account of failure of the supplier to meet his contractual obligations is _____ (here indicate the amount).

Signature of Consignee with date: _____

Name and designation of Consignee: _____

Seal of the Consignee: _____

SECTION – XVI

LIST OF ITEMS QUOTED

FORMAT OF SUBMISSION OF VALID REVISED SCHEDULE -M/ WHO-GMP/IMPORT LICENSE/ COPP/ MANUFACTURING LICENSE (STRICT COMPLIANCE).

Sr. No.	Item' serial no. as per tender list	Name of Drugs	Page no. Tender where valid WHO-GMP/ Revised Schedule M/ import license/ COPP/Public Sector undertakings enclosed	Page no. Tender where valid Manufacturing License/ Import license enclosed.

Strict Compliance: - All the bidders are directed to mention the page number of the tender document where WHO-GMP/ Revised Schedule 'M' & page number of manufacturing license for indigenous drugs / import license for imported drugs enclosed. Merely mentioning the word '**Enclosed**' may lead to rejection of tender / bid. Submission

- a) Participating Pharmaceutical firm should submit a notarized undertaking on an affidavit of Rs. 100/- (Rupees One Hundred only) stating that:
 - i) They will comply with all the statutes & legislation regarding manufacturing, import, sale and supply of drugs in India and in particular the following Acts/Enactments viz., the Drugs and Cosmetics Act, 1940, The Drugs and Cosmetics Rules, 1945 (as amended), The Legal Metrology Act, 2009, The Drugs (Control) Act, 1950, The Indian Statistical Institute Act, 1959, GST Act.
 - ii) To supply drugs of standard quality as prescribed under the provisions of Drug and Cosmetic Act, 1940 (as amended). The bidder shall also undertake not to supply items / drugs "not of standard", "Grossly sub-standard" and "Spurious and adulterated drugs" as per the guidelines issued by the Drug Controller of India from time to time.
- b) The participating pharmaceutical firm should submit an affidavit of Rs. 100/- (Rupees One Hundred only) duly signed by the Notary (Annexure T) as under:-
 - i) "The pharmaceutical firm hereby declare that the drugs/items sold to the AIIMS under this contract shall be of best quality and workmanship and shall be strictly in accordance with the specifications and particulars

contained/mentioned in the description clauses hereof and the pharmaceutical firm/bidder hereby guarantees that the said drugs/items would continue to conform to their description/ specification and the provisions of law as stated in the contract and that notwithstanding the fact that the purchaser (inspector) may have inspected and/or approved the said drugs/items. If the same be discovered not to conform to the description and quality aforesaid or have deteriorated, the decision of the AIIMS in that behalf will be final and conclusive. AIIMS will be entitled to reject said drugs/items or such portion thereof as may be discovered not to conform to the said description and quality in the manner as prescribed. Such rejection of the drugs/items will be at the seller's risk and all the provisions herein contained relating to rejection of drugs/items etc. or such portion thereof if is rejected by the purchaser. Nothing herein contained shall prejudice any other right of AIIMS in that behalf under this contract or otherwise".

- ii) The Bidder submits stating that the drugs, which are being quoted, are not banned under Section 26 (A) of Drugs & Cosmetics Act. Or any other provision of law prevailing in India.
- iii) It is declared that the firm / company/ corporation and any of its director / proprietors/ partners/ Authorised signatories are not convicted/ or a criminal case filed against or pending in any court of India by any department of Govt. under prevention of Corruption Act or for cheating/ defrauding Govt/ embezzlement of Govt fund or any criminal conspiracy in the said matter.
- iv) The Bidder submits an undertaking that it is not submitting bid for any drug/ combination of drugs which is not approved by DCGI".
- v) Company/Authorised Signatory has to submit an affidavit giving address of manufacturing unit

SIGNATURE AND ADDRESS OF THE BIDDER

SECTION – XVII

PROFORMA TO BE FILLED BY THE TENDERER

I. GENERAL INFORMATION

- a)** Name of the firm :
- b)** Address & Telephone No. :
- c)** Whether the firm is Indian / Multi- national :
- d)** Whether Small / Medium/Large Scale Co. :
- e)** Person responsible for conduct of Business :
- f)** Particulars of Licenses held under Drugs & Cosmetic Act & the details. (If the license is under renewal, certificate from the Drug Controller that the license is under renewal and deemed to be enforced) :
- g)** Procurement agency with which registered and the agencies to whom drugs supplied during last one year :
- h)** Has the firm been convicted ever, if yes, give details:
 - i)** Any case pending in the Court with details:
 - j)** Has the firm ever been debarred / black-listed by any Govt. Hospital for poor quality or late supply of drugs? If yes, give details.
- k)** Fax No :
- l)** E- Mail Address :
- m)** Name & Mobile No of person/ authorized signatory to be contacted for this tender :

II. TECHNICAL

- a)** Equipments for material handling, manufacturing of drugs and quality-control of drugs :
- b)** Specialized testing facilities such as microbiological testing and Biological testing:
- c)** Details of Technical Staff
 - i)** Manufacturing Staff :
 - ii)** Quality Control Staff :

- d) Has the firm carried out stability study for drugs quoted : _____
- e) Is the firm basic manufacturer of the drug quoted, if yes, details : _____
- f) Has the firm following
 - i) WHO GMP Certificate /Schedule-M : _____
 - ii) ISO Certificate : _____
 - iii) FDA Certificate : _____
 - iv) Import License : _____
- g) Installed capacity and actual production details for different forms of drugs:
 - i) Tablets : _____
 - ii) Capsules : _____
 - iii) Syrups/ Suspension : _____
 - iv) Injections : _____
 - v) Powder : _____
 - vi) Inhalation : _____
 - vii) Topical : _____
- h) Drugs declared and sub-standard / re-called during the last three years. Give details with reasons and the remedial action taken : _____

III. FINANCIAL

- a) Turnover during last three financial years (year wise) of the pharmaceutical products. Firms should furnish copies of audited Balance-sheet / Sales Tax clearance certificate.
- b) Name & Address of the Bankers to the Firm and the facilities available from the bank.
- c) Income-tax No./ Central Sales-tax No./ State Sales-tax No.

DECLARATION

I, _____ Proprietor/Partner/Director of M/s _____ hereby declare that the information given in this form is true and correct to the best of my knowledge and belief.

(Signature)

(Name & Designation with Stamp)

WARNING: If the information furnished in this form is found to be incorrect at any point of time, the bidder may be debarred.

SECTION – XVIII
MANUFACTURING & MARKETING CERTIFICATE

This is to certify that M/s _____ are holding valid Manufacturing license No. _____ dated _____ of the _____ State and they are manufacturing and marketing, the following products for last three (3) years.

The products are as follows:

S. No.	Name of the Product	Pharmacopoeia Specification	Strength
1.			
2.			
3.			
4.			
5.			
6.			
7.			
8.			
9.			
10.			

**Signature and seal of
Drug Controller of
the Centre/State.**

Dated:

Note: This certificate is to be signed by the Drug Controller of **Centre/State**. Certificate issued by Inspector of Drugs will not be accepted unless an authorization by the concerned centre/State Drug Controller to this effect is supported by adequate documentary proof.

SECTION – XIX

PRODUCTION-CAPACITY ASSESSMENT CERTIFICATE

Item no. & name of items: _____

Indicate details of production of the items quoted at least two years from **2022-23** **2023-24** and **2024-25** duly certified by the **Chartered Accountant/ Centre/State Drug Controller.**

S. No. of the item as in Tender Enquiry	Name & Specification of the item	Date of issue of Mfg. License for the product	Date of marketing the 1st batch
1.	2.	3.	4.

2022-23		2023-24		2024-25		REMARKS
Batch No.	Size	Batch No.	Size	Batch No.	Size	

Signature of the Manufacturer:

**Signature of the Chartered Accountant/
Centre/State Drug Controller along with address & Seal**

SECTION - XX
CHECKLIST

Sr. No.	Documents to be submitted alongwith the techno-commercial bid	Attached at page number
a.	Scanned copy of “EMD/Bid Security” furnished in accordance with GIB alternatively, documentary evidence as per GIT for claiming exemption from payment of EMD/Bid security to be uploaded.	
b.	Scanned copy of “List of Items Quoted” as per SECTION - XVI of Tender Enquiry Document.	
c.	Scanned copy of “Tender Acceptance Form” as per Section IX to be uploaded	
d.	Scanned Copy of GST Registration Certificate. Copy of GST No dues Certificate Scanned copies of last 2 year's GST returns submitted to the concerned department	
e.	Scanned copy of Documents confirming to Sole Proprietorship/ Partnership/Private Limited Firm in the country of origin as the case may be to be uploaded.	
f.	Scanned copy of Manufacturing & Market standing/ experience certificate of minimum “Three Years” of the molecule quoted by them duly certified by centre/ State Drug Controller in the Performa Section- XVIII . The certificate should have been issued recently i.e. not more than one year old from the date of the opening of the tender.	
g.	Scanned Copy of Valid WHO-GMP certificate/ Valid Schedule ‘M’ certificate clearly indicating the products (molecule/drug) issued by Centre/ State Drug Controller and should not have been issued more than five years old.	
h.	<p>In case of imported drugs (i.e. not manufactured in India), COPP (Certificate of Pharmaceutical Products)/ import license and copy of the import registration of that particular molecule quoted in the tender indicating the list of products should be submitted as per WHO norms and ‘3-years’ Marketing experience certificate issued by the Drug Controller.</p> <p>In case of the newly off-patent molecules wherein MMC of 03 years is not cleared/ completed, it will be relaxed in accordance with the time from which the molecule has been declared off-patent.</p> <p>In case, renewal of MMC is pending, the application for renewal will be read in continuation with the last MMC provided there is no break between the two. However, the renewed MMC will have to be provided by the firm before completion of technical evaluation; failing which the bid will summarily be rejected.</p>	

i.	Scanned copy of valid manufacturing license issued by Centre/State Drug Controller indicating the list of products should be submitted. Public Sector Undertakings with at least “3-years” market standing having manufacturing license issued by Centre/ State Drug Controller.	
j.	In case of newly introduced drugs/molecules, the manufacturer can be eligible provided the firm submits a certificate from the DCGI, in this regard. In such cases, the firm has to submit an MMC of the molecule concerned from the date of issue of Certificate by the DCGI of the new drug to that firm. In such case MMC of 03 years is not cleared/ completed, it will be relaxed accordingly.	
k.	Manufacturing firms should submit scanned copy of performance certificate(s) of at least 02 years in last 05 years (2020-21, 2021-22, 2022-23, 2023-24 and 2024-25) , from other similar two Hospital, out of which one must be from Government/Public Sector from the Competent Authority.	
l.	Production-Capacity assessment certificate as per section-XIX	
m.	<p>Turnover:</p> <p>(a) Participating pharmaceutical Firms will have to submit audited financial statement by registered Chartered Accountant for last three preceding financial years (i.e.2022-23, 2023-24 and 2024-25) in support of the annual turnover.</p> <p>(b) Group turnover (other than drugs and their formulations) will not be considered for determining the eligibility and such tenders will be rejected summarily.</p> <p>(c) The manufacturing firm quoting for the items mention below have to Submit the documents of annual turnover of the company audited by a Chartered Accountant of the pharmaceutical products during any three consecutive financial years (Financial year 2020-21, 2021-22, 2022-23, 2023-24 and 2024-25) :</p> <p>i) Narcotic drugs, Enemas should have minimum annual turnover of Rs. 1.5 Crores.</p> <p>Niche products/Patented Products/MSE have minimum annual turnover of Rs. 1.5 Crores.</p> <p>ii) Cream/Ointment, lotion, eye/ear drops, mouth wash/Gargles , Contrast media, I.V fluids(large volume parenterals) should have a minimum annual turnover of Rs. 30.00 Crores.</p> <p>iii)Tablets, Capsules, Injections should have a minimum annual turnover of at least Rs. 150.00 Crore</p>	

n.	Scanned copy of Non-conviction certificate	
o.	Certificate on self attested non-judicial stamp paper of Rs.10/- stating that there is no vigilance/ CBI case pending against the firm/supplier and the firm has not been blacklisted/debarred on the date of submission of the bid by any Central Govt./State Govt. department/hospital/PSUs etc. Bidder should also provide information regarding blacklisting/debarring of the firm in last three years (2022-23, 2023-24 and 2024-25) by any Government or Private organization/Hospital. In case of any false information provided or concealed the information by any bidder, the bidder shall be debarred for two years and EMD/Bid Security/Performance Security submitted by the firm shall be forfeited.	
p.	The firms should give an undertaking to the effect that they will be legally bound to supply the medicines/drugs, for which they have quoted the rates in the tender during the validity of the contract. In case, they fail to execute any supply-order placed to them within 45 days from the date of placement of purchase order, they will be liable for action against them as per tender terms.	
q.	Scanned copy of Information as per the format enclosed (Section-XVII) should be submitted with the tender. Furnishing of false information will make the bidder ineligible and the firm will stand blacklisted.	
r.	At least one analysis batch report per year for any two of the last three years for each molecule quoted (i.e. minimum of two reports of at least 2-different years of the last three financial years (2022-23, 2023-24 and 2024-25) in the enclosed Performa at Section-XIX .	
s.	Name and address of the authorized distributor for supplying against the subject tender along with the OEM authorization certificate to be uploaded.	

ANNEXURE
List of Drugs/Medicines

ANNEXURE

S.no.	Item Name	Annual Consumption
1	Tablet Linezolid 600mg	13000
2	Tablet Ofloxacin + Ornidazole 200mg+500mg	11000
3	Injection Amoxycillin+Clavulanic Acid 1.2gm	500000
4	Injection Azithromycin 500mg	85905
5	Injection Clindamycin 600 mg 4ml	139851
6	Injection Teicoplanin 400mg	275765
7	Injection Doxorubicin 10mg	2632
8	Injection Doxorubicin 50mg	3051
9	Tablet Moxifloxacin 400mg	2000
10	Tablet Clarithromycin 250mg	15663
11	Injection Cefuroxime 750 mg	333630
12	Injection Colistimethate Sod. 10,00,000 IU/10ml	20000
13	Injection Colistimethate Sod. 20,00,000 IU	12000
14	Injection Vancomycin 500mg	201526
15	Injection Cefoperazone + Sulbactam 1gm	960746
16	Injection Meropenam 500mg	212959
17	Tablet Amoxycillin + Clavulanic Acid 875+ 125mg	As & when required
18	Tablet Azithromycin 500mg	513763
19	Tablet Cefuroxime 250 mg	As & when required
20	Tablet Cefuroxime 500 mg	7096
21	Injection Piperacillin + Tazobactum 4.5 gm	100000
22	Injection Ceftriaxone Sodium 1gm	604460
23	Injection Impenem + Cilastatin 500 mg	As & when required
24	Injection Tigecycline 50 mg	128651
25	Injection Polymyxin B 500000 units	86994
26	Injection Levofloxacin 500mg 100ml	135056
27	Injection Moxifloxacin 400mg 100ml	As & when required
28	Injection Minocycline 100mg	28805
29	Capsule Cyclosporine 25mg	1250
30	Injection Doxorubicin RTU 10mg	2632
31	Injection Doxorubicin RTU 50mg	3051
32	Capsule Rifabutin 150mg	100
33	Bottle Amoxycillin + Clavulanic Acid (200mg + 28.5mg/5ml) 30ml	23303
34	Syp Ofloxacin 60ml	143
35	Tablet Levofloxacin 750mg	10000
36	Injection Daunorubicin 20mg	10068
37	Injection Daptomycin 350mg	3583
38	Injection Tobramycin 80mg 2ml	431
39	Injection Ceftazidime 1gm	8422
40	Injection Doripenem 500mg	As & when required

41	Injection Meropenam + Sulbactum 1.5gm	570
42	Injection Linezolid 200mg 100ml	89532
43	Injection Cefotazime 1gm	26000
44	Capsule Amoxycillin 500mg	69289
45	Capsule Amoxycillin 250mg	25484
46	Tablet Ciprofloxacin 500mg	137611
47	Tablet Levofloxacin 500mg	182631
48	Tablet Cefixime 100mg	10000
49	Injection Ceftriaxone + Disodium Eddate + Salbactum 3.0gm	34
50	Injection Ceftriaxone + Disodium Eddate + Salbactum 1.5gm	14
51	Injection Aztreonam 1gm	156526
52	Injection Ciprofloxacin I.V 200mg 100ml	As & when required
53	Injection Meropenam 1gm	570699
54	Injection Docetaxel 20mg. 0.5ml	500
55	Tablet Anastrazole 1mg	44277
56	Injection Zoledronic acid 4mg	1832
57	Injection Bortezomib 2mg	2695
58	Injection Bevacizumab 100 mg	700
59	Injection Bevacizumab 400 mg	14
60	Injection Trastuzumab 440 mg	50
61	Tablet Bicalutamide 50 mg	2800
62	Injection Filgrastim 300 mcg	5000
63	Injection Pemetrexed Disodium 100 mg	1500
64	Injection Decitabine 50mg	36
65	Tablet Erlotinib 150mg	240
66	Tablet Imatinib 400mg	3100
67	Injection Nano-Albumin Bound Paclitaxel 100mg	148
68	Tablet Pazopanib 200mg	700
69	Tablet Sunitinib 12.5mg	As & when required
70	Tablet Sunitinib 25mg	As & when required
71	Tablet Sunitinib 50mg	As & when required
72	Tablet Imatinib 400mg	10278
73	Injection Durvalumab 120mg	50
74	Injection Durvalumab 500mg	100
75	Tablet Dasatanib 50mg	6192
76	Tablet Everolimus 5mg	280
77	Tablet Everolimus 10mg	60
78	Tablet Lapatinib 250mg	1530
79	Tablet Lenalidomide 250mg	As & when required
80	Capsule Nintedanib 150mg	13076
81	Tablet Thalidomide 50mg	920
82	Tablet Thalidomide 100mg	1722
83	Tablet Abiraterone Acetate 500mg	As & when required
84	Injection Leuprolide Acetate 22.5mg	196
85	Capsule Everolimus 0.25mg	142

86	Capsule Everolimus 0.50mg	1334
87	Capsule Ibrutinib 140mg	720
88	Tablet Abiraterone Acetate 250mg	As & when required
89	Injection Nimotuzumab 50mg	As & when required
90	Injection Eribulin 1mg	As & when required
91	Injection L-Asparaginase 5000IU	40
92	Injection Ifosfamide 1gm	100
93	Injection Cyclophosphamide 200mg	300
94	Injection Irinotecan 100mg	50
95	Injection Busulfan 60mg	20
96	Injection Cytarabine 1000mg	1800
97	Injection Carfilzomib 60mg	170
98	Injection Bleomycin 15mg	450
99	Injection Epirubicin 10mg	100
100	Tablet Hydroxyurea 500mg	10538
101	Tablet Tamoxifen 10mg	620
102	Injection Cisplatin 50mg	500
103	Injection Obinutuzumab 1000 mg	300
104	Capsule Nilotinib 150mg	4048
105	Capsule Nilotinib 200mg	814
106	Tablet Ruxolitinib 5mg	2220
107	Tablet Ruxolitinib 20mg	67
108	Injection Leucovorin Calcium Folinate 50mg	400
109	Injection Leucovorin Calcium Folinate 15mg	300
110	Tablet Leucovorin Calcium Folinate 15mg	240
111	Tablet Gefitinib 250mg	7
112	Capsule Palbociclib 100mg	84
113	Tablet Crizotinib 250mg	339
114	Injection Trastuzumab Emtansine 100mg	2
115	Injection Trastuzumab Emtansine 160mg	2
116	Injection Gadoteric Acid 10ml	50
117	Injection Gadoteric Acid 20ml	As & when required
118	Injection Iopromide 370mg 50ml	100
119	Injection Gadobutrol (1.0mmol eq. to 6047.72mg gadobutrol) 5ml	50
120	Injection Iopromide 370mg 100ml	2190
121	Injection Iopromide 300mg/ml (50ml)	50
122	Injection Diatrizoate Sod 41.7% w/v 30ml	2
123	Injection Gadopentetate dimeglumine 469mg + 78.6mg (20ml)	As & when required
124	Injection Iopamidol 300mg/ml (50ml)	100
125	Injection Iohexol 300mg/ml (50ml)	6000
126	Injection Iodized Oil 480mg/ml (10ml)	As & when required
127	Injection Gadobenate dimeglumine 334mg + 195mg/ml (15ml)	As & when required
128	Injection Iomeprol 400mg/ml (100ml)	600
129	Injection Diatrizoate Sod 41.7%w/v 100ml	40
130	Injection Gadodiamide 20ml	20

131	Injection Iohexol 300mg/ml (100ml)	4500
132	Injection Iopamidol 370mg/ml (50ml)	80
133	Injection Iopamidol 370mg/ml (100ml)	As & when required
134	Injection Sodium Diatrizoate & Meglumine Diatrizoate 76% pack of 20ml	1050
135	Injection Iodixanol USP 320mg/50ml	25000
136	Injection Atracurium 50mg 5ml	As & when required
137	Injection Propofol 200mg 20ml	35000
138	Injection Glycopyrrolate 0.2mg 1ml	75000
139	Injection Desflurane 240ml	220
140	Injection Rocuronium Bromide 50mg/5ml	1700
141	Injection Bupivacaine 0.5% hyperbaric (Heavy) 0.5% 4ml	7000
142	Injection Dexmedetomidine HCL 200mcg 2ml	2000
143	Injection Etomidate 10ml 2mg/ml 10ml	6000
144	Jelly Lubricating Jelly 20gm	As & when required
145	Jelly Lubricating Jelly 50mg	7151
146	Injection Lignocaine 2% 30ml	25000
147	Injection Lignocaine HCL + Adrenaline 21.3mg/ml + 0.005mg/ml (30ml)	As & when required
148	Jelly Lignocaine Jelly 2% 30gm	90000
149	Spray Lignocaine Throat spray 10% 50ml	As & when required
150	Cream Lignocaine + Prilocaine (dermal anaesthetic cream) 2.5% + 2.5% (30gm)	1172
151	Injection Neostigmine 5ml 2.5mg	40000
152	Injection Neostigmine 1ml 0.5mg	As & when required
153	Injection Ropivacaine HCL 0.75% 20ml	800
154	Injection Thiopentone 1000mg	500
155	Injection Levobupivacaine 0.50%	305
156	Injection Isoflurane 250ml	400
157	Injection Isoflurane 100ml	1600
158	Injection Vecuronium Bromide 4mg/2ml	7500
159	Injection Bupivacaine HCL Preservative free 0.50% 20ml. Amp.	6500
160	Lignocaine Lozenges	As & when required
161	Injection Sevoflurane 99.97% 250ml	500
162	Injection Ketamine I.V 100mg 2ml	16000
163	Injection Inactivated Influenza Quadrivalent 0.5ml	400
164	Injection Botulinum Toxin Type A 50U	As & when required
165	Injection Botulinum Toxin 100U	90
166	Injection Diphtheria Antitoxin 1000 IU/ml (10ml)	170
167	Injection Rabies Antiserum 1500IU 1500 IU	500
168	Injection Hepatitis B Immunoglobulin 100IU	160
169	Injection Human IV Gamma globulins 50ml 2.5gm	As & when required
170	Injection Balanced Solution of low Chlorites 500ml	120000
171	Two Port Close system container DNS (Dex + Sod. Chloride) 0.45% 500ml	As & when required
172	Two Port Close system container Compound Sodium Lactate 500ml	As & when required
173	Two Port Close system container Dextrose 5% 500ml	10000
174	Two Port Close system container Multiple Electrolytes & Dextrose Type 3 (500ml)	As & when required

175	Plastic Normal Saline (Sod. Chloride) (FFS) 0.9% 100ml	300000
176	Plastic Ringer Lactate Solution (Compound Sod. Lactate Injection)(FFS) 500ml	120000
177	Two Port Close system container Sodium Chloride 0.9% w/v 500ml	260000
178	Two Port Close system container Sodium Chloride 0.9% w/v 100ml	300000
179	Two Port Close system container Sodium Chloride + Dextrose 0.9% w/v + 5% w/v	As & when required
180	Plastic P.D fluid Low calcium (CAPD Bag)	As & when required
181	IV Fluid PD Fluid 1.5% Dextrose 5 Ltr	1000
182	IV Fluid PD Fluid 2.5% Dextrose 5 Ltr	1000
183	I.V. bag Normal Saline (Sod. Chloride) (self-collapsible bag without requiring any airway) 0.9% 100ml	As & when required
184	I.V. bag Normal Saline (Sodium Chloride) double pack Non-PVC bag 0.9% 500ml	10000
185	I.V. bag Ringer Lactate Solution (compound Sod. Lactate Injection) (self-collapsible bag without requiring any airway)500ml	As & when required
186	Two Port Close system contain Mannitol 20% 100ml	As & when required
187	Plastic Dextrose Saline N/3 (Dex. + Sod Chloride)(FFS) 5gm + 0.33gm/100ml (500ml)	60000
188	Plastic Glycine Irrigation Solution (FFS) 1.5% 3000ml	400
189	Plastic Mannitol (FFS) 20% 100ml	As & when required
190	Plastic Normal saline (Sod. Chloride) (FFs) 0.9% 3Ltr	18000
191	Glass Dextrose 5% 500ml	50000
192	Glass Dextrose 50% 100ml	40000
193	Glass Dextrose Saline N/2 (Dex+ Sod. Chloride) 5gm + 0.45gm/100ml (500ml)	10000
194	Glass Dextrose Saline N/3 (Dex. + Sod Chloride) 5gm+ 0.33gm/100ml (500ml)	8000
195	Glass Dextrose saline N/5 (Dex + Sod. Chloride) 5gm + 0.20gm/100ml (500ml)	5000
196	Glass DNS (Dex + Sod. Chloride) 5gm + 0.9gm/100ml (500ml)	55000
197	Glass Mannitol 20% 100ml	20000
198	Glass Mannitol 20% 350ml	3500
199	Glass Glass Normal Saline (Hypertonic) 3% 100ml	40000
200	Glass Normal Saline (Sod. Chloride) 0.9% 100ml	400000
201	Glass Normal Saline (Sod. Chloride) 0.9% 500ml	350000
202	Glass Ringer Lactate Solution (Compound Sod. Lactate Injection) 500ml	55000
203	Plastic Distilled Water 1Ltr	As & when required
204	Plastic Distilled Water 1Ltr	70000
205	Plastic Dextrose (FFS) 10% 500ml	8000
206	Plastic Dextrose (FFS) 5% 500ml	As & when required
207	Plastic Dextrose Saline N/2 (Dex + Sod. Chloride)(FFS) 5gm + 0.45gm/ 100ml (500ml)	As & when required
208	Plastic DNS (Dex + Sod. Chloride)(FFS) 5gm + 0.9gm/100ml(500ml)	60000
209	Two Port Close system containe Normal Saline (Sod. Chloride) 0.45% 500ml	As & when required
210	Two Port Close system container Normal Saline (Sod Chloride) .45% 100ml	As & when required
211	Tablet Ramipril+ hydrochlorthiazide 2.5mg+12.5mg	1575
212	PD Fluid 2.5% Dextrose 2 Ltr	2500
213	PD Fluid 1.5% Dextrose 2 Ltr	2500
214	PD Fluid 7.5% Dextrose 2 Ltr	2500
215	Tablet Nebivolol 5mg	6000
216	Tablet Atorvastatin 40mg	84500

217	Tablet Clopidrogrel 75mg	As & when required
218	Tablet Olmesartan + hydrochlorthiazide 20+12.5	6880
219	Tablet Olmesartan + hydrochlorthiazide 40mg+12.5mg	6000
220	Tablet Olmesartan 20mg	25000
221	Tablet Torsemide 20mg	27000
222	Injection Amiodarone HCL 150 3ml	7000
223	Tablet Amlodipine 5mg	792468
224	Tablet Metoprolol 50mg	14000
225	Tablet Metoprolol 25mg	12050
226	Tablet Metoprolol extended release 50mg	161001
227	Tablet Metoprolol extended release 25mg	158434
228	Tablet Trimetazidine M.R 35	As & when required
229	Tablet Finerenone 10mg	20
230	Injection Alteplase Tissue Plasminogen 50mg	As & when required
231	Injection Alteplase Tissue Plasminogen 20mg	5
232	Injection Tenecteplase 20mg	15
233	Injection Dalteparin 5000 I.U 0.2ml	900
234	Tablet Propranolol 10mg	30000
235	Tablet Propranolol 40mg	23494
236	Tablet Chlorthalidone 12.5mg	23000
237	Tablet Metoprolol + Amlodipine 50mg + 5mg	17000
238	Tablet Clopidrogrel + Asprin ASA 75mg + 75mg	81500
239	Tablet Isosorbide Dinitrite + Hydralazine 20mg + 37.5mg	5000
240	Tablet Ramipril 2.5mg	45000
241	Tablet Ramipril 5mg	50000
242	Tablet Rosuvastatin 5mg	60000
243	Tablet Telmidartan 80mg	20000
244	Tablet Telmidartan + Hydrochlorthiazide 80mg + 12.5mg	191557
245	Tablet Atorvastatin 5mg	17120
246	Tablet Losartan + Hyclochlorthiazide 50mg + 12.5mg	35000
247	Tablet Amlodipine 2.5mg	67115
248	Tablet Cilnidipine 10mg	80000
249	Tablet Bisoprolol 2.5mg	11784
250	Tablet Amiodarone 100mg	17929
251	Tablet Atorvastatin 80mg	9370
252	Tablet Clinidipine 10mg	78639
253	Tablet Isosorbide Mononitrate SR 30mg	25000
254	Tablet Ranolazine 500mg	50000
255	Tablet Risperidone 2mg	10000
256	Tablet Trimetazidine 20mg	As & when required
257	Tablet Clinipine 5mg	22000
258	Injection Adenosine 6mg 2ml	2500
259	Injection Clonidine 150mcg 1ml	1500
260	Injection Dopamine 40mg 5ml	As & when required
261	Injection Esmolol 100mg 10ml	3000

262	Injection Terlipressin 1mg 10ml	4500
263	Injection Dobutamine HCL 250mg/20ml	6500
264	Injection Tenecteplase 30mg	6
265	Injection Tenecteplase 40mg	As & when required
266	Tablet Eplerenone 25mg	500
267	Tablet Spironolactone + Fresemide 50mg + 20mg	30000
268	Tablet Perindopril 2mg	600
269	Tablet Atenolol 50mg	9632
270	Tablet Atenolol 25mg	13424
271	Tablet Atorvastatin 10mg	294490
272	Tablet Atorvastatin 20mg	138398
273	Tablet Carvedilol 6.25mg	14536
274	Tablet Diltiazem 30mg	3000
275	Tablet Indapamide SR 1.5mg	12000
276	Tablet Losartan Potassium 25mg	60000
277	Tablet Losartan Potassium 50mg	45000
278	Tablet Telmisartan 20mg	320000
279	Tablet Telmisartan + Hydrochlorthiazide 40mg + 12.5mg	200000
280	Tablet Telmisartan 40mg	13000
281	Tablet Nicorandil 5mg	25000
282	Tablet Torsemide 10mg	50000
283	Tablet Nifadepin SR 10mg	2500
284	Injection Labetalol 20mg 4ml	20000
285	Tablet Nitroglycerine 2.6mg	21000
286	Tablet Aspirin ASA 150mg	As & when required
287	Tablet Aspirin ASA 75mg	As & when required
288	Tablet Nimodipine 30mg	5000
289	Tablet Clobazam 5mg	11320
290	Tablet Clobazam 10mg	27895
291	Tablet Clonazepam 0.25mg	59330
292	Tablet Alprazolam 0.5mg	25000
293	Tablet Amitriptyline 25mg	18150
294	Tablet Amitriptyline 10mg	49230
295	Tablet Haloperidol 1.5mg	As & when required
296	Tablet Haloperidol 5mg	300
297	Tablet Lorazepam 1mg	20000
298	Tablet Atomoxatine 25mg	As & when required
299	Tablet Modafinil 100mg	As & when required
300	Tablet Quetiapine 25mg	16000
301	Tablet Quetiapine 50mg	As & when required
302	Tablet Zolpidem 5mg	10000
303	Capsule Flupirtine Maleate 100mg	As & when required
304	Bottle Levetiracetam 100mg/ml(100ml)	As & when required
305	Tablet Betahistine 8mg	28160
306	Capsule D- Penicillamine 250mg	As & when required

307	Tablet Pregabalin + Methylcobalamin 75mg + 750mg	102000
308	Tablet Clonazepam 0.5mg	59525
309	Tablet Piracetam 400mg	As & when required
310	Tablet Sertraline HCL 50mg	20000
311	Tablet Sodium Valproate + Valproic Acid CR 300mg	10000
312	Tablet Sodium Valproate + Valproic Acid CR 500mg	15000
313	Tablet Trihexyphenidyl Benhexol 2mg	As & when required
314	Injection Fosphenytoin 150mg	30
315	Injection Sodium Valproate 100mg/ ml (5ml)	3300
316	Injection Tramadol 100mg 2ml	35000
317	Injection Midazolam 5mg/5ml vial	100000
318	Tablet Clozapine 50mg	5000
319	Tablet Imipramine 25mg	As & when required
320	Tablet Olanzapine 5mg	As & when required
321	Tablet Pregabalin 75mg	As & when required
322	Tablet Venlafaxine 37.5mg	As & when required
323	Tablet Topiramate 25mg	As & when required
324	Tablet Carbamazepine 200mg	11920
325	Tablet Tramadol 50mg	35000
326	Tablet Phenytoin Sodium 100mg	20000
327	Tablet Montelukast 10mg	20000
328	Tablet Montelukast + Levocetirizine 10mg + 5mg	400000
329	Inhaler Salmetrol + Fluticasone 50mcg + 250mcg	62
330	Rotacap Formetrol + Fluticasone 6mcg+ 250mcg	As & when required
331	Inhaler Capsule Indacaterol / Glycopyronium 110/50mcg	390
332	Inhaler Formoterol + Budesonide 6mcg + 200mcg	500
333	Rotacaps Salmetrol + Fluticasonerotacaps 250mcg + 50mcg	As & when required
334	Tablet Pirfenidone 200mg	As & when required
335	Tablet Ebastine 10mg	As & when required
336	Tablet Vericiguat 2.5mg	50
337	Inhaler Budesonide 100mcg	1000
338	Inhaler Formeterol + Budesonide 6mcg + 100mcg	500
339	Inhaler Formeterol + Budesonide 6mcg + 400mcg	As & when required
340	Inhaler Formetrol + Fluticasone Inhaler 250mcg	200
341	Inhaler Budesonide 200mcg	1200
342	Inhaler Fluticasone + Salmetrol 250mcg + 25mcg	As & when required
343	Inhaler Fluticasone + Salmetrol 125mcg + 25mcg	As & when required
344	Inhaler Ipratropium 20mcg	As & when required
345	Inhaler Ipratropium Br + Levosalbutamol 20mcg + 50mcg	As & when required
346	Inhaler Salbutamol 100mcg	As & when required
347	Inhaler Tiotropium 9 mcg	As & when required
348	Injection N-Acetylcysteine 20% 200mg/ml (5ml)	As & when required
349	Injection N-Acetylcysteine 1gm / 5ml	30000
350	Tablet Acetylcysteine 600mg	As & when required
351	Injection Ramucirumab 100mg/10ml	As & when required

352	Injection Isoprenaline 2mg 1ml	800
353	Tablet Metformin SR 100mg	As & when required
354	Injection Human Biphasic Isophane Insulin 30/70 Pen (Permanent) & Penfill. Note Lowest firm shall supply pen, free of cost as per requirement. 100IU/ml(3ml)	As & when required
355	Tablet Teneligliptine 20mg	310865
356	Tablet Vildagliptine 50mg	1200
357	Tablet Vildagliptine + Metformin 50mg+500Mg	900
358	Tablet Metformin SR 500mg	35000
359	Tablet Glimipride 1mg	350000
360	Tablet Glimipride+ Metformin 1mg+500mg	65000
361	Tablet Glimipride+ Metformin 2mg+500mg	62000
362	Tablet Metformin 500mg	330000
363	Tablet Acarbose 50mg	800
364	Tablet Glimipride 2mg	1267055
365	Injection Human Biphasic Isophane Insulin 30/70 vial 10ml	5600
366	Injection Human Neutral soluble insulin vial 40IU ()10ml)	As & when required
367	Tablet Sitagliptin 100mg	95000
368	Tablet Sitagliptin 50mg	20440
369	Tablet Sitagliptin + Metformin 50mg + 1000mg	150000
370	Tablet Sitagliptin + Metformin 50mg + 500mg	130000
371	Tablet Dapagliflozin 10mg	205000
372	Tablet Dapagliflozin + Metformin 10mg + 500mg	500
373	Tablet Dapagliflozin + Metformin 10mg + 100mg	As & when required
374	Tablet Gliclazide 80mg	25000
375	Tablet Gliclazide moderate release 30mg	30000
376	Tablet Gliclazide MR 30mg	As & when required
377	Tablet Glibenclamide 5mg	600
378	Tablet Pioglitazone 15mg	16000
379	Injection Human Biphasic Isophane Insulin 30/70 pen (Disposable) & Penfill 300IU(3ml)	44423
380	Injection Human Isophane Insulin 40IU(10ml)	60472
381	Injection Human Neutral soluble insulin pen fill 100IU/ml (3ml)	30677
382	Injection Insulin Glargin Pen 300IU (3ml)	18406
383	Eye Drop Polyethylene glycol + Propylene Glycol 0.4%w/V+0.3%w/v(10ml)	55029
384	Eye Drop Cyclosporin emulsion with glycerin castor oil polysorbate 80 carbomer 1342 preservative free 0.05% 0.4ml	As & when required
385	Eye Drop bimatoprost with BAK 0.01% w/v + 0.02% 3ml	136
386	Intra- Vitreal Implant Dexamethasone Intravitreal Implant 0.7mg	4
387	Eye Drop Dorolamide 2.0% (5ml)	As & when required
388	Eye Drop Dorzolamide Hydrocoride + Timolol Maleate Combination 2% + 0.5% (5ml)	467
389	Eye Drop Flurbiprofen 0.03%	76
390	Eye Drop Travoprost 0.004%	1275
391	Tablet Acetazolamide 250mg	9700
392	Eye Drop Carboxymethyl Cellulose 0.5% 10ml	201698
393	Eye Drop Nepafenac 0.1% w/v 5ml	8158
394	Eye Drop Olapatadine 0.1% 5ml	As & when required

395	Eye Drop Timolol Eye drop 0.5% 10ml	443
396	Eye Drops Moxifloxacin 0.5% 5ml	25000
397	Injection Ranibizumab 10mg/ml	4
398	Solution for intravitreal Injection Pre-filled Syringe Brolucizumab 6mg/0.05ml(120mg/ml)	As & when required
399	Injection Iron Sucrose 100mg 5ml	110
400	Bottle Oral iron preparations containing Ferrous sulphate/ Ferrous fumarate/Ferrous gluconate/ Ferrous ammonium citrate 200ml	6000
401	Injection Biological Tissue Glue 2ml	244
402	Injection Biological Tissue Glue 4ml	As & when required
403	Injection Hemostatic Matrix - in prefilled syringe with calcium chloride & thrombin in vials (Hemostatic matrix)5ml	65
404	Injection Concizumab 150mg/1.5ml	10
405	Tablet Sevelamer Carbonate 800mg	20000
406	Injection Total parenteral nutrition solution in three chambered beg for central infusion containing glucose lipids and amino acids. Volume 1850- 2100ml, total calories at least 1800 Kcal, osmolality 1000-1500 mOsm/Kg. 1850-2100ml.	As & when required
407	Injection Total parenteral nutrition solution in three chambered beg for peripheral infusion containing glucose, lipids and amino acids, Volume 1000-1250ml, total calories at least 750Kcal, Osmolality/ Osmolarity 600- 1000mOsm/Kg 1000-1250ml	As & when required
408	Injection Dipeptide of N2 Alanyl glutamin 20gm 50ml	As & when required
409	Injection Intralipid Phospholipids & fat emulsion solution 100ml 20%	As & when required
410	Combination of Insulin Glargine 100 IU/ML + Lixisenatide 50mcg/ml	15
411	Injection Intralipid Amino acid solution 500ml 10%	As & when required
412	Injection Omega-3 fatty acid 50ml	469
413	Injection Erythropoietin Beta 5000 I.U Pfs	272
414	Injection Erythropoietin Alpha 4000IU 1ml	8132
415	Injection Guselkumab 100mg	10
416	Tablet Sevelamer Hydrochloride 400mg	21000
417	Capsule Mecobalamin + Calcitriol + Calcium Carbonate + Ecosapentaenoic Acid + Docosahexaenoic Acid + Folic Acid + Boron	As & when required
418	Drops Vitamin D3 400 IU/ml	5000
419	Injection Potassium Chloride 10ml	20000
420	Injection Recombinant Factor VIII 250IU	13040
421	Injection Thiamine 100mg 2ml	20000
422	Bottle Multivitamins & Minerals Combination Oral preparations containing Vit. B Complex C with or without Vit A,D Minerals & trace elements 200ml.	115061
423	Tablet Entecavir 0.5mg	21967
424	Tablet Declatasvir 60mg	As & when required
425	Tablet Ledipasvir + Sofosbuvir 90mg + 400mg	As & when required
426	Tablet Tenoforvir 300mg	4000
427	Tablet Tenofovir Alafenamide 25mg	2000
428	Tablet Sofosbuvir 400mg	500
429	Tablet Sofosbuvir + Velpatasvir 400mg + 100mg	1000
430	Injection Acyclovir 250mg 10ml	As & when required
431	Syrup Methadone 5mg/ml	As & when required

432	Tablet Entecavir 1mg	200
433	Tube Mupirocin 2% (5mg to 7.5gm)	60000
434	Cream Luliconazole 30gm	1000
435	Capsule Acitretin 10mg	As & when required
436	Capsule Acitretin 25mg	250
437	Ointment Betamethasone valerate 0.12% (20gm)	50
438	Ointment Betamethasone valerate+ neomycin 0.12% + 0.5% (20gm)	1506
439	Tube Adaplene gel 0.1% (15gm)	As & when required
440	Ointment Flucinolone 0.1% (20gm)	As & when required
441	Ointment Flucinolone 0.025% (15gm)	As & when required
442	Ointment Silver Sulphadiazine 1% w/w (25gm)	2
443	Ointment Silver Sulphadiazine 1% w/w (250gm)	6025
444	Tube Clotrimazole 1% w/w (15gm)	10000
445	Ointment Sodium fusidate 2% 10gm	100
446	Lotion Ketoconazole Lotion 2% (60ml)	As & when required
447	Solution Minoxidil Solution 60ml 2%	50
448	Solution Minoxidil Solution 60ml 5%	2500
449	Ointment Polymyxin B Sulph. Equivalent to neomycin 5000IU + 3400IU per gm (20gm-30gm)	As & when required
450	Ointment Silver Sulphadiazine + Chlorhexidine 1% + 0.2% (25gm)	1500
451	Tube Povidone Iodine 5% w/w (15gm)	15000
452	Ointment Tacrolimus 0.1%	As & when required
453	Ointment Tacrolimus 0.03%	20
454	Injection Ondansetron 4mg 2ml	400000
455	Tablet Rabeprazole Sodium 20mg	30000
456	Tablet Rabeprazole + Domperidone SR 20mg + 30mg	200
457	Tablet Pantoprazole + Domperidome SR 40mg + 30mg	708402
458	Tablet Pantoprazole 40mg	1449651
459	Tablet Ursodeoxycholic Acid 300mg	30000
460	Tablet Metronidazole 400mg	80000
461	Capsule Lansoprazole 15mg	80020
462	Tablet Lansaprazole 30mg	15000
463	Tablet Mozapride 5mg	200
464	Tablet Sulfasalazine 500mg	94420
465	Tablet Metoclopramide 10mg	8500
466	Tablet Ondansetron 4mg	160000
467	Tablet Trypsin + Bromelain + Rutoside 48mg + 90mg + 100mg	31000
468	Injection Ondansetron 8mg 4ml	45000
469	(Colonic Preparation) Polyethylene Glycol 3350	As & when required
470	Capsule Billion cfu of L.rhamnosus GR-1 and L. reuteri- probiotic capsule	As & when required
471	Sachet Polyethylene glycol, activated dimethicone with electrolytes for oral solution 17.1gm	1100
472	Tablet Ursodeoxycholic Acid 150mg	2700
473	Injection Pantoprazole 40mg	230000
474	Bottle Lactulose sugar free 10gm/15ml 100ml	35000

475	Capsule Omeprazole 20mg	324163
476	Tablet Ranitidine 150mg	210000
477	Tablet Ondansetron 8mg	10000
478	Tablet Domperidone 10mg	36000
479	Bottle Ondansetron 2mg/ 5ml	2000
480	Tablet Drotaverine 40mg	10000
481	Tablet Drotaverine 80mg	6500
482	Tablet Esmoprazole 40mg	25000
483	Tablet Tinidazole 500mg	1000
484	Injection Tranexamic Acid 500mg 5ml amp	551524
485	Tablet Rivaroxaban 2.5mg	As & when required
486	Tablet Rivaroxaban 10mg	300
487	Tablet Rivaroxaban 15mg	1000
488	Tablet Rivaroxaban 20mg	As & when required
489	Tablet Deferasirox 250mg	500
490	Tablet Deferasirox 400mg	500
491	Tablet Apixaban 2.5mg	20000
492	Injection Heparin Lock Flush solution 10IU/ml vial of 10.0ml	1000
493	Injection Protamine Sulphate Note: Injection Heparin 25000 IU and Injection Protamine will be purchased from single firm who are quoting both the items and selection shall be made on overall lowest (composite) basis after adding total cost of both the items 50mg 5ml amp	1200
494	Injection Tirofiban 5mg/ 100ml	5
495	Injection (PFS) Darbepoietin Alpha 60mcg	100
496	Tablet Fenofibrate 160mg	8500
497	Injection (PFS) Pegylated Recombinant erythropoietin 100mcg	150
498	Injection Iron Isomaltoside 1000 5ml vial	10
499	Injection Emicizumab 60mg/0.4ml	250
500	Capsule Dabigatram 75mg	As & when required
501	Capsule Dabigatram 150mg	As & when required
502	Injection Menandione	As & when required
503	Injection Heparin 5000IU 5ml	50000
504	Injection Fibrinogen Concentrate 1gm	As & when required
505	Injection (PFS) Darbepoietin Alpha 40mcg	500
506	Injection Enoxaparin, PF Syringe 80mg/0.8ml	As & when required
507	Injection Enoxaparin, PF Syringe 40mg/0.4ml	25000
508	Injection Enoxaparin, PF Syringe 60mg/0.6ml	16000
509	Tablet Nicoumalone 1mg	4500
510	Tablet Nicoumalone 2mg	6500
511	Tablet Etoricoxib 90m	12500
512	Tablet Acelofenac + Paracetamol 100mg+325mg	75000
513	Tablet Etoricoxib 60mg	6200
514	Injection Paracetamol 1gm 100ml	1029459
515	Tablet Ibuprofen 400mg	15000
516	Tablet Pataacetamol + Tramadol 325mg + 37.5mg	160000

517	Suspensiun (Drops) Paracetamol 100mg	500
518	Tablet Paracetamol (Dispersible) 125mg	550
519	Tablet Paracetamol (Dispersible) 250mg	As & when required
520	Tablet Paracetamol 500mg	710000
521	Injection Paracetamol I.V. 500mg 50ml	5500
522	Tablet Paracetamol 650mg	510000
523	Tablet Naproxen + Domperidone 500mg + 10mg	45000
524	Spray Diclofenac+ Linseed + Menthol + Methyl Salicylate 1.16% W/V + 3%W/V + 5% W/V + 10% W/V	4999
525	Injection Diclofenac 25mg/3ml Vial	2500
526	Injection Paracetamol 300mg 2ml	As & when required
527	Injection Diclofenac (without Cremophor & Transcutol - P) 75mg/1ml	492026
528	Injection Rituximab 100mg 10m	150
529	Injection Methyl prednisolone Sodium succinate 1gm	4500
530	Injection Ranibizumab 10mg	5000
531	Tablet Fluconazole 150 mg	5300
532	Injection Rituximab 500mg 50ml	2000
533	Injection Daratumumab and hyaluronidase fihj 1800mg	10
534	Tablet Mycophenolate mofetil 500mg	25000
535	Tablet Mycophenolate mofetil 250mg	2000
536	Tablet Mycophenolate sodium 360mg	1000
537	Tablet Mycophenolate sodium 180mg	100
538	Tablet Tacrolimus 0.5mg	1000
539	Tablet Tacrolimus 1mg	1200
540	Injection Liposomol Amphotericin- B (dextrose) 50mg/50ml	25000
541	Injection Anidulafungin 100mg	1500
542	Injection Micafungin Sod. 50mg	150
543	Tablet Mycophenolate Mofetil 750mg	80
544	Tablet Tacrolimus 2mg	900
545	Injection Methoxy Polyethyleneglycol Epoietin Beta 50 mcg 0.3ml	40
546	Injection Methoxy Polyethyleneglycol Epoietin Beta 100 mcg 0.3ml	30
547	Tablet Tamsulosin 0.2mg	3000
548	Tablet Tolteridene 4mg	20
549	Injection Infliximab 100mg	2508
550	Tablet Sirolimus 1mg	As & when required
551	Powder Clotrimazole Powder 1% (30MG)	12000
552	Injection Basiliximab 20mg	20
553	Capsule Isavuconazole 100mg	700
554	Injection Isavuconazole 200mg	1000
555	Injection Caspofungin 50mg	11000
556	Injection Caspofungin 70mg	1500
557	Injection Hydrocortisone Succinate 100mg / vial	60000
558	Injection Methyl prednisolone Acetate 40mg 1ml	7421
559	Injection Omalizumab 150mg	100
560	Capsule Tamsulosin 0.4mg	21500

561	Tablet Tamsulosin + Dutasteride 0.4+0.5mg	30000
562	Injection Pralidoxime (PAM Chloride) 1gm	100
563	Injection Naloxone 400mcg	1500
564	Injection Noradrenaline 2mg/ml	215000
565	Syrup Elemental Iron, Folic Acid 200ml	3200
566	Injection Sugammadex	500
567	Spray Fluticasone Nasal Spray 100mcg	100
568	Tablet Alfuzosin 10mg	6000
569	Injection Albumin 20% Bottle 100ml	32000
570	PFS Filgrastim 150mcg	600
571	Oral Tablet Tofacitinib 5mg	2000
572	Tablet Methylcobalmine + Gabapentin 500mg+300mg	As & when required
573	Mouth Wash Chlorhexidine 0.2% 150-160ml	12000
574	Liquid Clotrimazole Mouth Paint 1% 15ml	1000
575	Mouth Wash Povidone-Iodine Mouthwash/Gargle 2% 100ml	12000
576	Solution Cetrimide+ Chlorhexidine 15% +7.5% (500ml)	As & when required
577	Solution Cetrimide+ Chlorhexidine 15% +7.5% (1000ml)	As & when required
578	Insulin Glargine (rDNA origin) 1.5ml Prefilled Pen 300IU	20
579	Injection Human Chorionic Gonadotrophin Recombinant HCG 2000 IU/ml	99
580	Injection Caffeine Citrate 20mg 1ml	800
581	Injection Ephedrine 30mg 1ml	As & when required
582	Suppository Rectal Diclofenac 100mg	As & when required
583	Injection Neostigmine + Glycopyrrolate 2.5mg +0.5mg	2100
584	Injection Piracetam 200mg	As & when required
585	Suppository Paracetamol Suppository 250mg	35
586	Injection Buprenorphine 0.3mg 1ml	30
587	Injection Tocilizumab 400mg	100
588	Injection Etenarcept 25mg 0.5ml	150
589	Injection Etenarcept 50mg/ 1ml	55
590	Tablet Megestrol Acetate 40mg	70
591	Tablet Megestrol Acetate 160mg	600
592	Injection Alprostadil 500mcg 1ml	150
593	Injection Leucovarin Calcium Folinate 50mg	500
594	Tablet Naltrexone 50mg	300
595	Tablet Buprenorphine 0.4mg	8000
596	Tablet Buprenorphine + Naloxone 2mg + 0.5mg	10
597	Tablet Zinc Acetate 50mg	10000
598	Injection Cetrorelix Acetate GnRH Antagonist 0.25mg	3000
599	Injection Cefepime plus enmetazobactam 2.5gm	10
600	Tablet Empagliflozin 25mg	42000
601	Injection Teriparatide (PFS) 600mcg/ 2.4ml	500
602	Injection ACTH 1ml/ 60IU	As & when required
603	Tablet Calcium Citrate Maleate 250mg + folic Acid 250mg + 50mcg	9787426
604	Tablet Thyroxine 100mcg	23500
605	Tablet Thyroxine 25mcg	180000

606	Tablet Thyroxine 50mcg	170000
607	Tablet Artesunate + Sulphadoxine + Pyrimethamine 200mg + 500mg + 25mg	200
608	Tablet Atremether + Lumefantrine 80mg + 480mg	80
609	Tablet Solifenacin 5mg	200
610	Tablet Levocetirizine 5mg	550000
611	Tablet Estradiol Valerate 2mg	As & when required
612	Injection Teriparatide (Cartridge) 250mcg/ml (3ml)	As & when required
613	Tablet Baclofen	16000
614	Injection Urinary Gonadotropin (Urofollitropin) 75IU	As & when required
615	Injection Urinary Gonadotropin Menotrophin 75IU	As & when required
616	Tablet Cinacalcet 30mg	1000
617	Injection Human Chorionic Gonadotrophin HCG Urinary Highly purified 5000IU	981375
618	Injection Recombinant FSH Follitropin a b M.D 1200IU - 1500 Iu (1.75ml)	As & when required
619	Syrup Hydroxyzine 10mg/5ml	183
620	Tablet Alfacalcidol 0.25mcg	18000
621	Tablet Misoprostol (Prostaglandine E1) 200mcg	3500
622	Tablet Norethisterone 5mg	As & when required
623	Injection Recombinant FSH Follitropin a B M.D 900IU - 1100IU (1.75ml) Multi dose	As & when required
624	Injection Recombinant LH 75IU	As & when required
625	Injection Human Chorionic Gonadotropin HCG (Recombinant) 250IU	As & when required
626	Tablet Solifenacin 10mg	As & when required
627	Tablet Mirabegron 25mg	7000
628	Tablet Mirabegron 50mg	As & when required
629	Tablet Fenofibrate 40mg	As & when required
630	Injection Phenylephrine 10mg 1ml	13645
631	Tablet Calcium Acetate 667mg	15000
632	Tablet Baclofen 10mg	16000
633	Tablet Progesterone 200mg	6600
634	Tablet Hydroxychloroquine 200mg	421189
635	Syrup Levetiracetam 100 mg/mL	650
636	Syrup Brivaracetam 10mg/ml	As & when required
637	Drops Cholecalciferol 1ml/800U	2000
638	Capsule Vitamin D3 60000IU	As & when required
639	Tablet Methyl cobalamine +Alpha Lipoic Acid+ Folic Acid+ pyridoxine 750mcg+ 100mg+15mg +3mg	386072
640	Powder Cholecalciferol Sachet Vit. D3 60000 units 1gm	40000
641	Injection Vit.B1, B6, B12 combination 2 ml	As & when required
642	Capsule Elemental calcium+ calcitriol++zinc sulphate 500mg+0.25mcg+7.5mg	As & when required
643	Capsule All-trans retinoic acid 10 mg	As & when required
644	Tablet Calcium carbonate 625 mg	As & when required
645	Tablet Calcium folinate 15 mg	As & when required
646	Solid oral dosage form ergocalciferol 1.25 mg (50 000 IU)	As & when required
647	Tablet Ferrous salts(a) Iron Dextran;(b) Iron sorbitol citrate complex equivalent to 60 mg of elemental iron	As & when required
648	Tablet Pyridoxine 40 mg	9800

649	Tablet Pyridoxine 50 mg	As & when required
650	Tablet Riboflavin 10 mg	As & when required
651	Tablet Thiamine 100 mg	As & when required
652	Capsule/Tablet (including Chewable Tablet) Vitamin A 50000 IU	As & when required
653	Dispersible Tablet Zinc Sulphate 20 mg	As & when required
654	Tablet Ferrous Ascorbate 300 MG (elemental iron 60 mg)	As & when required
655	Solid oral dosage form Cholecalciferol 1000 IU	As & when required
656	Oral oily solution Retinol 100 000 IU/mL (as palmitate) in multidose dispenser.	As & when required
657	Syrup Vitamin A 100000 IU/ml	As & when required
658	Sachet Vitamin A+ Carbohydrate+ Energy+ Protein+ Vegetable fat+ Vit C+ Vit D	As & when required
659	Tablet Calcium Carbonate , L-Arginine , L- Lysine , Vitamin B6, Vitamin C , Vitamin D3 , and Zinc sulphate 250 mg+500 mg+200 mg 1.5mg+75 mg+200 IU+61.8 mg.	As & when required
660	Tablet Glucosamine Sulphate+ Chondroitin Sulphate 250 mg + 200 mg	As & when required
661	Sachet L-Arginine 5gm	As & when required
662	Capsule Oral preparation containing Ferrous sulphate with Vit. B Complex sustained release 150 mg	As & when required
663	Oral L- Glutamine 10gm sachet	As & when required
664	Sachet Amino Acids, Vitamins, Minerals and Lycopene Powder	As & when required
665	Eye Drop Atropine 1% 5ml	30
666	Eye Drop Hydroxypropyl Methylcellulose 0.3% 10 ml	As & when required
667	Eye Drop Ofloxacin 0.3%, 5ml	As & when required
668	Eye Drop Ofloxacin +Dexamethasone 10ml (0.3% + 0.1%)	As & when required
669	Eye Drop Polyvinyl Alcohol + Povidone + Chlorbutol eye drops 1.4%+0.06%+0.5% (10ml)	As & when required
670	Eye Drop Povidone Iodine 5% -pack of 5 ml	As & when required
671	Eye Drop Prednisolone 1% 10ml	As & when required
672	Eye Drop Proparacaine HCL 0.5% 5ml	As & when required
673	Eye Drop Tobramycin 0.3% - pack of 5ml	27
674	Eye Drop Tropicamide eye drops 1% 5ml	1561
675	Ophthalmic solution-based eye drop Ketorolac tromethamine 0.4% 5ml	As & when required
676	Injection Tropicamide-Phenylephrine-Lidocaine 0.2mg/ml+3.1mg/ml+10mg/ml	As & when required
677	Injection Hydroxy Propyl Methyl Cellulose with PFS 2% w/v 2ml	As & when required
678	Injection Hydroxy Propyl Methyl Cellulose 2% 2ml	As & when required
679	Injection Trypan blue Dye	As & when required
680	E/d Fluromethalone 0.001	65
681	Tablet Isoniazid 300mg	7000
682	Tablet Bilastine 20mg	As & when required
683	Tablet Ethambutol 400 mg	As & when required
684	Tablet Ethambutol 800 mg	27687
685	Tablet Ethionamide 250 mg	As & when required
686	Tablet ethambutol + isoniazid + pyrazinamide + rifampicin 275 mg + 75 mg + 400 mg + 150 mg	As & when required
687	Tablet ethambutol + isoniazid + rifampicin 275 mg + 75 mg + 150 mg	As & when required
688	Tablet (dispersible) isoniazid + pyrazinamide + rifampicin : 50 mg + 150 mg + 75 mg	As & when required
689	Tablet isoniazid + rifampicin 50 mg + 75 mg	As & when required
690	Tablet isoniazid 100 mg	200

691	Capsule Rifampicin 150 mg	100
692	Tablet Promethazine 25mg	As & when required
693	Capsule Isoniazid+ Rifampicin 300mg+450mg	3062
694	Capsule Rifampicin 450mg	11103
695	Injection Sodium hyaluronate10mg1ml	20
696	Tablet Posaconazole 100mg	2200
697	Capsule Atra Tretinoin 10mg	3200
698	Injection Posaconazole 18mg	1608
699	Injection Treosulfan 5gm	200
700	Combi pack Artesunate 200 mg + Sulphadoxine -Pyrimethamine (750 mg + 37.5 mg)	720
701	Combi pack Artesunate 100 mg + Sulphadoxine -Pyrimethamine (750 mg + 37.5 mg)	As & when required
702	Injection Artesunate 60mg	3346
703	Tablet Combo Pack (Mifepristone + Misoprostol) 200 mg+200mcg	As & when required
704	Injection Oxytocin 0.5mg	As & when required
705	Tablet Glipizide+Metformin5mg+500mg	As & when required
706	Tablet Mifepristone 200mg	As & when required
707	Tablet Combo Pack (Mifepristone + Misoprostol) 200mg+400mcg	As & when required
708	Tablet Glibenclamide + Metformine 2.5mg + 500mg	As & when required
709	Tablet Progesterone 100mg	As & when required
710	Capsule Progesterone 100mg	4000
711	Tube Feracrylum gel 3% 15gm	1000
712	Injection Iohexol 240mg/ml (50ml)	As & when required
713	Injection Iopamidol 300mg/ml (75ml)	100
714	Plastic Haemodialysis Fluid with Bicarbonate powder BP (FFS) 5 Ltr	As & when required
715	Plastic Haemodialysis Fluid with Bicarbonate powder BP (Low calcium) (FFS) 5 Ltr	As & when required
716	Plastic Haemodialysis Fluid with high concentration of sodium (103mgdl) 5 Ltr	As & when required
717	Plastic Dextrose Saline N/3 (Dex.+Sod. Chloride)(FFS) 5gm+0.33gm/100ml (500 ml)	As & when required
718	Glass Dextrose 50% 100 ml	As & when required
719	Glass Dextrose Saline N/3 (Dex.+Sod. Chloride) 5gm+0.33gm/100ml (500 ml)	As & when required
720	Glass Dextrose Saline N/5 (Dex.+Sod. Chloride) 5gm+0.20gm/100ml (500 ml)	As & when required
721	Glass Distilled water 1 Ltr	As & when required
722	Injection Glucose 50%	As & when required
723	Glass Mannitol 20% 100ml	As & when required
724	Glass Mannitol 20% 350ml	As & when required
725	Glass Dextrose 10% 500ml	As & when required