अखिल भारतीय आयुर्विज्ञान संस्थान, नागपुर

ALL INDIA INSTITUTE OF MEDICAL SCIENCES, NAGPUR

Address: Plot No. 2, Sector-20, MIHAN, Nagpur– 441108

Email: - hospitalstore@aiimsnagpur.edu.in Ph.No.0710-3295590

Notice Inviting Tender

For

" Annual Rate Contract for Supply of Drugs"

At

All India Institute of Medical Sciences, Nagpur

CRITICAL DATE SHEET

Published Date	13/06/2024 at 05:00 PM
Bid Document Download Start Date	13/06/2024 at 05:05 PM
Bid Submission Start Date	14/06/2024 at 09:00 AM
Bid Submission End Date	11/07/2024 at 03:00 PM
Bid Opening Date	12/07/2024 at 03:00 PM

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Signature and Stamp of Bidder

Introduction

- Online bids are invited on single stage two bid systems for "Annual Rate Contract for Supply of Drugs". **Manual bids shall not be accepted**.
- Tender document may be downloaded from AIIMS web site www.aiimsnagpur.edu.in (for reference only) and CPPP site https://eprocure.gov.in/eprocure/app as per the schedule as given in CRITICAL DATE SHEET asunder.
- Bid shall be submitted online at CPPP website: https://eprocure.gov.in/eprocure/app
- Bid documents may be scanned with 100 dpi with black and white option which helps in reducing size of the scanned document.
- Tenderer who has downloaded the tender from the AIIMS web site **www.aiimsnagpur.edu.in** and Central Public Procurement Portal (CPPP) e-procurement website https://eprocure.gov.in/eprocure/app shall not tamper/modify the tender form including downloaded price bid template in any manner. In case if the same is found to be tampered/modified in any manner, tender shall be completely rejected and tenderer is liable to be banned from doing business with AIIMS Nagpur.
- The Technical bid should include the detailed specifications of item. All items should be numbered as indicated in the Annexure-I (Any deviation should be clearly mentioned and supporting document should be submitted).

1. TENDERING SYSTEM:

The Bids are to be submitted in two Parts i.e.

i. Technical Bid **(Cover "A")**
ii. Financial Bid / Price Bid **(Cover "B")**

i. The **TECHNICAL BID** shall contain the complete technical details of the firm and the documents to provide the eligibility and competency of the bidder and shall be submitted online only in the manner prescribed in Bid document.
Bid Security shall be submitted before the specified schedule at the office of AIIMS, Nagpur super scribed, **"Tender Documents for Tender Reference No. AIIMS-NAG/Drugs/RC/OTE/HOS/24-25/04 dated 13/06/2024 for Annual Rate Contract for Supply of Drugs"**

ii. **The Bid shall be valid for a period of 180 days from the date of opening of Bid. Prior to the expiry of the bid validity**, the Tender Inviting Authority may request the Tenderers to extend the bid validity for further period as deemed fit on their original quoted prices and all terms & conditions. However, AIIMS, Nagpur reserves the right to place purchase orders at the quoted rate till such period or till completion of Rate contract.

a) The Tenderer shall fill in the rate per unit size, rate of GST in respective column of BOQ for the items quoted**.**
b) In determining the lowest evaluated price, the rate quoted per unit size exclusive of GST as indicated in the **BOQ** shall be taken into consideration.
c) Tender has been called for the **generic drugs.**
d) Rates (inclusive of customs duty, packing & forwarding charges, transportation, insurance and any incidental charges, but exclusive **GST** should be quoted for each of the required

drugs, medicines etc., on door delivery basis according to the unit ordered. Tender for the supply of drugs, medicines, etc. with cross conditions like "AT CURRENT MARKET RATES" shall not be accepted. Handling, clearing, transport charges etc., will not be paid separately. The delivery should be made as stipulated in the purchase order placed with Tenderers.

e) The price quoted by the tenderers shall not, in any case exceed the DPCO controlled price, if any, fixed by the Central/State Government, the Maximum Retail Price (MRP) and the selling price of the tenderer.

In case any tenderer quotes higher than the DPCO controlled price, competent authority shall be informed for appropriate action.

2. LAST DATE AND TIME FOR SUBMISSION OF ONLINE TENDER:

i. (a) Online Bids [in two separate Cover {Technical bid (Cover "A") and price/financial bid (Cover "B")}] shall be submitted till 15:00 Hours Up to **11/07/2024** on CPP portal i.e., https://eprocure.gov.in.

(b) Original Earnest Money Deposit shall be submitted on or before the date of opening of bid at the below mentioned address of AIIMS, Nagpur with super scribed, **"Earnest Money Deposit (EMD) for Tender Reference No.- AIIMS-NAG/Drugs/RC/OTE/HOS/24-25/04 dated 13/06/2024 for Annual Rate Contract for Supply of Drugs"**

"To,
Store Office (Hospital Store)
IPD First Floor,
All India Institute of Medical Sciences,
Plot No. 2, Sector20, MIHAN,
Nagpur– 441108"

ii. **Late Tender: -**There is NO PROVISION of uploading late tender beyond stipulated date & time in the e-tendering system.

3. ELIGIBILITY CRITERIA (TECHNICAL BID -COVER "A"):

Minimum Eligibility criteria along with list of documents to be submitted in Cover 'A'. Bidders should meet the following criteria to be eligible for bidding and relevant papers/documents must be submitted by them in their technical bid (Cover- 'A') in support of their eligibility for the tender.

I. The bidder needs to submit Earnest Money Deposit: Earnest money by means of a Bank Demand Draft/ FD, a scanned copy to be enclosed. It is also clarified that the bids submitted without earnest money will be summarily rejected. The DD/FD may be prepared in the name of "The Director, AIIMS Nagpur". The EMD cost must reach at officer of the Store Office (Hospital Store), IPD First Floor, AIIMS, Plot No. 2, Sector-20, MIHAN, Nagpur.

Note: *The Micro and Small enterprises (MSEs) are exempted from submitting the Bid Security as per prevailing rules. However, they have to submit the valid* ***"Udyam Registration Certificate"*** *in reference to* ***Gazette notification CG-DL-E-26062020-220191 dated 26.06.2020 and Office Memorandum no. 21(5)/2019-P&G/Policy (pt. IV), issued by Ministry of MSME dated 06.08.2020*** *with the technical bid.*

II. Documentary evidence of the constitution of the company/firm/Proprietorship such as Memorandum and Articles of Association, Partnership Deed etc. should be submitted with details of the Name, Address, Telephone Number, Fax Number, e-mail address of the firm and of the Managing Director / Partners / Proprietor should be submitted.

III. Power of Attorney or Resolution of the Board by which the authorized signatory has been authorized by the bidding firm to sign the documents should be submitted.

IV. mention that the bidder is Manufacture /Distributor /Dealer / Trader/Supplier relevant document should be uploaded

V. firm/company registration certificate

VI. In case of distributor/dealer/trader/supplier must upload tender specific authorization certificate from OEM/ manufacturer (**Annexure-V**)

VII. Copies of the Audited Balance Sheet, Profit and Loss statement showing details of their annual average turnover for last three consecutive financial years (2020-21, 2021-22,2022-23) not less than **₹50 Lakh** certified by charted accountant.

VIII. Copy of Income Tax Return for last three Consecutive financial years (FY 2020-21, 2021-22, 2022-23) should be submitted (self-attested).

IX. Tenderer must provide evidence of having supplied government hospital / reputed private hospital organizations in India similar nature of items of at least **₹ 50 Lakh** of Supply of Drug and Medicine of Tender value in the last three years i.e. 2020-21, 2021-22,2022-23 and the copy of the same should be uploaded

X. Relevant brochure / catalogue pertaining to the items quote with full specification etc.

XI. Technical compliance report

XII. WHO –GMP or cGMP as per schedule M (license no. and MFG. unit address highlighted)

XIII. All the documents enclosed with the bid document should also be seal signed and page numbered given to each and every page

XIV. In case bidder is Manufacturer:

a. An undertaking by the manufacturer that they have their own testing laboratories and in built quality assurance facilities and shall carry out batch-wise pre-inspection of the items (covered under the drug and cosmetic act) and submit such reports along with the supplies for each batch. The drug product should have compliances as per specification (IP/BP/USP etc.), if In-house (INH) specification than provision of reference standard and testing protocol for quoted items shall be submitted.

b. Authorization letter (with tender reference No.) nominating a responsible person (Name, Address, designation contact No. and E-mail) of the bidder to transact the business with the Tender Inviting Authority.

c. Self-attested Manufacturing/Repacking Licenses, the license must have been duly renewed up to date and the items quoted shall be clearly highlighted in the license.

d. A valid WHO-GMP or cGMP as per Schedule 'M' certificate as per requirement of the Drugs and cosmetic act and rules of Govt. of India from Competent Authority.

e. Non-conviction certificate, not more than six-month-old.

f. Instruments such as power of attorney etc.

XV. In case bidder is Distributer:

a. Details of Manufacturer. Distinct documents for each manufacturer

b. Authorization letter (with tender reference No.) given by the manufacturers to the distributer. Nominating a responsible person (Name, Address, designation contact No. and E-mail) of the bidder to transact the business with the Tender Inviting Authority. Distinct documents for each manufacturer.

c. Self-attested Valid Drug Licenses issued to distributer by local FDA. The license must have been duly renewed up to date.

d. Self-attested Manufacturing/Repacking Licenses (issued to manufacturer), the licenses must have been duly renewed up to date and the items quoted shall be clearly highlighted in the license.

e. WHO-GMP or GMP as per Schedule 'M' certificate as per requirement of the Drugs and cosmetic act and rules of Govt. of India from Competent Authority.

Signature and Stamp of Bidder

f. Non-conviction certificate (for Manufacturer), not more than six months old. Distinct documents for each manufacturer.
g. Copies of the Balance Sheet and Profit and Loss Account (for manufacturer) for the last three years. Distinct documents for each manufacturer.
h. GST No.(Distributer).
i. Declaration Form with affidavit.
j. An undertaking by distributer that it has not been deregistered, debarred or black listed by any govt. /autonomous institution, hospital or body in India.
k. The instruments such as power of attorney, Permanent Account No. (PAN) etc.
l. All pages of documents enclosed with the bid document should also be seal-signed and page numbered given to each and every pages.
m. An undertaking by the manufacturer that they have their own testing laboratories and in built quality assurance facilities and shall carry out batch-wise pre-inspection of the items (covered under the drug and cosmetic act) and submit such reports along with the supplies for each batch. The drug product should have compliances as per specification (IP/BP/USP etc.), if In-house (INH) specification than provision of reference standard and testing protocol for quoted items shall be submitted. Distinct documents for each manufacturer

REQUIREMENT FOR PRINCIPLE MANUFACTURER

a. Manufacturer should have valid License (issued by F.D.A) under the Drugs and Cosmetics Act, 1940 (including its amendments) and should comply with Schedule M (cGMP) of Drugs and Cosmetic Act. The bidder should furnish self-attested photocopy of manufacturing License (Own license/ loan license/ third party license) for the product duly approved by the Licensing authority for each and every product quoted as per specification in the tender. The license must have been duly renewed up to date and the items quoted shall be clearly highlighted in the license. Items for which FDA do not issue manufacturing license but issue repacking license, firm should submit valid repacking license. If revalidation of drug license has been applied, copy of application to State Drug / Licensing authority should be attached.
b. The manufacturer should have received a valid cGMP as per revised Schedule 'M' issued by Licensing Authority or WHO-GMPinspectioncertificateinlinewiththe WHOcertificationscheme, on pharmaceuticals being quoted. Bidder should submit a valid product wise Certificate of cGMP as per schedule "M" or WHO-GMP issued by Regulatory Authority, for each item offered. License no. and manufacturing unit address clearly highlighted on cGMP certificate.
c. A certificate issued by the Licensing Authority that the Manufacturer is not currently under conviction (Non- conviction) under the Drugs & Cosmetics Act for manufacturing/supplying sub-standard drugs or on any other grounds. The certificate should not be more than six month old on the day of opening of the tender.
d. Tender should not be submitted for the product/ products for which the concern manufacturer / company / authorized dealer / distributer has been blacklisted on quality grounds by any Government organization.
e. The bidder should submit a notarized affidavit stating that the manufacturing company have not been blacklisted for the quoted product/firm by any State Government or Central Government Organization and has not been found guilty of supplying spurious drugs in last three years and are eligible to participate in the present tender." If the information provided in the affidavit is found to be incorrect at any stage, during and after the tender, action will be initiated as per the tender conditions apart from forfeiture of performance security deposit (if any).

Requirement For DISTRIBUTER

a. Distributor, Should have valid licenses for sale (under the Drugs and Cosmetics Act, 1940 including its amendments) of concern items. The license must have been duly renewed up to date.
b. GST No.(Distributer).
c. Manufacturer must have valid License (issued by F.D.A) under the Drugs and Cosmetics Act, 1940 (including its amendments) and should comply with Schedule M (cGMP) of Drugs and Cosmetic Act. The bidder should furnish self-attested photocopy of manufacturing License (Own

license/loan license/third party license) for the product duly approved by the Licensing authority for each and every product quoted as per specification in the tender. The license must have been duly renewed up to date and the items quoted shall be clearly highlighted in the license. Items for which FDA do not issue manufacturing license but issue repacking license, firm should submit valid repacking license. If revalidation of drug license has been applied, copy of application to State Drug / Licensing authority may be attached.

d. The manufacturer should have received a valid cGMP as per revised Schedule 'M' issued by Licensing Authority or WHO-GMP inspection certificate in line with the WHO certification scheme, on pharmaceuticals being quoted. Bidder should submit a valid product wise Certificate of cGMP as per schedule "M" or WHO-GMP issued by Regulatory Authority, for each item offered. License no. and manufacturing unit address clearly highlighted on GMP certificate.

e. A certificate issued by the Licensing Authority that the Manufacturer is not currently under conviction (Non- conviction certificate) under the Drugs & Cosmetics Act for manufacturing/supplying sub-standard drugs or on any other grounds. The certificate should not be more than six month old on the day of opening of the tender.

f. Tender should not be submitted for the product/ products for which the distributer has been blacklisted on quality grounds by any Government organization.

g. The bidder should give a notarized affidavit stating that distributer has not been blacklisted for the quoted product/firm by any State Government or Central Government Organization and has not been found guilty of supplying spurious drugs in last three years and are eligible to participate in the present tender." If the information provided in the affidavit is found to be incorrect at any stage, during and after the tender, action will be initiated as per the tender conditions apart from forfeiture of performance security deposit (if any).

h. Authorization letter given by the manufacturers to the distributor for individual molecule.

XVI. Declaration on **Non-Judicial Stamp Paper** for eligibility in participating the tender for quoted drugs in prescribed format as per **Annexure-II**

XVII. Tenderers shall furnish Company's bank details as per **Annexure-III** (Mandate Form) with cancelled cheque.

XVIII. Duly attested Checklist as per (**ANNEXURE- I**) shall be submitted.

XIX. Copy of PAN Card of the bidder company should be submitted (self-attested).

XX. Copy of certificate of valid GST registration of the bidder company should be submitted (self-attested).

XXI. Copy of Non-Conviction Certificate issued by the concerned Licensing Authority from Drug Controller Administration of the State, not older than 12 months

XXII. Copy of Market Standing Certificate issued by the Licensing Authority from Drug Controller Administration of the State of quoted product for last 3 Years

XXIII. The vendor should provide an acceptance letter that if AIIMS, Nagpur unable to consume the supplied drugs, near its expiry vendor needs to replace the same with long expiry drugs without any charge (in companies letter head duly)

XXIV. **Bidder shall declare the % age of local content used in the manufacturing of quoted item in accordance with the calculations for local content of Order issued by Ministry of Chemicals & Fertilizers, Department of Pharmaceuticals vide no. 31026/4/2018-Policy dated 01.01.2019**

Note: -

(i) The certificates/ reports / annexure submitted with the bid document should be self-attested by the authorized signatory of the firm with official seal.

ii) Technical evaluation of the Bid will be done on the basis of the above-mentioned criteria and documents mentioned in Clause no. 3 (TECHNICAL BID- COVER 'A') are Mandatory Documents and shall be submitted online only at CPPP portal: https://eprocure.gov.in Failing which the bid will not be considered for technical evaluation.

iii) The technical evaluation shall be done only on the basis of documents/papers submitted by the bidder on https://eprocure.gov.in

iv) Clear copy of valid drug license and approval list highlighting the quoted drugs with AIIMS, Nagpur drug code should be uploaded. In case scanned copy of license uploaded is not visible or tempered or quoted drugs are not highlighted, AIIMS, Nagpur shall not consider the license for such drug.

4. GENERAL CONDITIONS:

(i) Bidders are advised to quote only for such drugs which meets the drug specification as mentioned in **Annexure VIII.** Do not quote if it differs with regard to any parameter.

(ii) The bidder shall submit the complete stability data (long term stability studies and accelerated stability studies) for all awarded drugs whenever required by the AIIMS, NAGPUR. For New drugs, complete stability data of 6 months' period shall be acceptable. (If manufacturer has licensed a formula from another company and such licensed formula is used for the product, then the stability data of the licensor should be submitted along with licensing agreement.)

(iii) The Tenderer should have not been blacklisted/debarred/de-registered/banned due to quality failure for the quoted product /firm by any State Government / Central Government/ AIIMS/Central or State Government's Drug procurement agencies **at the time of submission of bid**. Further, quoted drugs have not been failed to testing by any State Government/Central Government / its Drug procurement agencies during last two years. If any tenderer has been blacklisted/debarred/de-registered/banned due to quality failure, such tenderer or their Partner/Director/Owner shall not be permitted to participate in the tender.

(iv) During the validity of the tender if the firm / Company is blacklisted/debarred/de-registered/banned by any State Government/ Central Government/ AIIMS/ Central or State Government's Drug procurement agencies / convicted by any Court of law in India, it shall be intimated to AIIMS, Nagpur along with relevant authentic document by the tenderer firm/ company within one month.

(i) **If the tenderer/bidder fails to inform about blacklisting/ debarring/ de-registration/ban or punitive actions taken by the state or central government or AIIMS or any state/central drugs procurement agency against the bidder to AIIMS, Nagpur within 30 days and the bidder continues the contract and the said action is brought/comes to the knowledge of AIIMS, Nagpur, AIIMS Nagpur may ask an explanation from the tenderer which shall be clarified by the tenderer within 7 days of receiving the letter from AIIMS, Nagpur. If the tenderer fails to reply within 7 days or reply is not to the satisfaction of AIIMS, Nagpur, all the contracts awarded to the tenderer will be terminated, performance security deposits will be forfeited, and further tenderer will be backlisted up to 5 years from participating into any tender invited by AIIMS, Nagpur.**

(v) During tender or Rate Contract period, if L1 bidder is debarred/deregistered /blacklisted/ banned by any Central Government or State Government or its procurement agencies due to

quality failure, AIIMS, NAGPUR may purchase the drugs from other bidders at L1 rate or may go for fresh tender as per discretion of AIIMS, Nagpur.

(vi) The Tenderer should confirm that they have read tender document including Amendment(s) to Tender document (if any) along with terms and condition and these terms and condition of tender document including Amendment(s) to Tender document (if any) are acceptable unconditionally to them.

(vii) **Validity of Rate Contract: -The rate contract will be applicable for one year from the date of issuance of Rate contract. The validity of contract may be extended one or more years with mutual consent, if necessary.**

(viii) Only authorized employee of the Company/Tenderer will be allowed to transact the business with the Tender Inviting Authority.

(ix) The drug formulation like injection, oral drugs and Tablets, rates should be quoted only for the composition stated in the tender. Blood products should be supplied only after getting HIV and Hepatitis-B screening certificate. A copy of these Certificates should be sent with every consignment and every invoice.

(x) No bidder shall be allowed at any time on any ground, whatsoever it may be, to claim revision or modification in the rates quoted by him. Representation to make correction in the tender documents on the ground of Clerical error, typographical error, etc., committed by the bidders in the Bids shall not be entertained after submission of the tenders. Cross Conditions such as "SUBJECT TO AVAILABILITY" "SUPPLIES WILL BE MADE AS AND WHEN SUPPLIES ARE RECEIVED" etc., will not be entertained under any circumstances and the tenders of those who have given such conditions shall be treated as incomplete and accordingly the bid will be rejected.

(xi) **Purchase Preference to Local Suppliers**

In pursuance of Government of India Order No. P-45021/2/2017-B.E.-II dated 16th September 2020 (as amended from time to time) and F.No. Z.28018/67/2017-EPW dated 12th June 2018 purchase preference shall be given to local suppliers in all procurements undertaken in the manner specified hereunder and the procurement shall be made as per terms and conditions contained in the said order.

- **Minimum local content:** The minimum local content shall as per Government of India Order No. P- 45021/2/2017-B.E.-II dated 16th September 2020 (as amended from time to time) and F.No. Z.28018/67/2017-EPW dated 12/06/2018, till the Nodal Ministry prescribes a higher or lower percentage.
- **Margin of Purchase Preference:** The margin of purchase preference shall be 20%. The Local supplier whose quoted price falls in the margin of purchase preference desirous of claiming benefit of the Order No. P-45021/2/2017-B.E.-II dated 16th September 2020 shall submit an undertaking within 7 days of opening of financial bid, that he would be ready to supply the product at L1 price. In case of non-receipt of the same, he would not be given purchase preference.
- The bidders are required to submit the following annexure in compliance of public procumbent (Preference to Make in India) order, 2017: Affidavit of self-certification regarding local content (to be provided on Rs. 100/- stamp paper as per **Annexure VI.**
- **All other terms & conditions will be as per the Department of Industrial Policy and Promotion (DIPP) order No. P-45021/2/2017-B.E.-II dated 16th September 2020 (as amended from time to time).**

(xii) **Land Border Sharing** – as per Certificate regarding compliance Rule-144 (xi), any bidder from such countries sharing a land border with India will be eligible to bid in any procurement whether of goods, services (including consultancy services and non-consultancy services) or works (including turnkey projects) only if the bidder is registered with the Competent Authority. Bidders to submit self-declaration on their letter head as per **Annexure- VII**

5. PRICE BID – "COVER-B" (Financial Bid/BOQ)

A) Cover "B" (Financial Bid/BOQ) contains the Price Bid of the Tenderer. The Tenderer shall fill in the rate per unit size, rate of GST in respective column of BOQ for the items quoted**.**

B) Determination of L1 Bidder:

In determining the lowest evaluated price, the rate quoted per unit size for the given specification, exclusive of GST as indicated in the **BOQ** shall be taken into consideration. **The rates quoted should be in rupees and paisa up to 2 digits.**

C) GST (Goods and Services Tax)-The Tenderers must indicate the rate of GST applicable and payable by them. In case no information is given, it shall be presumed that rate is inclusive of GST and no GST shall be charged by them under any circumstances.

6. EARNEST MONEY DEPOSIT:

Money Deposit **(i.e. ₹ 2,00,000)** to be deposited in the form of Demand payee Demand Draft/ FDR/Banker's Cheque or BG (including e-Bank Guarantee) from any of the commercial Banks. Scanned copy to be enclosed with technical bid. It is also clarified that the bids submitted without earnest money will be summarily rejected. The Demand Draft/ FDR/Banker's Cheque or BG (including e-Bank Guarantee) may be prepared in the name of "The Director, AIIMS, Nagpur". The EMD (Demand Draft/ FDR/Banker's Cheque or BG (including e-Bank Guarantee) or any exemption certificate) must reach at Store Office (Hospital Store), First Floor, IPD, AIIMS, Plot No. 2, Sector- 20, MIHAN, Nagpur prior to opening of tender.

- No request for transfer of any pervious deposit of earnest money or security deposit or payment of any pending bill held by the AIIMS Nagpur in respect of any previous supply will be entertained. Tenderer shall not be permitted to withdraw his bid or modify the terms and conditions thereof. In case the tenderer fail to observe and comply with stipulations made herein or backs out after quoting the rates, the aforesaid amount of earnest money will be forfeited

- Tenders without Earnest Money will be summarily rejected.

- No claim shall lie against the AIIMS Nagpur in respect of erosion in the value or interest on the amount of EMD.

- If MSME firm (only Micro and Small Enterprises) is registered for above tendered item, then the firm will be exempted for submission of EMD amount. Firm must upload scanned copy of following documents in support of exemption.

a) District Industries Centers (DIC)

b) Khadi and Village Industries Commission (KVIC)

c) Khadi and Village Industries Board

d) Coir Board

e) National Small Industries Corporation (NSIC)

f) Directorate of Handicraft and Handloom

g) Any other body specified by Ministry of MSME (MoMSME)

h) Udyog Aadhaar Acknowledgment/Udyog Aadhaar Memorandum/Udyam issued by MoMSME.

i) Startups firms as recognized by Department of Industrial Policy & Promotion (DIPP) is also exempted for depositing of EMD amount. Valid documents should be uploaded.

- The earnest money will be returned/refund to the unsuccessful tenderers after the tender is decided.

- EMD should remain valid for a period of 180 days beyond the final bid validity period. When the tenderer agrees to extend the validity of bid, he shall also extend the validity of EMD suitably.

The Purchaser reserves the right to relax the Norms on Prior Experience for Start-ups and Micro & Small Enterprises in Public Procurement. The Start-ups are defined in Annexure-A of the "Action Plan for Start-ups in India". The same is available on the website of Department of Industrial policy and Promotion (DIPP), Ministry of Commerce & Industry.

7. GUIDELINES FOR THE PREPARATION OF TENDER:

A) The bidder shall bear all costs associated with the preparation and submission of its bid and Tender Inviting Authority will in no case be responsible or liable for these costs, regardless of the conduct or outcome of the bidding process.

B) **Language of Bid: -** The Bid prepared by the bidder and all correspondence and documents relating to the bid exchanged by the bidder and the Tender Inviting Authority, shall be in English language, Supporting documents furnished by the bidder may be in other languages provided they are accompanied by an authenticated (by the authority concerned) accurate translation of the relevant passages in the English language in which case, for purposes of interpretation of the Bid, the English translation shall alone govern. Failure to submit authentic translation of documents would result in rejection of bids. No bid can be partly in one language and partly in another language.

C) Power of Attorney or Resolution of the Board by which the authorized signatory has been authorized by the bidder firm should sign the documents in cases where person other than the Managing Director/Managing Partner or sole Proprietor signs the document.

8. PERIOD OF VALIDITY OF TENDER:

a) The tender must remain valid for minimum 180 days from the date of opening of Bid.

b) Prior to the expiration of the bid validity the Tender Inviting Authority may extend the bid validity for further period with mutual consent of the bidder.

c) The bidder who has extended the bid validity is not required or permitted to modify its bid.

d) The bidder cannot withdraw the bid within validity of Tender.

9. AMENDMENT OF TENDER DOCUMENTS:

At any time prior to the last date of submission of online bid, Tender Inviting Authority may, for any reason, whether on own initiative or in response to a clarification requested by a prospective Tenderer, may modify the condition in Tender documents by uploading an amendment on AIIMS, Nagpur website: ***www.aiimsnagpur.edu.in;*** and on CPP portal i.e., ***https://eprocure.gov.in*** will be binding on them. In order to provide reasonable time to take the amendment into account in preparing their bid, Tender Inviting Authority may at discretion, extend the date and time for submission of online bid.

A) Bidders are advised to check the *website of AIIMS, NAGPUR:* ***www.aiimsnagpur.gov.in;*** and CPP Portal i.e., ***https://eprocure.gov.in***; *regularly* for any corrigendum or amendment to the tender document.

B) AIIMS, Nagpur will not issue separate communication for any corrigendum or amendment.

10. MODIFICATION AND WITHDRAWAL OF BIDS:

A) The bidder may modify or withdraw its bid after the bid submission before last time and date of submission of online Technical Bid.

B) No bid will be allowed to be withdrawn after the last date & time of submission of online Technical Bids.

11. EVALUATION OF TENDER:

A) Technical evaluation of the Bid will be done on the basis of criteria and documents mentioned in
S.N. 3 (TECHNICAL BID-COVER A) & Annexure I (Check List) which are present on the CPP Portal i.e. https://eprocure.gov.in.

B) Bids of firms who have furnished all the required documents for each of the product quoted

will be considered.

C) If at any stage, it is found that the contract has been successfully obtained by the bidder by submitting forged/fabricated certificates/documents/licenses and/or by concealing the fact about blacklisting/debarring/de-registration of the firm by Govt. of India/Suspension/Cancellation/non- renewal of the manufacturing license of the bidder firm, the tender bid/rate contract may be rejected/terminated and suitable punitive action may be taken against the firm.

D) **In event of financial bid opening, due to provisions/compulsion of e-tendering system if complete quoted product list of financial bids of a bidder is opened then only those financial bids of quoted product shall be considered of whose technical bid has been found eligible by the Technical Evaluation Committee**.

E) **After evaluation of technical bid of tenderer/bidder, AIIMS, Nagpur may ask the objection/clarification from tenderer/ bidder**. (if found necessary by the committee)

12. ACCEPTANCE /REJECTION OF BIDS:

AIIMS, Nagpur reserves the right to accept or reject the tender for the supply of all or any one or more items of the drugs tendered for in a tender without assigning any reason.

13. PACKING

All primary packing containers/strips/blister should be strictly conforming to the Specification included in the relevant pharmacopoeia. Packing should be able to prevent damage or deterioration during transit. The labels in the case of injectable should clearly indicate whether the preparations are meant for Intravenous (IV), Intra Muscular (IM), Subcutaneous (SC), etc. Primary packing such as strips, labels, inner carton, outer carton etc. should bear the

following words “Govt. Supply- Not for Sale”

Secondary packing such as baby shipper (small corrugated box), outer corrugated boxed are labelled as under.

GOVT. SUPPLY-NOT FOR SALE Name of the Drug: Manufactured by: Batch no.: Mfg. Date : Exp. Date: Quantity:

14. AWARD OF CONTRACT:

A) The Purchaser will award the contract to the bidder whose quotation has been determined to be substantially responsive and who has bided the lowest evaluated quotation price.

B) Notwithstanding the above, the Purchaser reserves the right to accept or reject any quotations and to cancel the bidding process and reject all quotations at any time prior to the award of contract.

C) The bidder whose bid is accepted will be notified of the award of contract by the Purchaser prior to expiration of the bid validity period. The terms of the accepted bid shall be incorporated in the purchase order.

D) Rates should be quoted inclusive of GST & other charges (if applicable).

E) The competent authority reserves all rights to reject the goods if the same are not found in accordance with the required description/specifications/quality.

F) A brochure displaying clearly the product is to be attached with the tender if required.

G) Other terms and condition not mentioned above shall be applicable as per GFR-2017 and Manual for Procurement of Goods 2017.

15. PERFORMANCE SECURITY DEPOSIT:

- The successful bidder shall have to submit a performance guarantee (PG) within 30 days from the date of issue of Rate Contract. Extension of time for submission of PG beyond 30 days and up to 60 days from the date of issue of Rate Contract may be given by the competent authority to sign the contract agreement however a panel interest of 15% per annum shall be charged for the delay beyond 30 days. i.e. 31st day after the date of issue of Rate Contract. In case of the contract fails to submit the requisite PG even after 60 days from the date of issue of Rate Contract, the contract shall be terminated and other dues if any payable against the contract. The failed contractor shall be debarred from participating in re-tender (if any) for that item. Performance Guarantee Bond is mandatory.
- Successful supplier/firm should submit a performance security in the form of **Account pay Demand Draft, Fixed Deposit Receipt from a commercial bank, Bank Guarantee (including e-Bank Guarantee)** from any commercial Bank duly pledged in the name of the **"The Director, AIIMS Nagpur"** and to be received in the Store Office (Hospital Store) All India Institute of Medical Sciences, Plot No. 2, Sector20, MIHAN, Nagpur– 441108 before the date of commencement of supply or 30 days from the date of acceptance of the Rate Contract, whichever is earlier. The performance guarantee bond to be furnished in the form of Bank Guarantee as per given Proforma of the tender documents for an amount of 3% of the contract/projected purchase order value with minimum of **Rs. 1,00,000 to Rs. 3,00,000/-**
- The Performance Guarantee should be established in favour of "**The Director, AIIMS Nagpur**" through any commercial Bank with a clause to enforce the same on their local branch at Nagpur.
- Validity of the performance guarantee bond shall be for a period of 60 days beyond of entire contract period (i.e 14 Months), which needs to be extended if agreed for extension of Rate contract.

16. METHODOLOGY FOR PLACING ORDERS:

For the above purpose the following procedures will be adopted;

A) After the conclusion of Price Bid opening (Cover B), the rates offered by tenderers for each product are evaluated and lowest acceptable rate (L1 Rate) arrived at is declared and that tenderer is informed**.**

B) If two or more than two Tenderer's are declared as lowest suppliers for the same item(s), such Tenderers are eligible for Rate Contract and the placement of Purchase Orders for such item(s) for which they are declared as lowest.

C) The supplier shall supply the Drugs/Medicines to **AIIMS, Nagpur** along with copy of Purchase order, copy of test reports and 3 original copies of Invoice.

D) A purchase order is placed on supplier for supply of definite quantity in terms of Rate Contract during validity period of Rate Contract that purchase order is valid and binding contract.

E) No Minimum drawl is in the Rate Contract. The actual quantity may vary from nil to maximum required quantity during validity of Rate Contract.

F) The purchaser has the option to renegotiate the price with the rate contract holders. In case of emergency, the purchaser may purchase the same item through Ad hoc contract with a new supplier.

G) Purchase orders, incorporating definite quantity of drugs/products to be supplied along with all other required conditions following the rate contract terms, shall be issued for obtaining supplies through the rate contract.

H) The purchaser is entitled to place purchase orders up to the last day of the validity of the rate contract and, though supplies against such purchase orders will be affected beyond the validity period of the rate contract, all such supplies will be guided by the terms & conditions of the rate contract.

I) Any supply without a valid purchase order will not be acceptable by AIIMS, Nagpur and the AIIMS shall not be responsible for any loss on this account.

J) AIIMS, Nagpur reserve its rights to cancel the supply order without assigning any reason.

K) The rates quoted shall not be varied with the Purchase order quantity during the full contract period.

L) The rates quoted and accepted will be binding on the Tenderer for the full contract period of one year and any increase in the price will not be entertained till the completion of this contract period. Accordingly, this clause will be applicable for all orders placed during the contract period. However, Rate Contract validity period may be extended further with mutual concern with the both parties.

M) No Tenderer shall be allowed at any time and on any ground, whatsoever it may be, to claim revision or modification in the rates quoted by them. Representation to make correction in the tender documents on the ground of Clerical error, typographical error, etc., committed by the Tenderers in the Bids shall not be entertained after submission of the tenders. Cross Conditions such as "SUBJECT TO AVAILABILITY", "SUPPLIES WILL BE MADE AS AND WHEN SUPPLIES ARE RECEIVED" etc., will not be entertained under any circumstances and the tenders of those who have mentioned such conditions shall be treated as incomplete and accordingly the Tender will be summarily rejected.

N) The supplier shall take utmost care in supplying the quality Drugs/Medicines and ensure that the batch number mentioned in the packages of the Drugs/Medicines tally with the batch number mentioned in the Invoice produced to AIIMS, Nagpur for payment. Also, the supplier shall ensure the quantity relevant to the Batch Number of the Drugs/Medicines is mentioned in the invoice.

O) The Rate Contract (RC) awarded under the present tender enquiry will be in the nature of standing offer. Purchase Order (PO) may be placed from time to time against Rate Contract (RC).

P) The procurement agency has the liberty to place orders for drugs under MJPJAY or any other ongoing scheme at AIIMS Nagpur through the concerned office, with the approval of the Director, AIIMS Nagpur.

17) SUPPLY CONDITIONS:

(i) Purchase orders along with the delivery destinations will be placed on the successful bidder at the discretion ofthe Ordering Authority.

(ii) All supplies will be scheduled for the period from the date of acceptance till the completion of the tender in installments, as may be stipulated in the Purchase Order. The supplied medicines and Drugs (covered in Schedule P of Drugs & Cosmetics Act) should have a maximum potency throughout the shelf life period as prescribed inthe Drugs & Cosmetics Act 1940 and rules there under. All other items of drugs and medicines should have a shelf life period of minimum 1 year (Except in that drug where self-life is recommended less than 1 year as per drug & cosmetic act 1940 but not less than 75% of expected shelf life) from the date ofmanufacture.

(iii) The supply should be completed within 30 days from the date of purchase order. The supplier may continue the supply of unexecuted quantity after the issue of Amendment for Delivery period extension, however liquidated damages as specified in Penalty Provision of the tender conditions, will be levied on the quantity supplied after the expiry of original DP). Unloading of material will be done by supplier.

(iv) The supplier shall complete the earlier purchase order before commencing the supply of subsequent purchase orders. In case of non- execution, AIIMS Nagpur reserves the right to place purchase order (partially/ fully) on alternate source at the risk and cost of the defaulting bidder.

(v) The Bidder must submit a Test Analysis report (Certificate of Analysis) for every batch of drug along with other documents. If P.O. for single molecules is more than 50,000/- (Rupees Fifty Thousand), company has to be submit the Test Analysis report (Certificate of Analysis) from any of the recognized Govt. approved NABL accredited lab at the time of supply & if P.O. for single molecules is up to 50,000/- (Rupees Fifty thousand), In-house lab testing reports are mandatory.

In case of failure on part of the supplier to furnish such report, the batch of drugs will be returned back to the suppliers and the bidder is bound to replenish the same with Government approved lab test report. The Drugs and Medicines supplied by the successful bidder shall be of the best quality and shall comply with the specifications, stipulations and conditions specified in the tender.

(vi) If the bidder fails to execute the supply within the stipulated time, the Tender Inviting/Ordering Authority is at liberty to make alternative arrangement for purchase of the items of drugs and medicines for which the Purchase orders have been placed, from any other sources or in the open market or from any other bidder who might have quoted higher rates, at the risk and the cost of the supplier and in such cases the tender inviting authority /ordering authority has every right to recover the cost and impose the penalty without prejudice to the rights and remedies available with the Purchaser under the Law.

(vii) It shall be the responsibility of the Bidder for any shortages/damage at the time of receipt. Tender inviting authority is not responsible for the stock of the drug received, for which no order is placed.

(viii) The bidder needs to replace the expired Drugs which are not utilized by the tender inviting Authority within the shelf life period, all the cost regarding this needs to be borne by the bidder/vendor. If the bidder failed to do the same, value of expired drugs amount will be forfeited from the performance security deposit.

(ix) Bidder must comply to the shelf life of each quoted drugs in accordance with Schedule P of Drugs and Cosmetics Rules, 1945. In case drug/medicine not covered in Schedule P of Drugs and Cosmetics Rules, 1945, the bidder must supply the drug/medicine with **minimum 24 months shelf life**. Bidders must declare the required shelf-life detail in Para V of Annexure II.

18. Force Majeure:

If, at any time during the subsistence of this contract, the performance in whole or in part by either party of any obligation under this contract is prevented or delayed by reasons of any war or hostility, act of public enemy, civil commotion, sabotage, fire, floods, exception, epidemics, quarantine restriction, strikers lockout or Act of God (hereinafter referred to as events) provided notice of happening of any such eventuality is given by party to other within 21 days from the date of occurrence thereof, neither party hall by reason of such event be entitled to terminate this contract nor shall either party have any claim for damages against other in respect of such non- performance or delay in performance and deliveries have been so resumed or not shall be final and conclusive. Further, that if the performance in whole or in part of any obligation under this contract is prevented or delayed by reason of any such event for a period exceeding 60 days, AIIMS, Nagpur party may, at least option to terminate the contract.

19. QUALITY TESTING & QUALITY CONTROL:

A. All the batches of the drugs supplied shall be supported by test/ analysis reports furnished by independent NABL Accredited Drugs Testing Laboratory/Central Drug Testing Laboratory/In House Quality Control Laboratory. The Tender Inviting Authority has the right to get the drugs tested at the laboratories of his choice for further verifications, from AIIMS, NAGPUR empanelled laboratories. **Bidder should submit the Certificate of Analysis (CoA) including**

leakage test, hardness (for uncoated Tablets: NLT 2kg/cm^2, for filmcoated Tablets: NLT 3kg/cm^2) **and friability test** (limit of uncoated Tablet NMT 1%) **wherever applicable.**

B. Random samples of each supplied batch will be chosen at the point of supply or distribution/storage points for testing. The samples will be sent to different AIIMS, NAGPUR empaneled laboratories including Government Drugs Testing Laboratory/NABL accredited lab. Handling and testing charges will be deducted by AIIMS, NAGPUR for the above purpose.

C. The Drugs shall have the active ingredients at the prescribed level as indicated in official compendiums throughout the shelf-life period of the drug. The samples will be drawn periodically throughout the shelf-life period and if found "Not of Standard Quality", the cost of entire batch paid will be recovered whether consumed fully/partially. Also, action will be initiated for blacklisting as per the points mentioned in the tender document irrespective of the period of supply. The supplies will be deemed to be completed only upon receipt of the quality certificates from the laboratories. Samples which do not meet quality requirement shall render the relevant batches liable to be rejected. If the sample is declared to be "Not of Standard Quality" or spurious or adulterated or misbranded, such batch/batches will be deemed to be rejected goods.

D. In the event of the samples of Drugs supplied fails in quality tests or found to be not as per specifications, the AIIMS, NAGPUR is at liberty to make alternative purchase of the items of drugs for which the Purchase orders have been placed from any other sources or in the open market or from any other Tenderer who might have quoted higher rates, at the risk and the cost of the supplier and in such cases the AIIMS, NAGPUR has every right to recover the cost and impose penalty as mentioned in Tender document.

E. If the product / sample fails in quality test, every failed batch shall be taken back by the supplier at their own cost and AIIMS, NAGPUR shall not be responsible for any damage during this period.

F. The supplier shall furnish evidence of the basis for shelf life and other stability data concerning the commercial final package on request by the AIIMS, NAGPUR. In case of any complaint in the field, the B.M.R/ B.P.R for the particular batch of the product(s) supplied shall be produced when demanded. **For New drugs, complete stability data of 6 months' period shall be acceptable.**

G. **The** products should conform to the standards of IP/BP/USP/EP/JP as the case may be. **However, the drugs notified in the IP (amended up to date) shall be accepted only if supplied conforming to the standards outlined in the IP.** In case the product is not included in the any of the said compendiums, the supplier, upon award of the contract, must provide the reference standards and testing protocols for quality control testing.

H. The case of admixture of drugs will be treated as a violation of tender conditions and fine will be levied as per tender conditions. If such lapses happen more than twice in a tender period such cases will be treated as "Misbranded Drugs".

20. PAYMENT PROVISION:

1. Payment Terms: - Payment shall be made subject to recoveries, if any, by way of liquidated damages or any other charges as per terms & conditions of contract in the following manner.
2. 100% payment of the contract price shall be paid on receipt and acceptance of goods in good condition at the consignee premises and subject to recoveries, if any, either on account of defects/ deficiencies not attended by the supplier or otherwise and upon the submission of the following documents:
3. Four copies of suppliers invoice showing contract number, goods description, quantity, unit price and total amount with revenue stamp.

4. Two copies of packing list identifying contents of each package.
5. The supplier shall not claim any interest on payment under the contract.
6. Where there is a statutory requirement for tax deduction at source, such deduction towards income tax and other tax as applicable will be made from the bills payable to the supplier rates as notified from time to time.
7. No payment shall be made for rejected stores. Rejected item/equipment must be removed by the supplier within two weeks of the date of issue of rejection advice at their own cost & replace immediately. In case these are not removed these will be auctioned/disposed of at the risk and responsibility of the suppliers without notice.

21. LIQIDATED DAMAGES & OTHER PENALTIES:

a. If the suppliers fail to deliver and place any or all the item or perform the service by the specified date as mention in purchase order, penalty at the rate of 0.5% per week of delayed value of goods subject to the maximum of 10% of delayed goods value will be deducted, afterwards another penalty may be imposed.

b. If the complete supply or part thereof is received in damaged condition it shall not be accepted and shall be recorded on Delivery Challan. Such damaged material should be replaced by the supplier within 14 days from the date of noting on Delivery Challans or rejection advice issued by consignee or else subsequent to no replacement in 14 days the Performance security (SD) would be forfeited with a notice to the supplier. In case of damage only in the outer packing, the supply will be accepted only after levying penalty of 1% on the total value of the supply to that destination place. Further the Performance Security (SD) would be forfeited with a notice to the supplier.

c. Tender Inviting Authority will be at liberty to terminate, without assigning any reasons thereof, the contract either wholly or in part on 30 days' notice. The Bidder will not be entitled for any compensation whatsoever in respect of such termination. All litigations related to the supplier for any defaults will be done by Tender Inviting Authority and his decision will be final and binding.

22. DEDUCTION & OTHER PENALTIES ON ACCOUNT OF QUALITY FAILURE:

A) If the samples do not conform to statutory standards, the Tenderer will be liable for relevant action under the existing laws and the entire stock in such batch has to be taken back by the Tenderer within a period of 30 days of the issue of the letter from the AIIMS, NAGPUR. Such stock shall be taken back at the expense of the Tenderer. Further, actual handling and testing charges shall be paid to AIIMS, NAGPUR by the supplier otherwise these charges shall be recovered from their pending bill/Performance Security Deposit.

B) The AIIMS, NAGPUR has the right to destroy such "NOT OF STANDARD QUALITY DRUGS" if the Tenderer does not take back the goods within the stipulated time. The AIIMS, NAGPUR will arrange to destroy the "NOT OF STANDARD QUALITY DRUGS" after the expiry of 30 days mentioned above without further notice and shall also collect handling charges (in case the product is sent back to supplier on freight to pay basis)/ demurrage charges calculated at the rate of 2% per week on the value of the drugs rejected till such time stipulated. Further, the cost of disposal shall be recovered from the supplier.

C) If any items of Drugs/Medicines supplied by the Tenderer have been partially or wholly used or consumed after supply and are subsequently found to be in bad odour, unsound, inferior in quality or description (Adulterated/Spurious/Misbranded) or otherwise faulty or unfit for consumption, then the contract price or prices of total such batches supplied will be recovered from the Tenderer, if payment had already been made to him. In other words, the Tenderer will not be entitled to any payment whatsoever for Items of drugs found to be of "NOT OF STANDARD QUALITY" whether consumed or not consumed and the Tender Inviting Authority is entitled to deduct the cost of such batch of drugs from any amount payable to the Tenderer. On the basis of the nature of failure, action will be initiated to blacklist the

product/supplier.

D) For the supply of Adulterated/Spurious/Misbranded, as defined in the Drugs and Cosmetics Act, 1940, to AIIMS, NAGPUR. AIIMS, NAGPUR reserves the right to blacklist the supplier. No further supplies shall be accepted from the firm/company.

If the tenderer is blacklisted, the tenderer shall also not be eligible to participate in tenders of Tender Inviting Authority of AIIMS, NAGPUR for supply of Drugs for a period of 5 years from the date of blacklisting.

In case of supply of NOT OF STANDARD QUALITY drug(s) to AIIMS, NAGPUR, the product shall be blacklisted by AIIMS, NAGPUR, and no further supplies shall be accepted for the particular drug(s). The Tenderer shall also not be eligible to participate in tenders of AIIMS, NAGPUR for supply of such Drugs for a period of 2 years from the date of blacklisting.

In addition, the Director of Drugs Control of concerned State will be informed for initiating necessary action on the Tenderer in their state. Performance Security Deposit will also be forfeited without any intimation.

E) The Tenderer shall furnish the source of procurement of raw material utilized in the formulations, if required by the AIIMS, NAGPUR. The AIIMS, NAGPUR reserves the right to cancel the purchase orders if the source of supply is not furnished.

F) The decision of The Director, AIIMS, NAGPUR or any officer authorized by him/her, as to the quality of the supplied drugs, medicines etc., shall be final and binding. In such cases, the AIIMS, NAGPUR will be at liberty to terminate, the contract either wholly or in part on 30 days' notice. The Tenderer will not be entitled for any compensation whatsoever in respect of such termination besides forfeiture of Performance Security Deposit.

G) For contravention of the stipulations of the contract or for other justifiable reasons, the contract may be terminated by the AIIMS, NAGPUR, and the Tenderer shall be liable to pay for all losses sustained by the AIIMS, NAGPUR in consequence of the termination which may be recovered from the Tenderer, as per rules besides forfeiture of Performance Security Deposit.

H) Non-performance of any of the contract conditions and provisions will disqualify a firm from participating in the tender for the next 2 years/ Blacklisting the tenderer.

I) In the event of making Alternative Purchase due to failure of supply, partial supply, quality issue & etc, penalty will be imposed on the supplier. The excess expenditure over and above contracted prices incurred by the AIIMS, NAGPUR in making such purchases from any other sources or in the open market or from any other Tenderer who has quoted higher rates and other losses sustained in the process, shall be recovered from the Performance Security Deposit or from any other money due and become due to the supplier and in the event of such amount being insufficient, the balance will be recovered personally from the supplier as per rules.

J) In all the above conditions, the decision of the AIIMS, NAGPUR shall be final and binding.

23. BLACKLISTING IN THE EVENT OF WITHDRAWAL FROM THE TENDER, AND NON-ADHERENCE TO THE QUALITY STANDARDS AND SUPPLY SCHEDULE:

A) BLACKLISTING OF PRODUCT/TENDERER ON WITHDRAWAL OF TENDER

a) If the Tenderer fails to perform the obligations under the tender conditions / commits default in the performance of the Rate contract, such Tenderers will be blacklisted for a period of **2 years** by AIIMS, Nagpur from the date of intimation besides forfeiture of Performance security deposit.

 Signature and Stamp of Bidder

b) The Tenderers who have withdrawn after participating in the tender after the last date and time of submission of online bid, either fully or partially, **the entire firm/company** will be blacklisted for a period of **2 years** from the date of intimation by AIIMS, Nagpur apart from forfeiture of the Security Deposit.

B) BLACKLISTING FOR QUALITY FAILURE/QUALITY TEST BY THE EMPANELLED LABORATORIES OF AIIMS, Nagpur.

a) Each and every batch of drugs/medicines supplied by the supplier shall be subjected to quality test by the Empaneled laboratories or Government laboratory as per the procedure adopted by AIIMS, Nagpur. AIIMS, Nagpur shall also draw the samples of products supplied in the marketplace and get the same tested, to make sure the products are conforming to quality requirements till Self life.

b) If the sample of any batch fails in quality test and report is received stating "Not of standard quality "in any test the report along with the chromatograms etc. such batch of drugs shall be rejected & no further procurement of that drug from the supplier will be taken for two years from the date of sample being declared not of standard quality.

(i) If the supplier challenges and request for retesting, the sample shall be tested at government testing laboratory or reputed govt. institute like NIPER. The test report of govt. lab or NIPER will be final and will be binding to the supplier.

(ii) The cost of such Re-testing shall be recovered from the supplier.

(iv) If **2** batches of item/drug supplied by the same supplier is reported to NOT OF STANDARD QUALITY in specification, then the firm shall be blacklisted for 2 years after observing procedure besides forfeiture of Performance Security Deposit.

(v) If a single batch of any product(s) supplied by the company/firm declared as Adulterated/spurious/ Misbranded by the Government Authorities during the shelf life of the product supplied irrespective of tender period, the company/firm shall be blacklisted for a period of **2 years from the date of intimation & forfeiture of security deposit.**

C) Quality Test by Statutory Authorities:

(i) If any drug is declared "NOT OF STANDARD QUALITY", by any government agencies or drug licensing authority, the issue of available stock of the particular item will be stopped. Further, the available stock of the product in AIIMS, Nagpur will be retrieved.

(ii) If any batch of any product(s) supplied by the company/firm declared, NOT OF STANDARD QUALITY in specification as defined in the Drugs and Cosmetics Act, 1940, by the Government Authorities during the relevant tender period or during quality check within shelf-life period, the company/firm shall be blacklisted for a period of 2 **years from the date of blacklisting** after observing procedure.

D) Procedure for Blacklisting:

(i) On receipt of complaint from user department/patients or report from Govt. Analyst/Drug Testing Laboratory indicating that a particular Item/Drug is "**NOT OF STANDARD QUALITY/ ADULTERATED/ SPURIOUS/MISBRANDED" (**As the case may be), a show cause notice shall be issued to the supplier calling for explanation within 7 days from the date of notice. On receipt of explanation from the supplier, the director, AIIMS, Nagpur may take appropriate action on merits of the case and impose penalty including the blacklisting of the item of the product/company or firm as deemed fit besides forfeiture of Performance Security Deposit.

(ii) If a particular item of the drug has been blacklisted according to the procedure stated above, the

supplier is not eligible to participate in any of the tenders for such item floated by the AIIMS, Nagpur until the period of blacklisting is over.

(iii) If a supplier company/firm is blacklisted according to the procedure stated above, such supplier is not eligible to participate in any of the tenders floated by the AIIMS, Nagpur until the period of blacklisting is over.

E) BLACKLISTING FOR NON-SUPPLY:

Due to non-supply of item against any purchase order, 10% value of purchase order shall be recovered from the supplier in addition of other penal like risk purchase.

The competent authority of AIIMS, Nagpur have rights to blacklist supplier/suppliers for non-supply of items.

24. SAVING CLAUSE:

A) No suit, prosecution or any legal proceedings shall lie against the Tender Inviting Authority or any person for anything that is done in good faith or intended to be done in pursuance of the tender.

B) The tender is being invited following all the orders/rules issued on time to time regarding the supply of Drugs. However, during the course of the dealing, any directions/orders/rules brought before the AIIMS, Nagpur or any provision appears contrary to the orders/ rules passed regarding supply of Drugs, the AIIMS, Nagpur may amend the tender policy as required at any point of time. The said amendment shall be final and binding upon all the parties.

25. RESOLUTION OF DISPUTES

The AIIMS, Nagpur and the supplier shall make every effort to resolve, amicably by direct informal negotiation any disagreement or dispute arising between them under or in connection with the contract.

A) ARBITRATION AND JURISDICTION.

Normally, there should not be any scope of dispute between the AIIMS, Nagpur and the supplier after entering a mutually agreed valid contract/ Rate Contract.

However, due to various unforeseen reasons, problems may arise during the progress of the contract/ Rate Contract leading to disagreement AIIMS, Nagpur and the supplier shall first try to resolve the same amicably by mutual Consultation. If the parties fail to resolve the dispute by such mutual consultation within twenty-one days, then, depending on the position of the case, either the AIIMS, Nagpur or the supplier shall give notice to other party of its intension to commence Arbitration procedure as per Indian Arbitration and Conciliation Act, 1996. Such disputes/differences shall be referred to Sole Arbitrator to be appointed by the director, AIIMS, Nagpur. The venue of Arbitration Shall be at Nagpur. The award published by the Arbitrator shall be final and binding on the parties.

26. CONTACTING THE AIIMS, NAGPUR BY THE BIDDER:

A) No bidder shall contact the AIIMS, NAGPUR on any matter relating to its bid, from the time of bid opening to the time the contract is awarded.

B) Any effort by a bidder to influence the AIIMS, NAGPUR in the Purchaser's bid evaluation, bid comparison or contract award decisions may result in rejection of the bidder's bid.

C) The bidder shall not make any attempt to establish unsolicited and unauthorized contact with the Tender Accepting Authority, Tender Inviting Authority or Tender Scrutiny Committee after opening of the bids and prior to the notification of award and any attempt by any bidder to bring to bear extraneous pressures on the Tender Accepting Authority, Inviting Authority or Tender Scrutiny Committee, shall be sufficient reason to disqualify the bidder.

D) Notwithstanding anything contained in clause (C) above the Tender Inviting Authority or the Tender Accepting Authority, may seek bonafide clarifications from bidders relating to the bids submitted by them during the evaluation of bids.

27. FRAUDULENT AND CORRUPT PRACTICES:

A) For Bidders:

It is purchaser's policy to ensure that suppliers and their authorized representatives/agents observe the highest standard of ethics during the procurement and execution of such contracts. (In this context, any action taken by a bidder, supplier, contractor, or by their authorized representatives/agent, to influence the procurement process or contract execution for undue advantage is improper) 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" is the offering, giving, receiving, or soliciting, directly or indirectly, of anything of value to influence improperly the actions of another party ("another party" refers to a public official acting in relation to the procurement process or contract execution]. In this context, "public official" includes staff and employees of other organizations taking or reviewing procurement decisions.

(ii) "fraudulent practice" is any act or omission, including a misrepresentation, that knowingly or recklessly misleads, or attempts to mislead, a party to obtain a financial or other benefit or to avoid an obligation (a "party" refers to a public official; the terms "benefit" and "obligation" relate to the procurement process or contract execution; and the "act or omission" is intended to influence the procurement process or contract execution).

(iii) "collusive practice" is an arrangement between two or more parties designed to achieve an improper purpose, including to influence improperly the actions of another party ["parties" refers to participants in the procurement process (including public officials) attempting to establish bid prices at artificial, non-competitive level].

(iv) "coercive practice" is impairing or harming, or threatening to impair or harm, directly or indirectly, any party or the property of the party to influence improperly the actions of a party (a "party" refers to a participant in the procurement process or contract execution).

(v) "obstructive practice" is (a) deliberately destroying, falsifying, altering or concealing of evidence material to the investigation or making false statements to investigators in order to materially impede a investigation into allegations of a corrupt, fraudulent, coercive or collusive practice; and/or threatening, harassing or intimidating any party to prevent it from disclosing its knowledge of matters relevant to the investigation or from pursuing the investigation; or acts intended to materially impede the exercise of the purchaser's inspection and audit rights provided for under sub-clause (e) below.

b) Will reject a proposal for award if it determines that the bidder considered for award has, directly or through an agent, engaged in corrupt, fraudulent, collusive, coercive, or obstructive practices in competing for the contract in question.

c) will cancel the contract if the purchaser determines at any time that the bidder, supplier and contractors and their sub-contractors engaged in corrupt, fraudulent, collusive, or coercive practices

d) will sanction a firm or individual, including declaring in eligible, either indefinitely or for a stated period of time, to be awarded a contract if it at any time determines that the

firm has, directly or through an agent, engaged in corrupt, fraudulent, collusive, coercive or obstructive practices in competing for, or in executing, a contract; and

e) will have the right to inspect the accounts and records of the bidders, supplier, and contractors and their subcontractors/authorized representatives and to have them audited by auditors appointed by the purchaser.

B) For Suppliers:

If the AIIMS, NAGPUR determines that a Supplier has engaged in corrupt, fraudulent, collusive, coercive or obstructive practices, in competing for or in executing the Contract, then the AIIMS, NAGPUR may, after giving 7 days' notice to the Supplier, terminate the Supplier's engagement under the Contract and cancel the contract, and the procurement will be made at the risk and cost of the supplier besides

blacklisting the bidder for 5 years with forfeiture of Performance Security Deposit apart from other penal actions.

a) For the purposes of this Sub-Clause:

(i) "corrupt practice" is the offering, giving, receiving, or soliciting, directly or indirectly, of anything of value to influence improperly the actions of another party.

(ii) "fraudulent practice" is any act or omission, including a misrepresentation, that knowingly or recklessly misleads, or attempts to mislead, a party to obtain a financial or other benefit or to avoid an obligation;

(iii) "collusive practice" is an arrangement between two or more parties designed to achieve an improper purpose, including to influence improperly the actions of another party;

(iv) "coercive practice" is impairing or harming, or threatening to impair or harm, directly or indirectly, any party or the property of the party to influence improperly the actions of a party;

(v) "obstructive practice" is (aa) deliberately destroying, falsifying, altering or concealing of evidence material to the investigation or making false statements to investigators in order to materially impede a purchaser investigation into allegations of a corrupt, fraudulent, coercive or collusive practice; and/or threatening, harassing or intimidating any party to prevent it from disclosing its knowledge of matters relevant to the investigation or from pursuing the investigation; or (bb)acts intended to materially impede the exercise of the purchaser's inspection and audit rights provided for.

28. JURISDICTION:

In the event of any dispute arising out of the tender such dispute would subject to the jurisdiction of the Civil Court within the city of Nagpur only.

Annexure - I

CHECK-LIST (Whether uploaded the documents)

(Technical bid/cover "A")

(failure to submit the below mentioned documents, your bid will be rejected)

S.N.	Check List	YES /No	Page No.	Remarks
1	Check list – **ANNEXURE – I**			
2	**ANNEXURE –II** (Declaration on **non-judicial Stamp Paper of Rs. 100/-** for eligibility in participating the tender) to be submitted			
3	Earnest Money deposit for **Rs. 2,00,000/-** in the form of DD/FD/BG			
4	Udyam Registration Certificate (If claimed for EMD exemption) as per Clause No. 6 (B) Note.			
5	Copies of documentary evidence for the constitutions of the company / Firm/ Proprietorship such as Memorandum and Article of Association, Partnership deed with complete address (if applicable)			
6	Power of attorney or Resolution of board by which the authorized signatory has been authorized by the Tenderer to sign the tender documents as per clause (if applicable)			
7	Copy of valid Manufacturing License of the product quoted with latest license renewal certificate			
8	Self-attested valid drug licenses issued to distributer by local FDA. The license must have been duly renewed up to date			
9	Mention that the bidder is Manufacture /Distributor /Dealer / Trader/Supplier **relevant document** should be uploaded			
10	Firm/company registration certificate			
11	In case of distributor/dealer/trader/supplier must upload tender specific authorization certificate from OEM/ manufacturer (**Annexure-V**)			
12	Copies of the Audited Balance Sheet, Profit and Loss statement showing details of their **annual average turnover** for last three consecutive financial years (2020-21, 2021-22,2022-23) not less than **Rs. 50 Lakh** duly certified by charted accountant.			
13	Tenderer must provide evidence of having supplied government hospital / reputed private hospital organizations in India similar nature of items of **at least ₹ 50 Lakh** of Supply of Drug and Medicine of Tender value in the last three years and the copy of the same should be uploaded			
14	Relevant brochure / catalogue pertaining to the items quote with full specification etc.			
15	PAN and GST Registration certificate			
16	Copy of Market Standing Certificate issued by the Licensing Authority from Drug Controller Administration of the State of quoted product for last 3 Years			
17	Copy of Non-Conviction Certificate issued by the concerned Licensing Authority from Drug Controller Administration of the State, not older than 12 months			
18	Copies of **WHO-GMP** (WHO-Good Manufacturing Practice) certificate as per revised Schedule- 'M' of the manufacturing unit issued by the Licensing Authority/ Drugs Control Department/FDA. The WHO-GMP certificate must be valid as on the last date of submission of tender			
19	**ANNEXURE-III (Mandate form)** with cancelled cheque to furnish company bank details			
20	**ANNEXURE-IV** indicating manufacturing License, validity of license and market standing certificate details			
21	**ANNEXURE-VI** (Declaration of Local Content as per clause no.) ***should mention percentage of Local content if not mentioned the bid will be rejected***			

Signature and Stamp of Bidder

22	Authorization letter nominating an employee of the tenderer to transact the business with the Tender Inviting Authority			
23	All pages of documents enclosed with the bid document should also be seal-signed and page numbered given to each and every page.			
24	The vendor should provide an acceptance letter that if AIIMS, Nagpur unable to consume the supplied drugs within expiry period, near expiry drugs needs to replace by the vendor with long expiry without any charge (in firms letter head)			
25	Border sharing clause – **Annexure – VII** (If applicable)			
26	Technical compliance report (for the drugs quoted)			

Name of authorized signatory: …….

Signature of authorized signatory: …….

Company seal:

ANNEXURE –II

(On non-judicial Stamp Paper of Rs. 100/-)

DECLARATION

I/We M/s .....represented by its Proprietor/Managing Partner /Managing Director having its registered office at .....and its factory premises at……..……..…… ……do hereby declare as under: -

(I) that I/we have carefully read all the terms and conditions of tender in ref. no**. AIIMS-NAG/Drugs/RC/OTE/HOS/24-25/04 dated 13/06/2024** including Amendment(s) to Tender document (if any) issued by All India Institute of Medical Sciences (AIIMS, NAGPUR), Nagpur, 441108 and accept unconditionally all terms and condition of tender document including Amendment(s) to Tender document Also I/We have not tampered /modified the tender documents in any manner. In case the same is found tampered/ modified, I/We understand that my/our offer shall be summarily rejected and I/We are liable to be banned from doing business with AIIMS Nagpur and/or prosecuted as per laws (if any).

(II) A. that I/We are holding and have uploaded (a) valid drug license for quoted drugs,(b) valid WHO-GMP certificate , (c) 3 years market standing certificate for quoted products issued by licensing authority, (d) valid non conviction certificate not older than 12 months, (e) the copies of the specifications for all quoted drugs and STP (standard testing procedure) for Non- Pharmacopoeia quoted drugs and also enclosed all undertaking/declaration as per Annexure mentioned in the tender document.

B. that I/We shall submit the complete stability data (long term stability studies and accelerated stability studies) for all awarded drugs within 15 days from the date of issue Letter of Acceptance. (If manufacturer has licensed a formula from another company and such licensed formula is used for the product, then the stability data of the licensor shall be submitted along with licensing agreement.)

On the basis of above undertaking/declaration, the price bid shall be opened subsequently after opening of technical bid. However, any document uploaded with technical bid is not complying as per undertaking, the contract/ Rate Contract shall be cancelled with forfeiture Performance Security Deposit/Bank guarantee (if any) against tender no. **AIIMS-NAG/Drugs/RC/OTE/HOS/24-25/04 dated 13/06/2024** along with other action including suspension/disqualification of contract.

(III) A. I/We declare that we/manufacturer possess the valid drug manufacturing license for AIIMS, Nagpur tendered items as per details below:

Sr. No.	Drug Code	Description of Drug as per AIIMS, Nagpur Tender	Drug Lic. No.	Date of Issue	Validity of Drug Lic.	Address of (OEM) Manufacturing Unit

B. I/We declare that we possess the valid WHO-GMP (World Health Organization-Good Manufacturing Practices) Certificate issued by competent authority and complies and continue to comply with the condition lied in schedule M of Drug & cosmetic act, 1940 the rules made there under.

C. I/We declare that the information of local content provided in Annexure VI is correct.

I am / We are aware of the Tender inviting Authority's right to forfeit the Performance Security Deposit and suspending/disqualifying/blacklist me/us for a period of 5 years if, any information furnished by us proved to be false at time the of inspection and not complying the condition as per schedule M of the said Act for a period of five years.

(IV) that in pursuant to the conditions in Clause No. 6. (A) of the tender, the bids can be suspended/disqualified by the Tender Inviting Authority in case of violation of any of the conditions and non-performance of the obligation under tender document.

(V) that our company/applied items have not been blacklisted/debarred/de-registered/banned **due to quality failure for the quoted product /firm by any State Government / Central Government/ Central or State Government's Drug procurement agencies/PSU/AIIMS** for the following products quoted in the tender at

Signature and Stamp of Bidder

the time of submission of bid. Further, quoted drugs have not been failed in house testing or testing by any State Government/Central Government / its Drug procurement agencies during last two years. We are eligible to participate in the tender ref. No. **AIIMS-NAG/Drugs/RC/OTE/HOS/24-25/04 dated 13/06/2024** for the following quoted products with mentioned shelf life as per clause 17(ix): -

S. No.	**Drug Code (Refer Annexure- VIII)**	**Description of Drug as per AIIMS, Nagpur Tender**	**Shelf life complying the Schedule-P" of the Drugs and Cosmetics Rule, 1945.**

Signed.....

Name: ……..

Designation…….…….

(Company Seal)

Witness: -

(1) Signed: …….

Name: .....

Designation: …….

(2) Signed: …….

Name: .....

Designation: …….

ANNEXURE-III
MANDATE FORM

Sl. No.	Details Required	
1.	**Company Name**	
2.	Postal Address of the Company	
	GST No.	
	Telephone No.	
	Fax No.	
	E-mail ID	
3.	Name of the Managing Director / Director / Manager	
	Mobile No. / Phone No	
	E-mail ID	
4.	Name and Designation of the authorized company official	Name:
		Designation:
	Mobile No.	
	E-mail ID	
5.	Permanent E-mail ID	
	Permanent Mobile No.	
6.	**Bank Details**	
	a) Name of the Bank	
	b) Branch Name & address	
	c) Branch Code No.	
	d) Branch Manager Mobile No.	
	e) Branch Telephone no	
	f) Branch E-mail ID	
	g) 9-digit MICR code number of the bank and branch appearing on the MICR cheque issued by the bank	
	h) IFSC Code of the Branch	
	i) Type of Account (Current / Savings)	
	j) Account Number (as appear in cheque book)	

(In lieu of the bank certificate to be obtained, please **attach the original cancelled cheque** issued by your bank for verification of the above particulars).

I / We hereby declare that the particulars given above are correct and complete. If the transaction is delayed or not effected at all the reasons of incomplete or incorrect information, I would not hold **All India Institute of Medical sciences, Nagpur (AIIMS, Nagpur)** responsible. I have read the conditions of the tender / Rate contract entered and agree to discharge the responsibility expected of me / from the company as a tenderer / successful tenderer.

Signature and Stamp of Bidder

Date: Signature :

Name :

Designation:

Place: Company Seal (Name of the person signing & designation)

CERTIFIED THAT THE PARTICULARS FURNISHED ABOVE BY THE COMPANY ARE CORRECT AS PER OUR RECORDS.

Signature of the authorized official of the bank

Bank Seal with address:

Annexure-IV

Date:

(manufacturing License, validity of license and market standing certificate details)

S. N.	Drug Code (Only Quoted Drugs as mentioned in Annexure VIII)	Drug Specification (As per Tender Specification)	Drug Manufacturing License					Marketing standing Certificate (MSC)		
			Drug Manufacturing License No.	License Issue date	License Renewal Date	License Validity Date	Page no. of Document in uploaded Scan Copy (Do not put page nos. in range)	Market Standing Certificate Issue Date	Period of Marketing as per Marketing standing Certificate (MSC)	Page no. of Document in uploaded Scan Copy (Do not put page nos. in range)

Note:

(i) **In case any details as desired above is missing/not submitted against quoted drug, the bid for such drug are liable to be rejected.**

(ii) **It is directed to not put page nos. in range and should indicate the page nos. one by one for all respective quoted drug codes.**

Signature:

Name:

Authorized Signatory:

Seal of the Company:

Signature and Stamp of Bidder

Annexure-V

MANUFACTURER's / PRINCIPAL's AUTHORIZATION FORM

To
The Director,
All India Institute of Medical Sciences Nagpur

Dear Sir,

Tender No._____

We _____ who are established and reputable manufacture of _____ having factories at _____ and _____ hereby authorize Messrs. (Authorized Dealer/Sole Distributor/Supplier) _____
(name and address of agents) to bid, negotiate and conclude the contract with you against Tender No._____ _____ For the above goods manufactured by us. No company or firm or individual other than Messrs _____ are authorized to bid, negotiate and conclude the contract in regard to this business against this specific tender, We hereby extend our full guarantee and warranty as per the conditions of tender for the goods bided for supply against this tender by the above firm The authorization is valid up to _____

Yours faithfully,

(Name)

For and on behalfof
M/s._____
(Name of manufacturers)/Principal

Signature and Stamp of Bidder

Annexure-VI
Declaration of Local Content

Format for Affidavit of Self Certification regarding Local Content in drugs/consumables to be purchase on Rs. 100/- Stamp Paper.

I_____S/o, D/o, W/o_____ of do

_____

hereby solemnly affirm and declare as under:

That I will agree to abide by the terms and conditions of the policy of Government of India issued vide Notification No:

That the information furnished hereinafter is correct to best of my knowledge and belief and I undertake to produce relevant records before the procuring entity or any authority so nominated by the Department of Pharmaceuticals. Government of India for the purpose of assessing the local content.

That the local content for all inputs which constitute the said consumables has been verified by me and I am responsible for the correctness of the claims made therein.

That in the event of the domestic value addition of the product mentioned herein is found to be incorrect and not meeting the prescribed value-addition norms based on the assessment of an authority so nominated by the Department of Pharmaceutical. Government of India for the purpose of assessing the local content, action will be taken against me as per Order No. P-45021/2/2017-B.E-II dated 15.06.2017 and Guidelines issued vide letter no. 31026/36/2016- MD dated – 18.05.2018.

I agree to maintain the following information in the company's record for a period of 8 years and shall make this available for verification to any statutory authority.

i. Name and details of the Domestic Manufacturer (Registered Office, Manufacturing unit location, nature of legal entity).
ii. Date on which this certificate is issued.
iii. Consumable/drugs for which the certificate is produced
iv. Procuring entity to whom the certificate is furnished
v. Percentage of local content claimed **(to be calculated based on total items quoted by bidder) - ____ %**
vi. Name and contact details of the unit of the manufacturer
vii. Sale Price of the product
viii. Ex-Factory Price of the product
ix. Freight, insurance and handling
x. Total Bill of Material
xi. List and total cost value of inputs used for manufacture of the consumables.
xii. List and total cost of inputs which are domestically sourced Value addition certificates from suppliers. If the input is not in use attached.
xiii. List and cost of inputs which are imported, directly or indirectly.

For and on behalf of (Name of firm/entity)
Authorized signatory

Signature and Stamp of Bidder

Annexure-VII

(Border sharing clause)

The bidder should submit related undertaking for Restrictions on procurement from bidders from a county or countries, or a class of countries under Rule 144 (XI) of the General Financial Rules 2017 in compliance of office OM no. 6/18/2019-PPD dated 23rd July 2020. Ministry of Finance Department of Expenditure, Public Procurement Division on the basis of following Certificate given below, on the company letter head duly signed by authorized signatory for this tender.

Certificate for Tender

Tender no.:- _____

Item/drug name: - _____

'We have read the clause regarding restrictions on procurement from a bidder of a country which shares a land border with India; and solemnly certify that we are not from such a county or, if from such a country, we are registered with the Competent Authority (copy enclosed). We hereby certify that we fulfill all requirements in this regard and are eligible to be considered."

AND

We have read the clause regarding restrictions on procurement from a bidder of a county which shares a land border with India and on sub-contracting to contractors from such a country; and solemnly certify that we are not from such a county or, if from such a country, we are registered with the Competent Authority (copy enclosed) and we shall not subcontract any work to a contractor from such countries unless such contractor is registered with the Comps eat Authority. We hereby certify that we fulfil all requirement in this regard and are eligible to be considered."

It is to declare that if, our bid/offer is accepted by the purchaser, as per undertaking given by us as per aforementioned points on the basis of certificate are found to be false, in such case this would be a ground for immediate termination of our bid/offer and further legal action in accordance with the law to be initiating on us by the procuring entity i.e. AIIMS, Nagpur.

***Note:** This clause is not applicable for Indian bidders

[Signature with date, name and designation]
for and on behalf of Messrs_____

[Name & address of the manufacturers]

Annexure – VIII

(List of Drugs/medicines with Specification)

S.N	NAME OF DRUGS
1	Inj.Dexmedetomidine 100MCG/ML (1ml/Ampule)
2	5-Aminosalicylic acid Retention Enema (1 units)
3	5-Aminosalicylic acid Suppository 500 mg (1 units)
4	5-Aminosalicylic acid Tablet 400 mg (1 Tablet)
5	5-Fluorouracil Injection 250 mg/5 ml (5ml/ amp)
6	6-Mercaptopurine Tablet 50 mg (1 Tablet)
7	Abacavir (A) + Lamivudine (B) Tablet 60 mg (A) +30 mg (B) (1 Tablet)
8	Abacavir (A) + Lamivudine (B) Tablet 600mg (A) +300 mg (B) (1 Tablet)
9	Abacavir Tablet 300 mg (1 Tablet)
10	Abacavir Tablet 60 mg (1 Tablet)
11	Acetazolamide Tablet 250 mg (1 Tablet)
12	Acetretin Cap 10mg (1 Capsule)
13	Acetretin Cap 25mg (1 Capsule)
14	Acetylsalicylic acid Effervescent Tablet 300 mg (1 Tablet)
15	Acetylsalicylic acid Dispersible Tablet 300 mg (1 Tablet)
16	Acetylsalicylic acid Tablet 100mg, (1 Tablet)
17	Acetylsalicylic acid Tablet 150mg, (1 Tablet)
18	Acetylsalicylic acid Tablet 300mg, (1 Tablet)
19	Acetylsalicylic acid Tablet 75 mg, (1 Tablet)
20	Actinomycin D Powder for Injection 0.5 mg (0.5g/vial)
21	Activated charcoal Powder (as licensed) (100g bottles)
22	Acyclovir ointment 3% (5g unit)
23	Acyclovir Oral liquid 200 mg/5ml (125ml bottle)
24	Acyclovir Oral liquid 400 mg/5 ml (100ml bottle)
25	Acyclovir Powder for Injection 250 mg (250mg/vial)
26	Acyclovir Powder for Injection 500 mg (500mg/ vial)
27	Acyclovir Tablet 200 mg (1 Tablet)
28	Acyclovir Tablet 400 mg (1 Tablet)
29	Adenosine Injection 3 mg/ml (2ml Ampule)
30	Adrenaline Injection 1 mg/ml (1 ml Ampule)
31	Allopurinol Tablet 100 mg (1 Tablet)
32	Allopurinol Tablet 300 mg (1 Tablet)
33	All-trans retinoic acid Capsule 10 mg (1 Capsule)
34	All-trans retinoic acid Capsule 20mg (1 Capsule)
35	Alprostadil Injection 0.5 mg/ml (1ml/Ampule)
36	Amikacin drop 0.3% (5ml drop)
37	Amino Acid for injection 10 %
38	Amiodarone Injection 50 mg/ml (3ml Ampule)
39	Amiodarone Tablet 100 mg (1 Tablet)
40	Amiodarone Tablet 200 mg (1 Tablet)
41	Amitriptyline Tablet 10 mg (Tablet)
42	Amitriptyline Tablet 25 mg (Tablet)
43	Amlodipine Tablet 10 mg (1 Tablet)
44	Amlodipine Tablet 2.5 mg (1 Tablet)
45	Amoxicillin (A) + Clavulanic acid (B) Dry Syrup 125 mg (A) + 31.25 (B)/5 ml (30ml bottles)
46	Amoxicillin (A) + Clavulanic acid (B) Oral liquid 200 mg (A) + 28.5 mg (B)/5 ml (30 ml Bottle)
47	Amoxicillin (A) + Clavulanic acid (B) Powder for Injection 250 mg (A) + 50 mg (B)/vial

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48	Amoxicillin (A) + Clavulanic acid (B) Powder for Injection 500 mg (A) + 100 mg (B)/vial
49	Amoxicillin (A) + Clavulanic acid (B)Tablet 500 mg (A) + 125 mg (B) (1 Tablet)
50	Amoxicillin Oral liquid 250 mg/5 ml (100ml bottles)
51	Amphotericin B (conventional) 50mg/vial
52	Amphotericin B (Liposomal) 50mg/vial
53	Ampicillin Powder for Injection 1 g (1g/vial)
54	Ampicillin Powder for Injection 125 mg (125/vial)
55	Anti Rabies Immunoglobulin 150iu/ml/PF5 Vaccine
56	Anti-D immunoglobulin (1500IU/vial)
57	Anti-D immunoglobulin (300mcg/vial)
58	Anti-tetanus immunoglobulin (250IU/vial)
59	Arsenic trioxide Injection 1mg/ml (10ml/vial)
60	Artemether (A) + Lumefantrine (B) Oral liquid 80 mg (A) + 480 mg (B)/5 ml (30ml bottles)
61	Artemether (A) + Lumefantrine (B) Tablet 20 mg (A) + 120 mg (B) (1 Tablet)
62	Artemether (A) + Lumefantrine (B) Tablet 40 mg (A) + 240 mg(B) (1 Tablet)
63	Artemether (A) + Lumefantrine (B) Tablet 80 mg (A) + 480 mg (B) (1 Tablet)
64	Arterolone + piperaquine Each film coated Tablet contains; arterolane maleate eq.to arterolone-150mg, piperaquine phophate-750ml (1 Tablet)
65	Artesunate (A) + Sulphadoxine - Pyrimethamine (B) Combi pack (A+B) 1 Tablet 100 mg (A) + 1 Tablet(750mg+37.5mg) (B) (1 Tablet)
66	Artesunate (A) + Sulphadoxine - Pyrimethamine (B) Combi pack (A+B) 1 Tablet 150 mg (A) + 2 Tablet(500mg+25mg) (B) (1 Tablet)
67	Artesunate (A) + Sulphadoxine - Pyrimethamine (B) Combi pack (A+B) 1 Tablet 200 mg (A) + 2 Tablet(750mg+37.5mg) (B) (1 Tablet)
68	Artesunate (A) + Sulphadoxine - Pyrimethamine (B) Combi pack (A+B) 1 Tablet 25 mg (A) + 1 Tablet(250mg+12.5mg) (B) (1 Tablet)
69	Artesunate (A) + Sulphadoxine - Pyrimethamine (B) Combi pack (A+B) 1 Tablet 50 mg (A) + 1 Tablet(500mg+25mg) (B) (1 Tablet)
70	Artesunate Powder for Injection 120 mg (120mg/ Vial)
71	Artesunate Powder for Injection 60 mg (60mg/ Vial)
72	Atazanavir (A) + Ritonavir (B) Tablet 300 mg (A) + 100 mg (B) (1 Tablet)
73	Atorvastatin Tablet 40 mg (1 Tablet)
74	Atracurium Injection 10 mg/ml (2.5ml/Ampule)
75	Atropine Drops 1% (5 ml drops)
76	Atropine Injection 0.6 mg/ml (1 ml Ampule)
77	Atropine Injection 1 mg/ml (1 ml Ampule)
78	Atropine Ointment 1% (5g/tube)
79	Azacitidine Lyophilized powder for injcetion 100mg (1 vial)
80	Azacitidine Lyophilized powder for injcetion 50 mg (1 vial)
81	Azathioprine Tablet 50 mg (1 Tablet)
82	Azithromycin eye drops 1% (3ml drop)
83	Azithromycin Ointment 1% (5 gm/1tube)
84	Azithromycin Oral liquid 200 mg/5ml (15 ml bottle)
85	Azithromycin powder for injection 500 mg (500mg/vial)
86	Baclofen Tablet 10 mg (1 Tablet)
87	Baclofen Tablet 20 mg (1 Tablet)
88	Baclofen Tablet 5 mg (1 Tablet)
89	Barium sulphate Oral liquid 100% w/v (1lit)
90	Barium sulphate Oral liquid 250% w/v (1lit)
91	BCG vaccine (20ml/ vial)
92	Bedaquiline Tablet 100 mg (1 Tablet)
93	Bendamustine Hydrochloride Lyophilized powder for injection 100mg/vial (1vial)

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94	Benzoyl peroxide Gel 2.5% (20/30gunit)
95	Betamethasone Cream 0.05% (15gm unit)
96	Betamethasone Cream 0.1% (15gm)
97	Betamethasone Injection 4 mg/ml (1ml/Ampule)
98	Betaxolol Drops 0.25% (5ml drops)
99	Betaxolol drops 0.5% (5ml drops)
100	Bicalutamide Tablet 50 mg (1 Tablet)
101	Bimatoprost Drops, 0.03% (3ml drops)
102	Bisacodyl Suppository 5 mg (1 units)
103	Bisacodyl Tablet 5 mg (1 Tablet)
104	Bleomycin Powder for Injection 15 units (15 units/ vial)
105	Bortezomib Powder for Injection 2mg (2mg/vial)
106	Brimonidine tartarate + timolol maleate eye drop 0.15 w/v (5ml drop)
107	Brimonidine tartarate + timolol maleate eye drop 0.5 w/v (5ml drop)
108	Budesonide Respirator solution for use in nebulizer 0.5 mg/ml (2ml/respule)
109	Budesonide Respirator solution for use in nebulizer 1 mg/ml (2ml/respule)
110	Budesonide (A)+ Formoterol (B) Nasal Spray 100 mcg/dose (spray)
111	Budesonide (A)+ Formoterol (B) Nasal Spray 50 mcg/dose (spray)
112	Bupivacaine Injection 0.25% (20 ml /Vial)
113	Bupivacaine Injection 0.5% with 7.5 glucose (4 ml/ Ampule)
114	Buprenorphine (A)+ Naloxone (B) Tablet A-2 mg + B-0.5 mg (1 tab)
115	Buprenorphine transdermal patch 10 mg
116	Caffeine Injection 20 mg/ml (1ml/vial)
117	Caffeine Oral liquid 20 mg/ml (3 ml/vial)
118	Calamine Lotion (As per IP) (60/100ml unit)
119	Calcitriol oint. Each gm contains; calcitriol-0.003mg (3mcg) (20g tube)
120	Calcium carbonate suspension 500mg/ml (200ml bottle)
121	Calcium folinate Injection 3 mg/ml (5ml vial)
122	Calcium folinate Tablet 15 mg (1 Tablet)
123	Calcium gluconate Injection 100 mg/ml (10 ml Ampule)
124	Capecitabine Tablet 500 mg (1 Tablet)
125	Capreomycin Powder for Injection 1 g (1g Vial)
126	Carbamazepine Oral liquid 200 mg/5 ml (10 ml bottles)
127	Carbamazepine CR Tablet 200 mg (1 Tablet)
128	Carbamazepine CR Tablet 400 mg (1 Tablet)
129	Carbamazepine Oral liquid 100 mg/5 ml (100 ml bottle)
130	Carbamazepine Tablet 100 mg (1 Tablet)
131	Carbamazepine Tablet 200 mg (1 Tablet)
132	Carbamazepine Tablet 400 mg (1 Tablet)
133	Carbimazole Tablet 10 mg (1 Tablet)
134	Carbimazole Tablet 5 mg (1 Tablet)
135	Carboplatin Injection 10 mg/ml (10ml vial)
136	Carboprost 250mcg/ml in (1ml/Ampule)
137	Carboxymethylcellulose Drops 0.5 % (10 ml drops)
138	Carboxymethylcellulose Drops 1 % (10 ml drops)
139	Cefadroxil Oral liquid 125 mg/5 ml (60ml bottles)
140	Cefadroxil Tablet 1 g (1 Tablet)
141	Cefazolin Powder for Injection 1 g (1g vial)
142	Cefazolin Powder for Injection 500 mg (500mg vial)
143	Cefixime Oral liquid 100 mg/5 ml (60ml bottles)

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144	Cefixime Oral liquid 50 mg/5 ml (60ml bottles)
145	Cefixime Tablet 400 mg (1 Tablet)
146	Ceftriaxone Powder for injection 2 gm (1 vial)
147	Cefuroxime Powder for Injection 1.5 gm (1.5gm vial)
148	Cefuroxime Powder for Injection 750 mg (750 vial)
149	Chlorambucil Tablet 2 mg (1 Tablet)
150	Chlorambucil Tablet 5 mg (1 Tablet)
151	Chloramphenicol ointment 1 %
152	Chlorhexidine Solution 2% (Concentrate for dilution) (500 ml bottle)
153	Chlorhexidine Solution 5% (Concentrate for dilution) (500 ml bottle)
154	Chloroquine Oral liquid 50 mg/5 ml (60ml bottles)
155	Chloroquine Tablet 150 mg (1 Tablet)
156	Cholecalciferol Oral liquid 400 IU/ml (30ml bottles)
157	Cholecalciferol Tablet 1000 IU, (1 Tablet)
158	Cholecalciferol Tablet 60000 IU, (1 Tablet)
159	Ciprofloxacin drops 0.3% (2.5ml unit)
160	Ciprofloxacin Ointment 0.3% (10g unit)
161	Ciprofloxacin Oral liquid 250mg/5ml (100 ml bottle)
162	Cisplatin Injection 1mg/ml (50ml Vial)
163	Clarithromycin Oral liquid 125mg/5 ml (30ml bottle)
164	Clarithromycin Tablet 250 mg (1 Tablet)
165	Clarithromycin Tablet 500 mg (1 Tablet)
166	Clindamycin Capsule 150 mg (1 Capsule)
167	Clindamycin Capsule 300 mg (1 Capsule)
168	Clindamycin Injection 300mg/2ml
169	Clobazam Tablet 10 mg (1 Tablet)
170	Clobazam Tablet 5 mg (1 Tablet)
171	Clobetasol propionate 0.05% +Salicylic acid 3% Ointment (20g tube)
172	Clofazimine Capsule 100 mg (1 Capsule)
173	Clofazimine Capsule 50 mg (1 Capsule)
174	Clomiphene Tablet 100 mg (1 Tablet)
175	Clomiphene Tablet 50 mg (1 Tablet)
176	CLOMIPRAMINE 25MG SR (1 Tablet)
177	CLOMIPRAMINE 75MG SR (1 Tablet)
178	CLONAZEPAM 2MG (1 Tablet)
179	Clonazepam Tablet 0.5 mg (1 Tablet)
180	Clopidogrel Tablet 75mg (1 Tablet)
181	Clotrimazole Drops 1% (5ml drop)
182	Clotrimazole mouth paint 1 % (w/w)
183	Clotrimazole Pessary 100 mg (1 unit)
184	Clozapine Tablet 100 mg (1 Tablet)
185	Clozapine Tablet 25 mg (1 Tablet)
186	Clozapine Tablet 50 mg (1 Tablet)
187	Coagulation factor IX Powder for Injection 600 IU (600IU/Vial)
188	Coagulation factor VIII Powder for Injection 250 IU (250IU/vial)
189	Coagulation factor VIII Powder for Injection 500 IU (500IU/vial)
190	Coal tar Solution 5% (50ml bottle)
191	Colchicine Tablet 0.5 mg (1 Tablet)
192	Colistin Inj. 1 MU (1 mu vial)
193	Colistin Inj. 2 MU (2 mu vial)

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194	Condom As per the standards prescribed in Schedule R of Drugs and Cosmetics rules, 1945 (1 unit)
195	Co-trimoxazole [Sulphamethoxazol e (A) + Trimethoprim (B)] Tablet 400 mg (A) + 80 mg (B) (1 Tablet)
196	Cryoprecipitate As licensed (1 unit)
197	Cyclophosphamide Powder for Injection 500 mg (500mg /Vial)
198	Cyclophosphamide Tablet 200 mg (1 Tablet)
199	Cyclophosphamide Tablet 50 mg (1 Tablet)
200	Cycloserine Capsule 125 mg (1 Capsule)
201	Cycloserine Capsule 250 mg (1 Capsule)
202	Cyclosporine Capsule 25 mg (1 Capsule)
203	Cyclosporine Capsule 50 mg (1 Capsule)
204	Cyclosporine Capsule10 mg (1 Capsule)
205	Cyclosporine eye drop 0.5% (5ml drop)
206	Cyclosporine Injection 50 mg/ml (5ml/Ampule)
207	Cyclosporine Oral liquid 100 mg/ml (50ml bottles)
208	Cytosine arabinoside Injection 100 mg/ ml (1ml vial)
209	Cytosine arabinoside Powder for Injection 1000 mg (1g Vial)
210	Cytosine arabinoside Powder for Injection 500 mg (500mg/Vial)
211	Dabigatran etexilate mesylate Capsule 110 mg (1 Capsule)
212	Dabigatran etexilate mesylate Capsule 75 mg (1 Capsule)
213	Dabigatran etexilate mesylate Capsule150 mg (1 Capsule)
214	Dacarbazine ,Powder for Injection 200 mg (200mg Vial)
215	Dacarbazine Powder for Injection 500 mg (500ml Vial)
216	Daclatasvir Tablet 60 mg (1 Tablet)
217	Dapsone Tablet 100 mg (1 Tablet)
218	Dapsone Tablet 25 mg (1 Tablet)
219	Dapsone Tablet 50 mg (1 Tablet)
220	Darunavir (A) + Ritonavir (B) Tablet 400 mg (A) + 50 mg (B) (1Tablet)
221	Darunavir (A) + Ritonavir (B) Tablet 800 mg (A) + 100 mg (B) (1Tablet)
222	Darunavir Tablet 600 mg (1 Tablet)
223	Daunorubicin Injection 5 mg/ml (10ml vial)
224	Delamanid Tablet 50 mg (1 Tablet)
225	Desferrioxamine Powder for Injection 500 mg (500g/Vial)
226	Desloratadine Tablet 5mg (1 Tablet)
227	Desvenlafaxine Tablet 100 mg (1 Tablet)
228	Desvenlafaxine Tablet 50 mg (1 Tablet)
229	Dexamethasone Injection 4 mg/ml (2ml Ampule)
230	Dexamethasone Injection 4 mg/ml (5ml/vial)
231	Dexamethasone Tablet 0.5 mg (Tablet)
232	Dextran-40 Injection 10% (500 ml infusion)
233	Diazepam Injection 5 mg/ml (2ml Ampule)
234	Diazepam Oral liquid 5 mg/5ml, (100 ml bottles)
235	Diazepam Suppository 5 mg (1 unit)
236	Diazepam Tablet 2mg (1 Tablet)
237	Diazepam Tablet 5mg (1 Tablet)
238	Dicyclomine Injection 10 mg/ml (2ml Ampules)
239	Dicyclomine Oral Solution 10mg/5ml (30ml bottle)
240	Dicyclomine Tablet 10 mg (1 Tablet)
241	Diethylcarbamazine Oral liquid 120 mg/5 ml (100ml bottle)
242	Diethylcarbamazine Tablet 100 mg (1 Tablet)

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243	Diethylcarbamazine Tablet 50 mg (1 Tablet)
244	Digoxin Injection 0.25 mg/ml (2ml Ampule)
245	Digoxin Oral liquid 0.05 mg/ml (60ml bottles)
246	Digoxin Tablet 0.25 mg (1 Tablet)
247	Diloxanide furoate Tablet 500 mg (1 Tablet)
248	Diltiazem Injection 5 mg/ml (5ml Vial)
249	Diltiazem SR Tablet 90 mg (Tablet)
250	Diltiazem Tablet 30 mg, (Tablet)
251	Diltiazem Tablet 60 mg (Tablet)
252	Dimercaprol Injection 50 mg/ml (2ml Ampule)
253	Dinoprostone Gel 0.5 mg (1 Tablet)
254	Dinoprostone Tablet 0.5 mg (1 Tablet)
255	Diphtheria antitoxin (10ml/vial)
256	Divalproex sodium Tablet 250mg (1 Tablet)
257	Divalproex sodium Tablet 500mg CR(1 Tablet)
258	Dobutamine Injection 50 mg/ml (5ml Ampule)
259	Docetaxel Powder for Injection 20 mg (20mg/ Vial)
260	Docetaxel Powder for Injection 80 mg (2ml Vial)
261	Dolutegravir (A)+Lamivudine (B)+Tenofovir Disoproxil Fumarate Tablet 50 mg (A)+300mg (B)+ 300 mg (1 Tablet)
262	Dolutegravir Tablet 50 mg (1 Tablet)
263	Donepezil Tablet 10 mg (1 Tablet)
264	Donepezil Tablet 5 mg (1 Tablet)
265	Dopamine Injection 40 mg/ml (5ml Ampule)
266	Doxorubicin Injection 2 mg/ml (1ml/vial)
267	Doxycycline 50mg/ml dry syrup (50mg bottles) with sterile water
268	Doxycycline Powder for injection 100 mg/vial
269	DPT + Hib + Hep B vaccine (0.5ml Ampule)
270	DPT vaccine (10ml/vial)
271	Empaglifozin Tablet 25mg (1 Tablet)
272	Empaglifozin Tablet10mg, (1 Tablet)
273	Enalapril Tablet 2.5 mg (1 Tablet)
274	Enalapril Tablet 5 mg (1 Tablet)
275	Enoxaparin Injection 40 mg/0.4 ml (Syringe)
276	Enoxaparin Injection 60 mg/0.6 ml (Syringe)
277	Entecavir Tablet 0.5 mg (1 Tablet)
278	Entecavir Tablet 1 mg (1 Tablet)
279	Epleronone Tablet 25 mg (1 Tablet)
280	Erythropoietin Injection 10000 IU/ml (1ml syringe)
281	Erythropoietin Injection 2000 IU/ml (1ml Ampule)
282	ESCITALOPRAM 10MG (1 Tablet)
283	ESCITALOPRAM 5MG (1 Tablet)
284	Esmolol Injection 10 mg/ml (10ml Ampule)
285	Etanercept-25mg Injection (25mg/vial)
286	Ethambutol Tablet 200 mg (1 Tablet)
287	Ethambutol Tablet 400 mg (1 Tablet)
288	Ethambutol Tablet 600 mg (1 Tablet)
289	Ethambutol Tablet 800 mg (1 Tablet)
290	Ethinylestradiol Tablet 0.01 mg, (1 Tablet)
291	Ethinylestradiol Tablet 0.05 mg (1 Tablet)
292	Ethinylestradiol(A) + Levonorgestrel Tablet 0.03 mg (A) + 0.15 mg (B) (1 Tablet)

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293	Ethionamide Tablet 125 mg (1 Tablet)
294	Ethionamide Tablet 250 mg (1 Tablet)
295	Ethyl alcohol (Denatured) Solution 70% (500 ml)
296	Etoposide Capsule 100 mg (1 Capsule)
297	Etoposide Capsule 50 mg (1 Capsule)
298	Etoposide Injection 20 mg/ml (5ml vial)
299	Fentanyl Injection 50 mcg/ml (2ml Ampule)
300	Ferric Carboxy maltose Injection 500mg (10 ml vial)
301	Ferrous salt (A) + Folic acid (B) Oral liqiud 20 mg elemental iron(A) + 100 mcg (B)/ml (100ml bottle)
302	Ferrous salt (A) + Folic acid (B) Tablet 100mg elemental iron (A) + 500 mcg(B) (1 Tablet)
303	Ferrous salt (A) + Folic acid (B) Tablet 45mg elemental iron (A) + 400 mcg(B) (1 Tablet)
304	Ferrous salts Oral liquid equivalent to 25 mg of elemental iron/ml (200ml bottle)
305	Ferrous salts Oral liquid equivalent to 25 mg of elemental iron/ml (60ml bottle)
306	Ferrous salts Tablet equivalent to 60 mg of elemental iron (1 Tablet)
307	Fexofenadine Tablet 120mg (1 Tablet)
308	Fexofenadine Tablet 180mg (1 Tablet)
309	Fibrin sealant tissue glue 2ml
310	Filgrastim Injection 300 mcg (1ml vial)
311	Fluconazole Injection 200 mg /100 ml (100ml IVF) Plastic bottle Uro Head
312	Fluconazole Oral liquid 50 mg/5 ml (60ml bottles)
313	Fluconazole Tablet 100 mg (1 Tablet)
314	Fluconazole Tablet 200mg (1 Tablet)
315	Fluconazole Tablet 400 mg (1 Tablet)
316	Fludarabine Lyophilized powder for injection 50 mg/vial
317	Flunarizine Tablet 10mg (1 Tablet)
318	Flunarizine Tablet 5 mg (1 Tablet)
319	Fluorescein Eye drop 1% (15/25ml drop)
320	Fluorometholone acetate 0.1% +tobramycin Drops (5 ml drops)
321	Fluoxetine Capsule 10 mg (1 Capsule)
322	Fluoxetine Capsule 20 mg (1 Capsule)
323	Fluoxetine Capsule 20 mg (1 Capsule)
324	Fluoxetine Capsule 40 mg (1 Capsule)
325	Flurbiprofen DROPS (5ml drops)
326	Fluticasone propionate cream-0.05% (15g tube)
327	Fluticasone propionate ointment 0.005% (15g tube)
328	Folic acid Tablet 5 mg (1 Tablet)
329	Fresh frozen plasma As licensed (1 unit)
330	Fulvestrant Injection 250 mg /5ml
331	Furosemide Oral liquid 10 mg/ml (60 ml bottle)
332	Furosemide Tablet 40 mg (1 Tablet)
333	Fusidic acid cream 2 % (15 gm/tube)
334	Gadobenate Injection 529 mg/ml (5ml/vial)
335	Gadobutrol Injection 1mmol/ml
336	Gatifloxacin eye drop 5 % (5 ml drop)
337	Gefitinib Tablet 250 mg (1 Tablet)
338	Gelofusine 4 % w/v solution
339	Gemcitabine Powder for Injection 1 g (1 g/Vial)
340	Gemcitabine Powder for Injection 200 mg (200mg/vial)
341	Gentamycin 3% + dexamethasone0.1% Drops (5 ml drops)
342	Glicazide 30 mg MR (1 Tablet)

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343	Glicazide Tablet 80mg, (1 Tablet)
344	Glucose (A) + Sodium chloride (B) Injection 5% (A) + 0.9% (B) (500ml)
345	Glucose 50% 50ML GLASS BOTTLE (50ml vial)
346	Glucose Injection 10 % 500ml
347	Glucose Injection 25% (20ml/Ampule)
348	Glucose Injection 25% 100ml
349	Glucose Injection 5 % 500ml
350	Glucose Injection 50% (20ml/Ampule)
351	Glutaraldehyde Solution 2% (500 ml)
352	Glycerin Oral Liquid (50g bottle)
353	Glyceryltrinitrate Injection 5 mg/ml (10ml Ampule)
354	Glyceryltrinitrate Sublingual Tablet 0.5 mg (1 Tablet)
355	Glycopyrrolate Injection 0.2 mg/ml (1 ml Ampule)
356	Griseofulvin Tablet 125 mg (1 Tablet)
357	Griseofulvin Tablet 250 mg (1 Tablet)
358	Haemodialysis fluid As licensed (5lit can)
359	Haloperidol Injection 5 mg/ml (10ml vial)
360	Haloperidol Injection 5 mg/ml (1ml Ampule)
361	Haloperidol Tablet 1.5 mg (1 Tablet)
362	Haloperidol Tablet 5 mg (1 Tablet)
363	Heparin Injection 1000 IU/ml (5ml Vial)
364	Heparin Injection 5000 IU/ml (5ml Vial)
365	Hepatitis B immunoglobulin (100IU/ml vial)
366	Hepatitis B vaccine (10ml /vial)
367	Homatropine Drops 2% (5ml drops)
368	Hormone releasing IUD Contains 52 mg of Levonorgestrel (1 unit)
369	Human albumin Injection 20 % 100ml
370	Human chorionic gonadotropin Injection 1000 IU (1000IU/1 ml vial)
371	Human chorionic gonadotropin Injection 5000 IU
372	Human normal immunoglobulin 5% (100 ml)
373	Hydrochlorothiazide Tablet 12.5 mg (1 Tablet)
374	Hydrochlorothiazide Tablet 25 mg (1 Tablet)
375	Hydrochlorothiazide Tablet 50 mg (1 Tablet)
376	Hydrocortisone Injection 100 mg/ml (vial)
377	Hydrocortisone Tablet 10 mg (1 Tablet)
378	Hydrocortisone Tablet 5 mg (1 Tablet)
379	Hydrogen peroxide Solution 6% (500 ml bottle)
380	Hydroxocobalamin Injection 1 mg/ml (1ml Ampule)
381	HYDROXY ETHYL STARCH 6%/500ML (500ml Bottle)
382	Hydroxychloroquine Tablet 200 mg (1 Tablet)
383	Hydroxychloroquine Tablet 400 mg (1 Tablet)
384	Hydroxypropyl methylcellulose Injection 2% (2ml/Ampule)
385	Hydroxyurea Capsule 500 mg (Capsule)
386	Hydroxyurea Tablet 500 mg (1 Tablet)
387	Hyoscinebutylbromide Injection 20 mg/ml (1 ml Ampule)
388	Hyoscinebutylbromide Tablet 10 mg (1 Tablet)
389	Ibuprofen Oral liquid 100 mg/5 ml (100 ml bottles)
390	Ifosfamide Powder for Injection 1 g (1 g/Vial)
391	Ifosfamide Powder for Injection 2 g (2g/Vial)
392	Imatinib Tablet 100 mg (1 Tablet)

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393	Imatinib Tablet 400 mg (1 Tablet)
394	Imipenem + Cilastatin Injection Imipenem(500mg) + Cilastatin(500) (500mg vial)
395	INJ.BUPRENORPHINE 0.3MG/ML (2ml/Ampule)
396	Insulin (Soluble) Injection 40 IU/ml (10ml vial)
397	Insulin Glargine Injection 100 IU/ml (3ml cartrige)
398	Intermediate Acting (NPH) Insulin Injection 40 IU/ml (10ml vial)
399	Intraperitoneal dialysis solution As licensed (500ml bottle)
400	Iohexol Injection 300 mg iodine/ml (100ml bottle)
401	Iohexol Injection 350 mg iodine/ml (100ml bottle)
402	Ipratropium Inhalation Respirator solution for use in nebulizer 250 mcg/ml (250mcdg/dose)
403	Irinotecan Injection 100mg/5ml (5ml vial)
404	Iron sucrose Injection 20 mg/ml
405	Isoflurane Inhalation (250ml bottle)
406	Isoniazid Oral liquid 100 mg/5 ml (100ml bottle)
407	Isoniazid Tablet 100 mg (1 Tablet)
408	Isoniazid Tablet 300 mg (1 Tablet)
409	Isoniazid Tablet 50 mg (1 Tablet)
410	Isosorbide-5- mononitrate SR Tablet 30 mg (1 Tablet)
411	Isosorbide-5- mononitrate SR Tablet 60 mg (1 Tablet)
412	Isosorbide-5- mononitrate Tablet 10 mg (1 Tablet)
413	Isosorbide-5- mononitrate Tablet 20 mg (1 Tablet)
414	Isosorbidedinitrate Tablet 10 mg (1 Tablet)
415	Isosorbidedinitrate Tablet 5 mg (1 Tablet)
416	Ispaghula Granules/ Husk/ Powder (90g bottle)
417	Itraconazole Cap. 200mg (1 Capsule)
418	Itraconazole Cap.100mg (1 Capsule)
419	IUD containing Copper As licensed (1 unit)
420	Ivermectin Tablet 12mg (1 Tablet)
421	Ivermectin Tablet 6mg (1 Tablet)
422	Ivermectin Tablet 3mg, (1 Tablet)
423	Japanese encephalitis vaccine (20ml/ vial)
424	Kanamycin Powder for Injection 1 g (1g)
425	Kanamycin Powder for Injection 500 mg (500mg/vial)
426	Kanamycin Powder for Injection 750 mg (750mg/vial)
427	Ketamine Injection 10 mg/ml (10ml vial)
428	Ketamine Injection 50 mg/ml (10 ml vial)
429	Ketorolac tromethamine 0.40% Drops (5ml drops)
430	Labetalol 100mg (1 Tablet)
431	Labetalol 200mg (1 Tablet)
432	Labetalol Injection 5 mg/ml, (4ml/Ampule)
433	Labetalol Tablet 50mg (1 Tablet)
434	Lamivudine (A) + Zidovudine (B) Tablet 150 mg (A) + 300 mg (B) (1 Tablet)
435	Lamivudine (A) + Zidovudine (B) Tablet 30 mg (A) + 60 mg (B) (1 Tablet)
436	Lamivudine Oral solution 10mg/ml (60 ml)
437	Lamivudine Tablet 150 mg (1 Tablet)
438	Lamivudine Tablet 300 mg (1 Tablet)
439	L-Asparaginase Powder for Injection 10000 KU (10000KU/Vial)
440	L-Asparaginase Powder for Injection 5000 KU (5000KU/Vial)
441	Latanoprost drop 0.005%
442	Letrozole Tablet 2.5 mg (1 Tablet)

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443	Leuprolide acetate Injection 1 mg /0.5ml
444	Levetiracetam ER Tablet 750 mg (1 Tablet)
445	Levetiracetam Injection 100 mg/ml (5ml vial)
446	Levetiracetam Oral liquid 100 mg/ml (100ml bottles)
447	Levetiracetam Tablet 250 mg (1 Tablet)
448	Levetiracetam Tablet 500 mg (1 Tablet)
449	Levetiracetam Tablet 750 mg, (1 Tablet)
450	Levodopa (A) + Carbidopa (B) Tablet 250 mg (A) + 25 mg (B) (Tablet)
451	Levodopa (A) + Carbidopa (B) CR Tablet 100 mg (A) + 25 mg (B) (1 Tablet)
452	Levodopa (A) + Carbidopa (B) CR Tablet 200 mg (A) + 50 (B) mg (1 Tablet)
453	Levodopa (A) + Carbidopa (B) Tablet 100 mg (A) + 10 mg(B) (1 Tablet)
454	Levodopa (A) + Carbidopa (B) Tablet 100 mg (A) + 25 mg(B) (1 Tablet)
455	Levofloxacin Tablet 750 mg (1 Tablet)
456	Levonorgestrel Tablet 0.75 mg (1 Tablet)
457	Levothyroxine 12.5 mcg (1 Tablet)
458	Levothyroxine Tablet 100mcg (1 Tablet)
459	Levothyroxine Tablet 112mcg (1 Tablet)
460	Levothyroxine Tablet 125mcg (1 Tablet)
461	Levothyroxine Tablet 150mcg (1 Tablet)
462	Levothyroxine Tablet 25mcg (1 Tablet)
463	Levothyroxine Tablet 50mcg (1 Tablet)
464	Levothyroxine Tablet 62.5mcg (1 Tablet)
465	Levothyroxine Tablet 75mcg (1 Tablet)
466	Levothyroxine Tablet 88mcg (1 Tablet)
467	Lignocaine (A) + Adrenaline (B) Injection 1% (A) + 1:200000 (5 mcg/ml) (B) (30ml/ vial)
468	Lignocaine (A) + Adrenaline (B) Injection 2% (A) + 1:200000 (5 mcg/ml) (B) (30ml/ vial)
469	Lignocaine Eye drops-4% (10 ml drops)
470	Lignocaine Injection 1% (30ml/vial)
471	Lignocaine Injection 2% (30ml/vial)
472	Lignocaine Injection 5% with 7.5% Glucose (2 ml Ampule)
473	Lignocaine spray 5 %
474	Lignocaine Topical forms 2% (30g tube)
475	Linalidomide Capsule 10 mg (1 Capsule)
476	Linalidomide Capsule 15 mg (1 Capsule)
477	Linalidomide Capsule 2.5 mg (1 Capsule)
478	Linalidomide Capsule 20 mg (1 Capsule)
479	Linalidomide Capsule 25 mg (1 Capsule)
480	Linalidomide Capsule 5 mg (1 Capsule)
481	Linezolid injection 200mg/100ml
482	Linezolid Suspension 100mg/5ml
483	Linezolid Tablet 600 mg (1 Tablet)
484	Lithium Tablet 400mg SR(1 Tablet)
485	Loperamide Tablet 2 mg (1 Tablet)
486	Lopinavir (A) + Ritonavir (B) Oral liquid 400 mg (A) + 100 mg (B)/ 5ml (160ml bottles)
487	Lopinavir (A) + Ritonavir (B) Tablet 100 mg (A) + 25 mg(B) (1 Tablet)
488	Lopinavir (A) + Ritonavir (B) Tablet 200 mg (A) + 50 mg(B) (1 Tablet)
489	Lorazepam Injection 2 mg/ml (1ml Ampule)
490	Lorazepam Tablet 1 mg (1 Tablet)
491	Lorazepam Tablet 2 mg (1 Tablet)

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492	Loteprednoletabonate 0.20% Drops (5ml drops)
493	Luliconazole 1%,Each gram contains; luliconazole-10mg (15g tube)
494	Magnesium sulphate Injection 500mg/ml (2ml Ampule)
495	Mannitol Injection 10% (100ml bottle) Plastic Bottle Uro Head
496	Measles vaccine (20ml/vial)
497	Medroxyprogesterone acetate Injection ,104 mg medroxyprogesterone acetate in 0.65ml in pre filled syringe for subcutaneous injection (0.65 ml syringe)
498	Medroxyprogesterone acetate Tablet 10 mg, (1 Tablet)
499	Medroxyprogesterone acetate Tablet 5 mg (1 Tablet)
500	Mefenamic acid Capsule 250 mg (1 Capsule)
501	Mefenamic acid Capsule 500 mg (1 Capsule)
502	Mefenamic acid Oral liquid 100 mg/5 ml (60 ml bottles)
503	Mefloquine Tablet 250 mg (*Only for use as chemoprophylaxis for long term travellers like military and travel troops, travelling from low endemic to high endemic area.) (1 Tablet)
504	Meglumine diatrizoate Injection 60% w/v (20ml/Ampule)
505	Meglumine diatrizoate Injection 76% w/v (20ml/Ampule)
506	MELATONIN 3MG (1 Tablet)
507	Melphalan Tablet 2 mg (1 Tablet)
508	Melphalan Tablet 5 mg (1 Tablet)
509	Meropenam Powder for injection 125 mg (1 vial)
510	Mesna Injection 100 mg/ml (10ml/vial)
511	Mesna Injection 100 mg/ml (2ml Ampule)
512	Metformin tab 1000 mg (Immediate and controlled release) (1 Tablet)
513	Metformin Tablet 750 mg (1 Tablet)
514	Methotrexate Injection 25 mg/ ml (1 ml Ampule)
515	Methotrexate Tablet 10 mg (1 Tablet)
516	Methotrexate Tablet 2.5 mg (1 Tablet)
517	Methotrexate Tablet 5 mg (1 Tablet)
518	Methotrexate Tablet 7.5 mg (1 Tablet)
519	Methyldopa Tablet 250 mg (1 Tablet)
520	Methyldopa Tablet 500 mg (1 Tablet)
521	Methylergometrine Injection 0.2 mg/ml (1ml Ampule)
522	Methylergometrine Tablet 0.125 mg (1 Tablet)
523	Methylprednisolone Injection 40 mg/ml (40mg/vial)
524	Methylprednisolone Injection 500 mg (500mg/vial)
525	Methylprednisolone Tablet 16 mg (1 Tablet)
526	Methylprednisolone Tablet 32 mg (1 Tablet)
527	Methylprednisolone Tablet 8 mg (1 Tablet)
528	Methylrosanilinium chloride (Gentian Violet) Topial preparation 0.25% to 2% (30ml bottle)
529	Methylthioninium chloride(Methylene blue) Injection 10 mg/ml (10ml Ampule)
530	Metoclopramide Injection 5 mg/ml (2ml Ampule)
531	Metoclopramide oral liquied 5ml/ml (100ml bottles)
532	Metoclopramide Tablet 10 mg (1 Tablet)
533	Metoprolol SR Tablet 25 mg (1 Tablet)
534	Metoprolol SR Tablet 50 mg (1 Tablet)
535	Metoprolol Tablet 25 mg (1 Tablet)
536	Metoprolol Tablet 50 mg (1 Tablet)
537	Metronidazole Oral Liquid 200mg/5ml (60ml bottles)
538	Midazolam Injection 1 mg/ml (5 ml Ampule)
539	Midazolam Injection 5 mg/ml (2 ml Ampule)
540	Midazolam Oral liquid 2 mg/ml (118ml bottle)

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541	Midazolam Tablet 15 mg (1 Tablet)
542	Midazolam Tablet 7.5 mg (1 Tablet)
543	Mifepristone Tablet 200 mg (1 Tablet)
544	Milrinone lactate Injection 10mg/10 ml (10 ml vial)
545	Miltefosine Capsule 10 mg (1 Capsule)
546	Miltefosine Capsule 50 mg (1 Capsule)
547	Misoprostol Tablet 100 mcg (1 Tablet)
548	Misoprostol Tablet 200 mcg (1 Tablet)
549	Montelukast Tab 10 mg(1 Tablet)
550	Montelukast Tab 5 mg(1 Tablet)
551	Morphine 10 mg Tablet, (1 Tablet)
552	Morphine Injection 10 mg/ml (1ml Ampule)
553	Morphine Injection 15 mg/ml (1 ml Ampule)
554	Moxifloxacin Drop 0.5% (5ml drop)
555	Moxifloxacin Tablet 200 mg (1 Tablet)
556	Moxifloxacin Tablet 400 mg (1 Tablet)
557	Mupirocin ointment 2% w/v 15 gm/tube
558	Mycophenolatemofetil Tablet 250 mg (Tablet)
559	Mycophenolatemofetil Tablet 500 mg (Tablet)
560	N-acetylcysteine Injection 200 mg/ml (5ml Ampule)
561	N-acetylcysteine Sachet 200 mg (1 sachet)
562	Naloxone Injection 0.4 mg/ml (1 ml Ampule)
563	Natamycin Drops 5% (5ml drops)
564	Neostigmine Injection 0.5 mg/ml (1ml Ampule)
565	Neostigmine Tablet 15 mg (1 Tablet)
566	Nevirapine Tablet 200 mg (1 Tablet)
567	Nevirapine Dispersible Tablet 50 mg (1 Tablet)
568	Nevirapine Oral liquid 50 mg/5 ml (100ml bottle)
569	Nicotine gum 2 mg
570	Nicotine lozenges 2 mg
571	Nicotine lozenges 4 mg
572	Nicotine Transdermal patch 14mg/24hr
573	Nicotine Transdermal patch 21mg/24hr
574	Nicotine Transdermal patch 7mg/24hr
575	Nifedipine Tablet 10 mg (1 Tablet)
576	Nitazoxanide Oral suspension 100 mg/5ml (30 ml bottle)
577	Nitazoxanide Tablet 500 mg (1 Tablet)
578	Nitrofurantoin oral liquid 25mg/5 ml (100ml bottle)
579	Nitrofurantoin Tablet 100 mg (1 Tablet)
580	Noradrenaline Injection 2 mg/ml (2ml Ampule)
581	Norethisterone Tablet 5 mg (1 Tablet)
582	Ofloxacin 0.1% + dexamethasone0.1% Drops (10ml drops)
583	Olanzapine Tablet 10mg (1 Tablet)
584	Olanzapine Tablet 5mg (1 Tablet)
585	Olmesartan Tablet 20mg (1 Tablet)
586	Olmesartan Tablet 20mg (1 Tablet)
587	Olopatadine eye drop 2% (5/15ml drop)
588	Omeprazole Capsule 10 mg (1 Capsule)
589	Omeprazole Capsule 40 mg (1 Capsule)
590	Omeprazole Powder for oral liquid 20 mg (20mg sachet)

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591	Ondansetron Injection 2mg/ml (2ml Ampule)
592	Ondansetron oral liquid 2mg/5ml (100ml bottle)
593	Ondansetron Tablet 4mg (1 Tablet)
594	Ondansetron Tablet 8mg (1 Tablet)
595	Oral poliomyelitis vaccine (5ml)
596	Oxaliplatin Injection 5 mg/ml (10ml/ vial)
597	Oxygen Inhalation (Medicinal gas) (10 ltrs)
598	oxytocin Injection 10 IU/ml (1ml Ampule)
599	oxytocin Injection 5 IU/ml (1ml/Ampule)
600	Paclitaxel Injection 100 mg/16.7 ml (16.7 ml vial)
601	Paclitaxel Injection 30 mg/5 ml (5 ml Ampule)
602	Pantoprazole 40 mg Injection (vial)
603	Para- aminosalicylic acid Tablet 500 mg Granules (As licensed)
604	PARACETAMOL inj 1000MG/100ML (100 ml Infusion)
605	Paracetamol Injection 150 mg/ml (2 ml /Ampule)
606	Paracetamol Suppository 170 mg (1 unit)
607	Paracetamol Suppository 80 mg (1 unit)
608	Penicillamine Capsule 250 mg (1 Capsule)
609	Permethrin Cream 5% (30gm unit)
610	Permethrin Lotion 1% (60ml bottle)
611	PHENIRAMINE MALEATE 22.75MG/2ML INJECTION (2ml/Ampule)
612	Pheniramine maleate Tablet 25 mg (1 Tablet)
613	Pheniramine maleate Tablet 50 mg (1 Tablet)
614	Phenobarbitone Inj 100 mg/ml (2ml Ampule)
615	Phenobarbitone Inj 200mg/ml (1 ml Ampule)
616	Phenobarbitone oral liquid 20mg/5 ml (100ml bottle)
617	Phenobarbitone Tab 30mg (1 Tablet)
618	Phenobarbitone Tab 60mg (1 Tablet)
619	Phenoxy methyl Penicillin 125mg/5ml (100ml bottle)
620	Phenoxy methyl Penicillin Tablet 250 mg (1 Tablet)
621	Phenoxy methyl Penicillin Tablet 500 mg (1 Tablet)
622	Phenylephrine Drops 5% (5ml drops)
623	Phenytoin ER Tablet 300 mg (Tablet)
624	Phenytoin Injection 25 mg/ml (2ml/Ampule)
625	Phenytoin Injection 50 mg/ml (2ml/Ampule)
626	Phenytoin Oral liquid 125 mg/5 ml (200 ml bottles)
627	Phenytoin Oral liquid 30 mg/5 ml (120ml bottle)
628	Phenytoin Tablet 100 mg (1 Tablet)
629	Phenytoin Tablet 300 mg (1 Tablet)
630	Phenytoin Tablet 50 mg (1 Tablet)
631	Phynylephrine Drops 10 % (5ml drops)
632	Phytomenadione (Vitamin K1) Injection 10 mg/ml (1ml Ampule)
633	Phytomenadione (Vitamin K1) Tablet 10mg (1 Tablet)
634	Pilocarpine Drops 2% (5ml drops)
635	Pilocarpine Drops 4% (15ml drops)
636	Pioglitazone Tablet 15mg (1 Tablet)
637	Piperacillin (A) + Tazobactam (B) Powder for Injection 1 g (A) + 125 mg (B) (1.125g vial)
638	Piperacillin (A) + Tazobactam (B) Powder for Injection 2 g (A) + 250 mg (B) (2.25g vial)
639	Platelet rich plasma As licensed (1unit)
640	Podophyllin resin Solution 10% to 25% (10ml)

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641	Potassium chloride Injection 150 mg/ml (10ml/Ampule)
642	Potassium chloride Oral liquid 500 mg/5 ml (200ml bottles)
643	Potassium permanganate Crystals for topical solution (10g)
644	Pralidoxime chloride (2-PAM) Injection 25 mg/ml (10ml/Ampule)
645	Praziquantel Tablet 600 mg (1 Tablet)
646	Prednisolone Injection 20 mg/2 ml (10ml vial)
647	Prednisolone Oral liquid 15 mg/5 ml (60ml bottles)
648	Prednisolone Oral liquid 5 mg/5 ml (60ml bottles)
649	Prednisolone Tablet 10 mg (1 Tablet)
650	Prednisolone Tablet 20 mg (1 Tablet)
651	Premix Insulin 30:70 Injection (Regular:NPH) Injection 40 IU/ml (10ml vial)
652	Primaquine Tablet 15 mg (1 Tablet)
653	Primaquine Tablet 2.5 mg (1 Tablet)
654	Primaquine Tablet 7.5 mg (1 Tablet)
655	Procarbazine Capsule 50 mg (1 Capsule)
656	Proparacaine Drops 0.5% (15ml drops)
657	Propofol Injection 10 mg/ml (10 ML)
658	Propranolol Tablet 10 mg (1 Tablet)
659	Propranolol Tablet 40 mg (1 Tablet)
660	Protamine Injection 10 mg/ml (5ml Ampule)
661	Pyrazinamide Oral liquid 250 mg/5 ml (60ml bottles)
662	Pyrazinamide Tablet 1000 mg (1 Tablet)
663	Pyrazinamide Tablet 1500 mg (1 Tablet)
664	Pyrazinamide Tablet 500 mg (1 Tablet)
665	Pyrazinamide Tablet 750 mg (1 Tablet)
666	Pyridoxine Tablet 10 mg (1 Tablet)
667	Pyridoxine Tablet 100 mg (1 Tablet)
668	Pyridoxine Tablet 50 mg (1 Tablet)
669	QUETIAPINE 100MG (1 Tablet) SR
670	QUETIAPINE 25MG (1 Tablet)
671	Quinine Injection 300 mg/ml (2ml Ampule)
672	Quinine Tablet 300 mg (1 Tablet)
673	Rabies vaccine (1ml /vial)
674	Raltegravir Tablet 400 mg (1 Tablet)
675	Red blood cells As licensed (1unit)
676	Ribavirin Capsule 200 mg (1 Capsule)
677	Riboflavin Tablet 5 mg (1 Tablet)
678	Rifampicin Capsule 150 mg (1 Capsule)
679	Rifampicin Capsule 300 mg (1 Capsule)
680	Rifampicin Capsule 450 mg (1Capsule)
681	Rifampicin Capsule 600 mg (1 Capsule)
682	Rifampicin Oral liquid 100 mg/5 ml (60ml bottles)
683	RISPERIDONE 0.5mg (1 Tablet)
684	RISPERIDONE 2MG (1 Tablet)
685	Ritonavir Tablet 100 mg (1 Tablet)
686	Rituximab Injection 500 mg/50ml vial
687	Rosuvastatin Tablet 10 mg (1 Tablet)
688	Rosuvastatin Tablet 20 mg (1 Tablet)
689	Rotavirus vaccine
690	Salbutamol Oral liquid 2 mg/5 ml (60 ml bottle)

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691	Salbutamol pMDI 100mcg/puff
692	Salbutamol Respirator solution for use in nebulizer 5mg/ml (5ml bottle)
693	Salbutamol Tablet 2 mg (1 Tablet)
694	Salbutamol Tablet 4 mg (1 Tablet)
695	Salicylic acid Ointment 6% (30g/50gunit)
696	Secnidazole Oral granule 2 gm
697	Secnidazole Tablet 1g (1 Tablet)
698	Secnidazole Tablet 500 mg (1 Tablet)
699	Sertraline Tablet 100mg (1 Tablet)
700	Sertraline Tablet 50mg (1 Tablet)
701	Sevoflurane Inhalation (250 ml bottle)
702	Sildenafil Injection 10mg/12.5ml vial
703	Snake venom antiserum (Soluble/ liquid polyvalent) Injection (10 ml vial)
704	Snake venom antiserum I.P. (Lyophilized)/10ml Vial
705	Sodium bicarbonate Injection (as per IP) (25ml Ampule)
706	Sodium bicarbonate Tablet 1000 mg (1 Tablet)
707	Sodium bicarbonate Tablet 300 mg (1 Tablet)
708	Sodium bicarbonate Tablet 500 mg (1 Tablet)
709	Sodium chloride Injection 0.45% (500 ml Infusion) Plastic bottle Uro Head
710	Sodium chloride Injection 0.9% (100ml Infusion) Plastic bottle Uro Head
711	Sodium chloride Injection 0.9% (1000ml Infusion) Plastic bottle Uro Head
712	Sodium chloride Injection 0.9% (3000ml Infusion) Plastic bottle
713	Sodium chloride Injection 0.9% (500ml Infusion) glass bottle
714	Sodium chloride Injection 0.9% (500ml Infusion) Plastic bottle Uro Head
715	Sodium chloride Injection 3% (100 ml Infusion) Glass Bottle
716	Sodium chloride Injection 3% (1000ml Infusion) Plastic bottle Uro Head
717	Sodium cromoglycate 2.00% Drops (5 ml drops)
718	Sodium hypochlorite Solution 5 % (5 liter)
719	Sodium nitrite Injection 30 mg/ml (10 ml vial)
720	Sodium nitroprusside Injection 10 mg/ml (2ml/vial)
721	Sodium thiosulphate Injection 100 mg/ml (10ml vial)
722	Sodium valproate CR Tablet 300 mg (1 Tablet)
723	Sodium valproate CR Tablet 500 mg (1 Tablet)
724	Sodium valproate Injection 100 mg/ml (5ml Ampule/vial)
725	Sodium valproate Oral liquid 200 mg/5ml (100ml bottles)
726	Sodium valproate Tablet 200 mg (1 Tablet)
727	Sodium valproate Tablet 300 mg (1 Tablet)
728	Sofosbuvir Tablet 400 mg (1 Tablet)
729	Somatostatin Powder for Injection 3 mg (3g/Ampule)
730	Spironolactone Tablet 25 mg (1 Tablet)
731	Spironolactone Tablet 50 mg (1 Tablet)
732	Streptokinase Injection 15,00,000 IU (30ml vial)
733	Streptokinase Injection 750,000 IU (30 ml vial)
734	Streptomycin Powder for Injection 1 g (1g/ Vial)
735	Streptomycin Powder for Injection 750 mg (0.75g/vial)
736	Succinylcholine Injection 50 mg/ml (2ml/Ampule)
737	Sulfasalazine Tablet 500 mg (1 Tablet)
738	Sumatriptan Injection 6 mg/ 0.5 ml (0.5ml syringe)
739	Sumatriptan Tablet 25 mg (1 Tablet)
740	Sumatriptan Tablet 50 mg (1 Tablet)

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741	Surfactant Suspension for intratracheal instillation (As licensed) (4ml /vial)
742	Tacrolimus 0.1% eye ointment (3g/unit)
743	Tacrolimus Capsule 0.5 mg (1 Capsule)
744	Tacrolimus Capsule 1 mg (1 Capsule)
745	Tacrolimus Capsule 2 mg, (1 Capsule)
746	Tamoxifen Tablet 10 mg (1 Tablet)
747	Telmisartan Tablet 20 mg (1 Tablet)
748	Telmisartan Tablet 40 mg (1 Tablewt)
749	Temozolomide Capsule 100 mg (1 Capsule)
750	Temozolomide Capsule 20 mg (1 Capsule)
751	Tenecteplase Lyophilized powder for injection 40 mg/vial
752	Teneligliptin Tablet 20 mg (1 Tablet)
753	Tenofovir (A) + Lamivudine (B) + Efavirenz Tablet 300 mg (A) + 300 mg (B) + 600 mg (C) (1 Tablet)
754	Tenofovir (A) + Lamivudine (B) Tablet 300 mg (A) +300 mg (B) (1 Tablet)
755	Tenofovir Alafenamide Fumarate Tablet 25 mg (1 Tablet)
756	Tetanus toxoid (10ml/vial)
757	Thalidomid Capsule 100 mg (1 Capsule)
758	Thalidomid Capsule 50 mg (1 Capsule)
759	Thiamine Injection 100 mg/ml (2ml/Ampule)
760	Thiamine Tablet 100 mg (1 Tablet)
761	Thiopentone Powder for Injection 1 g (1 g/ vial)
762	Thiopentone Powder for Injection 0.5 g (0.5g/ vial)
763	Ticagrelor Tablet 90mg (1 Tablet)
764	Timolol Drops 0.25% (5 ml drops)
765	Timolol Drops 0.5% (5 ml drops)
766	Tiotropium Inhalation (DPI) 18 mcg/dose (18mcg/dose)
767	Tiotropium Inhalation (MDI) 9 mcg/dose (9mcg/dose)
768	Tobramycin eye drop 0.3% (5ml drop)
769	Torsemide Tablet 10 mg (1 Tablet)
770	Tramadol Capsule 50mg (1 Capsule)
771	Tramadol Injection 50 mg/ml (2 ml Ampule)
772	Tranexamic acid Injection 100 mg/ml (5ml Ampule)
773	Tranexamic acid Tablet 500 mg (1 Tablet)
774	Trastuzumab Lyophilized powder for injection 440mg/vial
775	Triamcinolone acetonide injection 40mg/ml
776	Trihexyphenidyl Tablet 2 mg (1 Tablet)
777	Tropicamide Eye drop 1% (5/15ml drop)
778	Tuberculin, Purified Protein derivative (5 ml/vial)
779	Valacyclovir Tablet 1 gm (1 Tablet)
780	Valacyclovir Tablet 500 mg (1 Tablet)
781	Valganciclovir Oral solution 50mg/ml (100ml bottle)
782	Valganciclovir Tablet 450mg (1 Tablet)
783	Vancomycin powder for inj 1 g (1g vial)
784	Vancomycin powder for inj 250mg (250mg vial)
785	Vancomycin powder for inj 500mg (500mg vial)
786	Vecuronium Powder for Injection 10 mg (10mg/vial)
787	Vecuronium Powder for Injection 4 mg (4mg/vial)
788	Verapamil Injection 2.5 mg/ml (2ml Vial)

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789	Verapamil Tablet 40 mg (1 Tablet)
790	Verapamil Tablet 80 mg (1 Tablet)
791	Vildagliptin Tablet 50mg (1 Tablet)
792	Vinblastine Injection 1 mg/ml (10ml Ampule)
793	Vincristine Injection 1 mg/ml (1ml Ampule)
794	Vitamin A Capsule 100000 IU (1 Capsule)
795	Vitamin A Capsule 5000 IU (1 Capsule)
796	Vitamin A Capsule 50000 IU (1 Capsule)
797	Vitamin A Injection 50000 IU/ml (2ml vial)
798	Vitamin A Oral liquid 100000 IU/ml (100ml bottles)
799	Voglibose Tab 0.2 mg (1 Tablet)
800	Voglibose Tab 0.3 mg (1 Tablet)
801	Voriconazole powder for eye drop 1 %
802	Warfarin Tablet 3 mg (1 Tablet)
803	Warfarin Tablet 1mg (1 Tablet)
804	Warfarin Tablet 2 mg (1 Tablet)
805	Warfarin Tablet 5 mg (1 Tablet)
806	Water for Injection Injection (5ml Ampule)
807	White Petrolatum Jelly 100% (50g bottle)
808	Whole blood As licensed (1 unit)
809	Xylometazoline nasal drops 0.05 % (10ml)
810	Xylometazoline nasal drops 0.1 % (10ml)
811	Zidovudine (A) + Lamivudine (B)+ Nevirapine Tablet 60 mg (A) + 30 mg (B) + 50 mg (C) (1 Tablet)
812	Zidovudine (A) + Lamivudine (B)+ Nevirapine Tablet 300 mg (A) + 150 mg (B) + 200 mg (1 Tablet)
813	Zidovudine Oral liquid 50 mg/5 ml (60 ml bottle)
814	Zidovudine Tablet 300 mg (1 Tablet)
815	Zinc sulphate Dispersible Tablet 20 mg (1 Tablet)
816	Zinc sulphate Drops 20 mg/mL
817	Zinc Sulphate Syrup 20 mg/100 mL
818	Zoledronic acid Powder for Injection 4 mg (4mg/vial)
819	Zolpidem Tablet 10 mg (1 Tablet)
820	Zolpidem Tablet 5 mg (1 Tablet)
821	Zuclopenthixol acetate Injection 50mg/ml

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PARTICULARS FOR PERFORMANCE GUARANTEE BOND

(To be typed on Non-judicial stamp paper of the value of Indian Rupees of Two Hundred)
(TO BE ESTABLISHED THROUGH ANY OF THE SCHEDULED BANK (WHETHER SITUATED AT NAGPUR OR OUTSTATION) WITH A CLAUSE TO ENFORCE THE SAME ON THEIR LOCAL BRANCH AT NAGPUR)

To,
The Director
All India Institute of Medical Sciences (AIIMS),
Nagpur-441108

LETTER OF GUARANTEE

WHERE AS All India Institute of Medical Sciences (AIIMS) Nagpur (Buyer) have invited Tenders vide Tender No.....Dt.....for purchase of..... .....AND WHERE AS the said tender document requires the supplier/firm(seller)whose tender is accepted for the supply of consumables etc. in response there to shall establish an irrevocable Performance Guarantee Bond in favour of "The Director, AIIMS Nagpur" in the form of Bank Guarantee for Rs.....[3% (Three percent) of the purchase value] which will be valid beyond 60 days of completion of warranty period from the date of supply, installation & commissioning, the said Performance Guarantee Bond is to be submitted within 30 (Thirty) days from the date of Acceptance of the Purchase Order.

NOW THIS BANKHERE BY GUARANTEES that in the event of the said supplier/firm (seller) failing to abide by any of the conditions referred to intender document/purchase order/performance/quality of the Injector Syringe, instrument/machinery, etc. This Bank shall pay to All India Institute of Medical Sciences (AIIMS) Nagpur on demand and without protest or demur..... (Rupees.).

This Bank further agrees that the decision of All India Institute of Medical Sciences (AIIMS) Nagpur (Buyer) as to whether the said supplier/firm (Seller) has committed a breach of any of the conditions referred in tender document/ purchase order shall be final and binding.

We,.....(name of the Bank& branch) here by further agree that the Guarantee herein contained shall not be affected by any change in the constitution of the supplier/firm (Seller)and/or All India Institute of Medical Sciences (AIIMS) Nagpur (Buyer).

Not with standing anything contained herein:

a. Our liability under this Bank Guarantee shall not exceed`.....(Indian Rupees.....only).

b. This Bank Guarantee shall be valid up-to.....(date) and date of claim should be beyond six months from the date of validity.

c. We are liable to pay the guaranteed amount or any part thereof under this bank guarantee only and only if AIIMS Nagpur serve upon us a written claim or demand on or before..... (Date). This should be beyond six months from validity as (b) above.

This Bank further agrees that the claims if any, against this Bank Guarantee shall be enforceable at our branch office at.....situated at.....(Address of local branch).

Yours truly,

Signature and seal of the Guarantor

Name of the Bank:…….

Complete Postal Address: .....

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Instructions for Online Bid Submission

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.

REGISTRATION

1) Bidders are required to enroll 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 Enrollment" on the CPP Portal which is free of charge.
2) As part of the enrolment process, the bidders will be required to choose a unique username and assign a password for their accounts.
3) 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.
4) 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.
5) 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.
6) 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.

SEARCHING FOR TENDER DOCUMENTS

1) 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.
2) 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.
3) 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.

PREPARATION OF BIDS

1) Bidder should take into account any corrigendum published on the tender document before submitting their bids.
2) Please go through the tender advertisement and the tender document carefully to

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understand the documents required to be submitted as part of the bid. Please note the

3) 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.

4) Bidder, in advance, should get ready the bid documents to be submitted as indicated in the tender document / schedule and generally, they can be in PDF / XLS / RAR / DWF/JPG formats. Bid documents may be scanned with 100 dpi with black and white option which helps in reducing size of the scanned document.

5) 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 (e.g. PAN card copy, annual reports, auditor certificates etc.) 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.

SUBMISSION OF BIDS

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.

2) The bidder has to digitally sign and upload the required bid documents one by one as indicated in the tender document.

3) 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.

4) 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.

5) The documents being submitted by the bidders would be encrypted using PKI encryption all 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 these cured 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.

6) Further this key is subjected to asymmetric encryption using buyers/bid opener's public keys. Overall, the uploaded tender documents become readable only after the tender opening by the authorized bid openers.

7) The uploaded tender documents become readable only after the tender opening by the authorized bid openers.

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8) Upon the successful and timely submission of bids (ie 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.
9) 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.

ASSISTANCE TO BIDDERS

1) Any queries relating to the tender 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 tender.
2) 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 number 0120-4200462,0120-4001002.

…….

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