

ALL INDIA INSTITUTE OF MEDICAL SCIENCES, RAIPUR (C.G.)

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**NIT FOR THE ANNUAL RATE CONTRACT CUM SUPPLY AND
EMPANELMENT FOR SUPPLY OF DRUGS AND MEDICINES FOR THE
YEAR 2013-14**



आरोग्यम् सुख सम्पदा

LAST DATE OF SUBMISSION OF BIDS:- 05.12.2013 UP-TO 3:00 P.M.

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Tender Enquiry No. ADMIN/Tender/DRUGS/1/2013

Date: 02.11.2013

Notice Inviting Tender

BIDS are invited upto 3.00 PM of 05.12.2013 for the Annual Rate Contract cum Supply and **Empanelment** for supply of **drug and medicines** for the year 2013-14. Details may be seen in the Bidding Documents at our office or at the website of State Public procurement Portal <http://www.aiimsraipur.edu.in> and may be downloaded from there.

**Director
AIIMS, Raipur**

ALL INDIA INSTITUTE OF MEDICAL SCIENCES, RAIPUR (C.G.)

**BID FOR THE ANNUAL RATE CONTRACT CUM SUPPLY AND
EMPANELMENT FOR SUPPLY OF DRUGS AND MEDICINES FOR THE
YEAR 2013-14**

Bid Reference	Tender Enquiry No. ADMIN/Tender/DRUGS/1/2013 Dated: 02.11.2013
Pre-bid conference	21.11.2013 at 11:00 AM (AIIMS, Raipur Meeting Hall)
Last date and time of submission of bids	05.12.2013 up-to 3:00 PM
Date and time of opening of technical bids	05.12.2013 at 3:30 PM
Cost of the Bid document	₹ 2000/- (Central State PSU, NSIC Listed SSI are Exempted from Cost)
Empanelment Fee (If applicable for Empanelment also)	₹ 5000/-

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GENERAL INSTRUCTION FOR BIDDERS

The bidders are instructed to read the complete bid document carefully. The following points may be noted so that mistakes/lapses/shortcomings during Bid submission may be avoided.

1. The turnover should be as per bid conditions. Do not submit Bid if the turnover of the firm is less.
2. Do not quote the products manufactured on Loan license basis.
3. Quote only for the products for which your Product Permission meets the Bid specifications. Do not quote if it differs with regard to any parameter.
4. Quote only for those products for which the bidder has Market Standing Certificates is there for last three years. (Three years means:- 3 x 365 days).
5. Quote rate in BOQ for the packing exactly given in annexure VII. For example if the packing is given for 10x10 tablets, the rate should be quoted for 10x10 tablets, and not for 1 tablet or 10 tablets, similarly if the packing unit in the Bid specifies 2 ml ampoule (10 ampoules), the rate should be for 10 ampoules and not for 1 ampoule or 25 ampoules.
6. Highlight the quoted items in the documents like Product Permission and Market Standing Certificate, and also mark the item code no. at appropriate place in the documents.
7. In case there is any suggestion regarding Bid conditions/specifications/shelf life, strength, packing/turn over etc. The suggestions should be submitted/sent/ E – Mailed one/two days earlier from the date of Pre-bid meeting so that the representation of the bidders may be well processed and decision could be taken well in time.
8. If there is any query in Bid document, you may contact

Deputy Director Administrator

AIIMS, Raipur

Tel No.- 07712573222, 2573555

E-mail:dda@aiimsraipur.edu.in

ALL INDIA INSTITUTE OF MEDICAL SCIENCES, RAIPUR (C.G.)

**BID FOR THE ANNUAL RATE CONTRACT CUM SUPPLY AND
EMPANELMENT FOR SUPPLY OF DRUGS AND MEDICINES FOR THE
YEAR 2013-14**

AIIMS, Raipur (hereinafter referred as Bids Inviting Authority unless the context otherwise requires) TENDER FOR THE ANNUAL RATE CONTRACT CUM SUPPLY AND EMPANELMENT FOR SUPPLY OF DRUGS AND MEDICINES FOR THE YEAR 2013-14

1. LAST DATE FOR RECEIPT OF BIDS, BID FEES, EMD AND EMPANELMENT FEES.

- (a) Bids [in three separate bids (Technical bid, Price Bid, Prequalification Bid)] will be received till 05.12.2013 up-to 3.00 PM by the All India Institute of Medical Sciences, Raipur, for the annual rate contract cum supply and empanelment for supply of drugs and medicines.
- (b) The bids shall be valid for a Period of 120 days from the date of opening of Technical Bid and prior to the expiration of the bid validity the Bid Inviting Authority may request the Bidders to extend the bid validity for another period of 30 days. The Bidder may refuse extension of bid validity without forfeiting the Earnest Money deposit.
- (c) The Bids will be received at AIIMS, Tatibandh GE Road Raipur. Every Bidder will be required to pay, EMD as applicable in Bid condition no. 8. The shall submit all the required documents Bank Draft/ DD/Bank Guarantee on Date: 05.12.2013 before 3:00 PM.
- (d) **Those who wish to apply for Empanelment as supplier for Drugs and Medicines** are required to deposit separately an Empanelment Fee of ₹ 5000 (Five Thousand rupees only) in the form of DD in favour of AIIMS, Raipur before due time and date of bid submission. Please see clause 21 and Annexure-IX in this regard.

2. ELIGIBILITY CRITERIA

- (a) Bidder shall be a manufacturer having valid own manufacturing license or direct importer holding valid import license. Distributors/ Suppliers / Agents/Loan licensee are not eligible to participate in the Bids.
- (b) Average Annual turnover (for drugs and medicines including Surgical

and sutures Business) in the last three financial years (2010-11, 2011-12, 2012-13 [as mentioned in clause no 5 (l) (m)] shall not be less than ₹ 10 Crores.

(c) Bidder should have permission to manufacture the item /drug quoted as per specification given in the Bid, from the competent authority. Product permission of brands shall be accepted in the Bid submitted, but the Bidder has to submit the product permission in generic names at the time of signing of the agreement/before supply.

(d) Bid should not be submitted for the product/products for which the concern/company stands blacklisted/banned/debarred either by Bid inviting Authority or any Govt. state organization/PSU or Autonomous body of Govt. of India.

The Bid should not be submitted for those products also for which the concern/company stands blacklisted/banned/debarred by any other State/Central Govt. or **it's any agencies** (central Drugs procurement agencies) on the ground of **conviction by court of law or the products being found spurious or adulterated on the ground of submission of fake or forged documents or false information /facts**

(e) The concern/company/firm which stands blacklisted/banned/debarred on any ground either by Bid Inviting Authority/ Govt. state organization/PSU or Autonomous body of Govt. of India or its department on the date of bid submission, shall not be eligible to participate in the Bid.

(f) The concern/company/firm which stands blacklisted/banned/debarred on any ground either by Bid Inviting Authority/ Govt. state organization/PSU or Autonomous body of Govt. of India or its department on the date of bid submission, shall not be eligible to participate in the Bid.

The concern/company/firm which stands blacklisted/banned/debarred on the ground of **conviction by court of law or the products being found spurious or adulterated** by any other State /Central Government or **it's any agencies** (central Drugs procurement agencies) shall also not be eligible to participate in the Bid.

(g) If any product/products of a company/firm have been declared as not of standard quality, as per Drugs & Cosmetics Act during last 2 years anywhere, such concern/company/firm shall not be eligible to participate in Bid for such product/products. If any company/firm is

found to have any such product quoted in the Bid, the product shall be blacklisted for 2 years and a penalty equivalent to EMD shall also be levied. In such situation, the bid will be considered further only if the amount of penalty is deposited before the completion of technical evaluation.

- (h) The concern/firm/company whose product has been declared as of spurious or adulterated quality and any criminal case is filed and pending in any court shall not be eligible to participate for that particular product, in the Bid. Similarly convicted firm/company shall also not be eligible to participate in the Bid.
- (i) If a company has two or more separate manufacturing units at different sites/states, the company will be allowed to submit only one Bid for all units but necessary document regarding separate manufacturing units will be submitted as a separate set with the same Bid. But a bidder will be allowed to submit only one offer for one product.

3. PURCHASE PREFERENCE

- i. Purchase preference admissible to the Central Govt./State Govt. PSUs, shall not exceed 15%. However these units will be required to participate in Bidding process and match L-1 price. **Decision of Director in this regard will be final.**

4. GENERAL CONDITIONS

- I. At any time prior to the date of submission of Bid, Bid Inviting Authority may, for any reason, whether on his own initiatives or in response to a clarification requested by a prospective Bidder, modify the condition in Bid documents by amendment. In order to provide reasonable time to take the amendment into account in preparing their bid, Bid Inviting Authority may at his discretion, extended the date and time for submission of Bids.
- II. Interested eligible Bidders may obtain further information in this regard from the office of the Bid Inviting Authority.
- III. In case any document submitted by the bidder or his authorized representative is found to be forged, false or fabricated, the bid will be rejected and EMD/SD will be forfeited. Bidder/his representative may also be blacklisted/banned/debarred. Report with police station may also be filed against such bidder/his representative.

5. TECHNICAL BID

The Bidder should furnish the following in technical bid:-

- (a) Bidders are allowed the option to quote for anyone item or more items as mentioned in tender (list of medicines proposed to be purchased at Annexure-VII). The amount of EMD will remain @ ₹ 10,000/- per item of drug quoted subject to minimum of ₹1.00 lacs and maximum of ₹10.00 lacs.
- (b) The required EMD / Tender fees may be in form of physical D.D./ BC/BG/FDR shall be in favour of AIIMS, Raipur.
- (c) Those who wish to apply for Empanelment as supplier for Drugs and Medicines are required to deposit separately an Empanelment Fee of ₹ 5000 (Five Thousand rupees only) in the form of DD in favour of AIIMS, Raipur before due time and date of bid submission.
- (d) Documentary evidence for the constitution of the company/Firm such as Memorandum and Articles of Association, Partnership Deed etc. with details of the Name, Address, Telephone Number, Fax Number, e-mail address of the firm and of the Director /Partners/Proprietor.
- (e) The Bidder should furnish attested copy of the valid License for the product duly approved by the Licensing authority for each and every product quoted as per specification in the Bid. The license must have been duly renewed/ valid up to date and the items quoted shall be clearly highlighted (**Bid item codes marked against each item**) in the license.
- (f) Attested photocopy of the valid import license in Form 10 with Form 41 (as per Rule 122A of Drugs and Cosmetics Act), if the product is imported. The license must have been renewed /valid up to date. A copy of a valid license for the sale of Drugs imported by the firms issued by the licensing authority shall be enclosed.
- (g) The instruments such as power of attorney, resolution of board etc., authorizing an officer of the Bidder should be enclosed.
- (h) Authorization/nominating a responsible person of the Bidder to transact the business with the Bid Inviting Authority **with photograph in Annexure VI**.
- (i) Market Standing Certificate issued by the Licensing Authority/ competent authority as a Manufacturer for the product for last 3 years (Certificate should be enclosed with list of items) should be enclosed. Items quoted should be highlighted in the market standing certificate. For imported items, the quoted item should have 3 years market

standing, for which bills of entry, sale invoices, etc should be submitted to establish the claim. The importing firm should have 3 years standing as importer / manufacturer of medicines in general. The manufacturer may submit his licence or MSC to establish 3 years standing; the importer firm may submit Bills of entry, etc of same or other drugs to establish 3 years for importing the items and to establish the market standing of the firm. The bidder shall submit valid import licence for import of the quoted item. The market standing of products containing Paracetamol 500 mg shall be accepted. However, the firm shall submit the product permission of the product as per the tender specification.

- (j) Non-conviction Certificate issued by the Drugs Controller of the State. It should be recent and not more than one year old.
- (k) Good manufacturing practices Certificate (GMP) as per revised Schedule -'M', or WHO-GMP Certificate issued by the Licensing Authority. The GMP certificate must not be older than one year from the due date of Bid submission in the case where validity is not mentioned in the certificate. The Bidder shall also furnish an undertaking in the format given in Annexure-VI point no.8 declaring that the Bidder complies with the requirements of GMP (as per revised Schedule-'M'). The Importer should produce WHO- GMP /COPP of the manufacturing firm or a certificate which is at par with WHO-GMP issued by exporting countries like US- FDA approval, etc. In the case of imported drugs, labels and product literature of all quoted products must be submitted.
- (l) Annual turnover statement for 3 years i.e., 2010-11, 2011-12 and 2012-13 in the format given in Annexure-II certified by the practicing Chartered Accountant. Provisional / Audited (by CA) Turnover, of financial year 2013-14 may be accepted but the firm has to submit audited turn over statement before execution of agreement. If the firm fails to produce audited turn over statement, it will be liable for action as applicable in the case of non execution of agreement.
- (m) Copies of the Balance Sheet and Profit and Loss Account for three years i.e. 2010-11, 2011-12 and 2012-13 duly certified by the practicing Chartered Accountant. Provisional / Audited (by CA) P&L , Balance sheet of financial year 2013-14 may be accepted but the firm has to submit audited final accounts before execution of agreement. If the firm fails to produce audited final accounts, it will be liable for action as applicable in the case of non execution of agreement.

- (n) VAT/Sales Tax Clearance certificate (copies of latest challans), as on 31.03.2013.
- (o) Registration with Excise Department, Govt. of India. The industries situated in excise free zones will be exempted from the registration provided they produce the copy of appropriate notification.
- (p) Undertaking **(as in Annexure-VI)** for embossment of logo on labels of bottles, etc as the case may be, as per conditions specified at Clause 15 herein.
- (q) Undertaking that the manufacturer has not been blacklisted, the product has not been declared as not of standard quality during last two years, it's manufacturing capacity and other details required on a format mentioned at Annexure-VI.
- (r) Details of technical personnel employed in the manufacture and testing of drugs (Employee Name, Qualification, and Experience) as enclosed in license.
- (s) List of items quoted to be shown in the **Annexure-VI** point number 6.
- (t) A **Checklist (Annexure-IV)** for the list of documents enclosed with their page number. The documents should be serially arranged as per **Annexure-IV**. Every bidder will also be required to submit details of product permission of the quoted item and the desired market standing, **in Annexure- V**.
- (u) An undertaking that the bidder complies with all the terms, conditions, amendments (if any) of bid document to be submitted in **Annexure-VI point no.11**.
- (v) All copies submitted should be attested and notarized. However, scanned copies of original documents will be accepted which obviously need be notary attested.
- (w) An undertaking in Annexure-IX that the bidder wishes to get empanelled as supplier for the quoted items and has submitted the necessary fee for the same. (This is only for those who apply for empanelment also).
- (x) A copy of PAN issued by Income Tax Department.

6. PRICE BID -

The price bid will also be known as financial document and every bidder will be required to submit its price in excel format attached to the bid document.

(BOQ). BOQ template must not be modified/ replaced by the bidder. and the same should be submitted after filling the relevant columns, else the bidder is liable to be rejected for this bid. Bidders are allowed to enter the bidder name and values only. The bidder should quote rate for the

mentioned packing unit only.

7. OPENING OF TECHNICAL AND FINANCIAL EVALUATION

The Bid will be scrutinized by Bid evaluation committee and inspection of manufacturing unit for compliance of GMP may be carried out by technical committee. Price Bid (BOQ) of the **Bidder** found eligible on satisfying the criteria for technical evaluation and inspection, will only be opened.

8. EARNEST MONEY DEPOSIT

The Earnest Money Deposit shall be @ ₹ 10,000/- for each item of Drugs & Medicines quoted subject to minimum of ₹ 1.00 lacs and maximum of ₹ 10.00 Lacs. In case Earnest money submitted by the bidder is at the minimum or more but number of quoted items is more than the earnest money submitted, the quoted items by the bidder will be counted in sequence up to the earnest money deposited. However without minimum earnest money the offer will not be considered at all. In case Earnest money submitted by the bidder is at the minimum or more but number of quoted items is more than the earnest money submitted, the quoted items by the bidder will be counted in sequence up to the number matching the earnest money deposited. However without minimum earnest money the offer will not be considered at all.

EMD will not be taken/Exempted from the organization who are registered with central purchase organization (CPO) state PSU or National Small Industry Corporation (NSIC).

The Bids submitted without sufficient EMD will be summarily rejected. The EMD will be forfeited, if the Bidder withdraws its Bid after last time & date fixed for receiving bids or in the case of a successful Bidder, if the Bidder fails within specified time to sign the contract agreement or fails to furnish the security deposit.

9. OTHER CONDITIONS

1. The orders will be placed by the Director or any officer designated by Director, All India Institute of Medical Sciences, Raipur, (hereinafter referred to as Ordering Authority).
2. The details of the required drugs, medicines, etc., are shown in **Annexure-VII**. The quantity mentioned is only the tentative requirement and may increase or decrease as per the decision of Ordering Authority. The rates quoted should not vary with the quantum of the order or the destination. **The commitment quantity for an item submitted by the bidder (in Annexure VI) shall be taken in to account and a bidder not having adequate capacity (as**

reflected in commitment quantity) may be technically disqualified.

3. Bid has been called for in the **generic names of drugs**. The Bidders should quote the rates for the generic products. The composition and strength of each product should be as per details given in **Annexure-VII**. Any variation, if found, will result in rejection of the Bid. The products should conform to the specified standards IP/BP/USP. In case the product is not included in the said compendium, the supplier, upon award of the contract, must provide the reference standards and testing protocols for quality control testing.
4. Rates (**inclusive of transportation, insurance, Packing and any incidental charges, Excise Duty, Customs duty & other statutory duties of the govt. except VAT and CST**) should be quoted for each of the required drugs, medicines etc., separately on door delivery basis according to the unit ordered. Bid for the supply of drugs, medicines, etc. with conditions like "AT CURRENT MARKET RATES" shall not be accepted. Handling, clearing, transport charges etc., will not be paid. The delivery should be made as stipulated in the purchase order placed with successful Bidders. No quantity or cash discount should be offered.
5.
 - a) To ensure sustained supply without any interruption, the Bid Inviting Authority reserves the right to fix more than one supplier to supply the requirement among the qualified BidderS.
 - b) **Orders will be placed periodically during rate contract period based on the stock positions only. Orders will be placed with L1 firms. However in order to ensure regular supply in case of any exigency at the discretion of the Bid Inviting Authority, the orders may also be placed with the other firms, in the ascending order, L-2, L-3 and so on who have matched the L1 rates.**
 - c) After the conclusion of Price Bid opening, the lowest offer of the Bidder, if required will be considered for negotiations, and rate arrived after negotiations will be L-1 rate and L-1 supplier for an item of drugs/medicines for which the Bid has been invited.
 - d) The Bidder who has been declared as L-1 supplier for certain item or items of drugs/medicines shall execute necessary agreement for the supply of the Bided quantity of such drugs/medicines as specified in the Bid document on depositing

the required amount as performance security and on execution of the agreement, such Bidder is eligible for the placement of purchaser orders.

- e) AIIMS, Raipur will inform the L1 rate to the Bidders who qualified for Price Bid opening, through AIIMS, Raipur web site or e-mail; willing bidders may inform in writing their consent to match with the L-1 rate for the item of the Drugs/Medicines quoted by them and the Bidders who agree to match L1 rate, will be considered as Matched L1.
- f) The Bidder, who agrees to match L-1 rate shall furnish the breakup detail of price (L-1 rate) as mentioned in BOQ & Price Bid **Annexure – VII**.
- g) The supplier upon receipt of the purchase order deems that the purchase orders exceeds the production capacity declared in the Bid documents and the delay would occur in executing the order, shall inform the Director, AIIMS Raipur immediately without loss of time and the purchase orders shall be returned within 7 days from the date of the order, failing which the supplier is estopped from disputing the imposition of liquidated damages, fine for the delayed supply.
- h) If the L1 supplier has failed to supply /intimated Director, AIIMS Raipur about his inability/delay in supply as per the purchase order, the required Drugs/ Medicines within the stipulated time or as the case may be, Director, AIIMS Raipur may also place purchase orders with the Matched L1 Bidder for purchase of the Drugs/Medicines, provided such matched L1 Bidders shall execute necessary agreement indicating the production capacity as specified in the Bid document on depositing the required amount. Such Bidder is eligible for the placement of purchase orders for the item or items of Drugs/Medicines quoted by them.
- i) Subject to Para (h) above, while Director, AIIMS Raipur has chosen to place purchase orders with Matched L1 supplier and there are more than one such matched L1 suppliers, then the purchase orders for the requirement of Drugs/Medicines will be placed with L-2 first on matched rates of L-1 and in case L-2 does not have the required capacity than L-3 would be considered on matched L-1 rates and the same order would be followed in case of L-3, L-4 etc.

- j) The matched L1 supplier, on placement of purchase orders, will be deemed as L-1 rate supplier for the purpose of the Bid and all provisions of the Bid document applicable to L-1 rate Bidder will apply mutatis mutandis to the matched L1 supplier.
6. The rates quoted and accepted will be binding on the Bidder during validity period of the bid and any increase in the price (except increase due to Excise Duty or any all other statutory taxes) will not be entertained.
 7. 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 after last date fixed for receipt of bid. Representation to make correction in the Bid 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 Bids. 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 Bids of those who have given such conditions shall be treated as incomplete and accordingly the Bid will be rejected.
 8. The rates should be quoted only for the composition stated in the Bid.
 9. Supplies should be made directly by the bidder and not through any other agency.
 10. The Bidder shall allow inspection of the factory at any time by a team of Experts/Officials of the Bid Inviting Authority and or the team designated by Director, AIIMS Raipur. The Bidder shall extend all facilities to the team to enable to inspect the manufacturing process, quality control measures adopted etc., in the manufacture of the items quoted. If a Company/Firm does not allow for any such inspection, its Bids will be rejected.

10. ACCEPTANCE OF BID

1. The Bid evaluation committee formed by Director, AIIMS Raipur will evaluate the Bid with reference to various criteria.
2. Bid Inviting Authority reserves the right to accept or reject the Bid for the supply of all or any one or more items of the drugs Bided for in a Bid without assigning any reason.
3. Bid Inviting Authority, or his authorized representative (s) has the right to inspect the factories of Bidders, before, accepting the rate quoted by them, or before releasing any purchase order(s), or at any

point of time during the continuance of Bid and also has the right to reject the Bid or terminate/cancel the purchase orders issued and or not to reorder, based on adverse reports brought out during such inspections.

4. The acceptance of the Bids will be communicated to the successful Bidders in writing by the Bid inviting authority. Immediately after receipt of acceptance letter, the successful Bidder will be required to deposit security deposit and agreement but not later than 15 days.
5. The approved rates of the successful Bidders would be valid for one year as Annual Rate contract, May extendable by 3 months with mutual consent.

11. The Tender Process

The tender process will be of 4 cover system, consisting:

- **Cover – A: EMD, Tender- Fee & Prequalification documents.**
- **Cover – B: Technical Bid**
- **Cover – C: Price Bid**
- **Cover – D: Big cover in which all three (A, B & C) covers are placed.**

- 1) Every cover should mention the name of cover like A/B/C/D.**
- 2) It should also mention its content like EMD/Technical BID/Price BID.**
- 3) Top cover should also mention the Tender for which submission is done.
“ Tender Enquiry No. ADMIN/Tender/DRUGS/1/2013”**

12. SECURITY DEPOSIT (PERFORMANCE SECURITY)

The Successful Bidders shall be required to pay performance Security Deposit @ 10% of the Contract value.

The performance security shall have an upper limit of ₹ 40 Lac to be deposited by a bidder at the time of signing of agreement (For one or many items). However when the actual purchase orders cross a threshold for requiring additional security (for a period of 24 month), the same will be required to be deposited by the supplier.

The performance guarantee should be paid upfront in respect of each contract on or before the due date fixed by Bid inviting authority in the form of Bank Guarantee/DD/FD (**Performa given in Annexure XII**) in favor of the Director, AIIMS Raipur Payable at Raipur, viz. Bid inviting authority before releasing the purchase order by the ordering authority. In case L-2, L-3 and so on, bidders who have agreed to match L-1 price, then the EMD of L-2, L-3 and so on bidders will be converted (**₹ 20000/- per item**) into security

deposit. In case of inability of L-1 bidder to supply the required quantity of drugs, in that case the L-2 and L-3 supplier (as the case may be) will be asked to supply the drugs. At the time of placing of order these matched suppliers will be asked to deposit amount of balance security for a period of 24 month.

13. AGREEMENT

- a) The successful Bidder shall execute an agreement on a non-judicial stamp paper of value mentioned in the Acceptance Letter (stamp duty to be paid by the Bidder) within 15 days from the date of the intimation letter of interest by the Bid Inviting Authority, viz., the **Director, AIIMS, Raipur**. The Specimen form of agreement is available in **Annexure-III, failing to submission of performance security and execution of agreement within 15 days as stipulated, will result in forfeiture of EMD & other consequential action.**
- b) The Bidder shall not, at any time, assign, sub-let or make over the contract or the benefit therefore or any part thereof to any person or persons whatsoever.
- c) All notices or communication relating to, or arising out of this agreement or any of the terms thereof shall be considered duly served on or given to the Bidder if delivered to him or left at the premises, places of business or abode.

14. SUPPLY CONDITIONS

1. Purchase orders along with the delivery destinations will be placed on the successful Bidder at the discretion of the Ordering Authority. Drugs and Medicines will be supplied at AIIMS, Raipur.
2. The supplier shall supply the entire ordered quantity before the end of 60 days from the date of issue of purchase order at the destinations mentioned in the purchase order, if the above day happened to be a holiday for AIIMS, Raipur the supply should be completed by 3.00 p.m. on the next working day. For drug items requiring sterility test and imported ones, the supply period will be 75 days from the date of issue of purchase order.
3. All supplies will be scheduled for the period from the date of purchase order till the completion of the tender in installments, as may be stipulated in the purchase order.
4. **Shelf Life: The shelf life of drugs supplied should be not less than 24 months except in those Drugs/Medicines/Consumable where**

self life is recommended lesser than 24 months as per Drugs & Consumable Act. 1940. Only those bidders shall quote who can manufacture and supply the product with the required shelf life. The product of labeled shelf life lesser than required shelf life will not be accepted.

Quality Assurance: The supplier shall guarantee that the products as packed for shipment.

- (a) Comply with all provisions of specifications and related documents.
 - (b) Meet the recognized standards for safety, efficacy and quality.
 - (c) Are fit for the purpose made.
 - (d) Are free from defects in workmanship and in materials and.
 - (e) The product has been manufactured as per cGMP included in Schedule M of Drugs & Cosmetic Rules.
5. The protocol of the tests should include the requirements given in I.P for tablets and those required specifically for the product specifications. The Bidder must submit its Test/ Analysis Report for every batch of drug along with invoice. In case of failure on the part of the supplier to furnish such report, the batch of drugs will be returned back to the supplier and he is bound to replenish the same with approved laboratory test report. The supplier shall provide the validation data of the analytical procedure used for assaying the components and shall provide the protocols of the tests applied.
 6. The Drugs and medicines supplied by the successful Bidder shall be of the best quality and shall comply with the specification, stipulations and conditions specified in the Bid documents.
 7. If supplies are not fully completed in 60 days from the date of the Purchase Order (75 days for drugs of the category of serum, vaccine, enzymes, blood grouping reagents, biological products, powder for injections and imported drugs), the provisions of liquidated damages of Tender conditions will come into force. The Supplier should supply the drugs at the AIIMS, Raipur Pharmacy Store.
 8. If the supplier fails to execute at least 50% of the quantity mentioned in single purchase order and such part supply continues for three consecutive Purchase orders, then the supplier will be ineligible to participate in any of the Bids for particular items of drugs/medicines for a period of one year immediately succeeding year in which supplier has been placed Purchase order.
 9. If the Bidder fails to execute the supply within the stipulated time, the

ordering authority is at liberty to make alternative 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 Ordering Authority/Bid inviting authority has every right to recover the cost and impose penalty as mentioned in Clause 20, apart from terminating the contract for the default.

10. The order stands cancelled after the expiration of delivery period, and if the extension is not granted with or without liquidated damages. Apart from risk/alternate purchase action, the Bidder shall also suffer forfeiture of the performance security and shall invite other penal action like blacklisting/Debaring disqualification from participating in present and future Bids of Bid Inviting Authority/ordering authority. . (As per guidelines for blacklisting/ debaring at annexure- VIII including amendment).
11. It shall be the responsibility of the supplier for any shortage/damage at the time of receipt at the designated places.
12. If at any time the Bidder has, in the opinion of the ordering authority, delayed in making any supply by reasons of any riots, mutinies, wars, fire, storm, tempest or other exceptional cause on a specific request made by the Bidder within 7 days from the date of such incident, the time for making supply may be extended by the ordering authority at its discretion for such period as may be considered reasonable. The exceptional causes do not include the scarcity of raw material, Power cut, labour disputes.
13. The supplier shall not be in any way interested in or concerned directly or indirectly with, any of the officers, subordinates or servants of the Bid Inviting Authority in any trade or business or transactions nor shall the supplier give or pay promise to give or pay any such officers, subordinates or servants directly or indirectly any money or fee or other considerations under designation of "Customs" or otherwise, nor shall the supplier permit any person or persons whomsoever to interfere in the management or performance hereof under the power of attorney or otherwise without the prior consent in writing of the Bidder Inviting Authority.

15. LOGOGRAMS / Markings

Logogram means, wherever the context occurs, the design as specified below:-

DESIGNS FOR LOGORAMS

INJECTIONS

Injection in ampoule form should be supplied in Double constructed neck ampoules with the lable bearing the words “**AIIMS, Raipur (C.G.) Supply- Not for Sale** निःशुल्क वितरण हेतु **QC - Passed**” overprinted and the following logogram which will distinguish from the normal trade packing. Name of drug should be printed in English and Hindi languages and should be legible and be printed more prominently. Storage directions should be clear, legible, preferably with yellow highlighted background.



The vials should be supplied with aluminum seals containing the following logogram:



LIQUIDS

Liquid preparations should be in bottles with pilfer-proof caps bearing the following logogram:



आरोग्यम् सुख सम्पदा

The top of the cap and the label to be affixed on the containers should bear a distinct colour different from the colour of the label of the trade packs and they should be overprinted in red colour with the words “**AIIMS, Raipur (C.G.) Supply- Not for Sale निःशुल्क वितरण हेतु QC – Passed**” and the logogram. Name of drug should be printed in English and Hindi languages and should be legible and be printed more prominently. Storage directions should be clear, legible, preferably with yellow highlighted background.



आरोग्यम् सुख सम्पदा

OINTMENTS & CREAMS

Ointments & Creams should be supplied in tubes bearing the following logograms and the words “**AIIMS, Raipur (C.G.) Supply- Not for Sale निःशुल्क वितरण हेतु QC – Passed**” overprinted. Name of drug should be printed in English and Hindi languages and should be legible and be printed more prominently. Storage directions should be clear, legible, preferably with yellow highlighted background.

TABLETS & CAPSULES

Tablets and Capsules should be supplied in Strips or Blisters or as mentioned in the list of items for tender. The strip, etc, should bear the following logograms and the words “**AIIMS, Raipur (C.G.) Supply- Not for Sale निःशुल्क वितरण हेतु QC – Passed**” overprinted.

Name of drug should be printed in English and Hindi languages and should be legible and be printed more prominently. Storage directions should be clear, legible, preferably with yellow highlighted background.



SPECIMEN LABEL FOR OUTER CARTON

SHALL BE OF DIFFERENT COLOURS FOR DIFFERENT CLASS OF DRUGS

<p>AIIMS, Raipur (C.G.) SUPPLY NOT FOR SALE</p> <p>(Name of Drugs etc.)</p> <p>CONSTITUENTS OF.....</p> <p>Name of the Drug,</p> <p>Manufactured by,</p> <p>Batch no.</p> <p>Mfg.Date,</p> <p>Exp. Date,</p> <p>Quantity/Kit</p> <p>Net. Weight:.....Kg</p>

The name of the drug shall be mentioned in Hindi and English and should be legible and be printed more prominently. **A uniform colour theme and artwork will be necessary.** Apart from this “**For AIIMS, Raipur (C.G.) – Not for Sale निःशुल्क वितरण हेतु QC – Passed**” along with logo of AIIMS, Raipur will be printed on each strip/label of the bottle. The storage directions should be clear, legible and preferably with yellow highlighted background.

1. Bids for the supply for Drugs and medicines etc., shall be considered only if the Bidder gives undertaking in his Bid that the supply will be prepared and packed with the logogram printed on the strips of tablets and capsules and labels of bottles, ampoules and vials etc., as per the design mentioned above.
2. All tablets and capsules have to be supplied in standard packing in aluminum strip or blisters with aluminium foil back with printed logogram and shall also conform to schedule P1 of the Drugs & Cosmetics Act & Rules wherever it applies. Affixing of stickers and rubber stamps shall not be accepted.
3. Labels of Vials, Ampoules and Bottles containing the items Bided for should also carry the logogram.
4. Failure to supply Drugs etc., with the logogram will be treated as breach of the terms of agreement and liquidated damages will be deducted from bills payable as per conditions in Clause 19.2 Bidders who are not willing to agree to conditions above will be summarily rejected.
5. In case of imported drugs affixing rubber stamp on the original label is allowed with indelible ink on inner most and outer packing.

16. PACKING

The item shall be supplied in the package schedule given below and the package shall carry the logogram specified in clause -15. The labeling of

1. The item shall be supplied in the package schedule given below and the package shall carry the logogram specified in clause-15. The labeling of different packages should be as specified below. The packing in each carton shall be strictly as per the specification mentioned. Failure to comply with this shall lead to non-acceptance of the goods besides imposition of penalties.
2. The pediatric drops should always be supplied with dropper. A measuring cap with suitable markings must be provided for other paediatric oral liquid preparations.

3. The labels in the case of injectables should clearly indicate whether the preparations are meant for IV, IM, SC, etc.
4. Injection vials should have flip off seals.
5. All plastic containers should be made of virgin grade plastic.
6. The name of the drug should be printed in clearly legible bold letters (It is advisable that the colour of font be different from other printed matter to make the name highly conspicuous.
7. It should be ensured that only first hand fresh packaging material of uniform size is used for packing. All packaging must be properly sealed and temper proof.
8. All packing containers should strictly conform to the specifications prescribed in the relevant pharmacopoeia/Act.
9. Packing should be able to prevent damages or deterioration during transit.
10. In the event of items supplied found to be not as per specifications in respect of their packing, the Ordering Authority is at liberty to make alternative purchase of the item for which the purchase orders have been placed from any other sources or from the open market or from any other Bidder who might have quoted higher rates at the risk and the cost of the supplier. In such cases the ordering authority has every right to recover the cost and impose penalty as mentioned in Clause 19.2 and 20.

**I. SCHEDULE FOR PACKAGING OF DRUGS AND MEDICINES
GENERAL SPECIFICATIONS**

No corrugate package should weigh over 15 kgs (i.e. product + inner carton + corrugated box).

All items should be packed only in first hand strong boxes only.

Every corrugated box should preferably be of single joint and not more than two joints.

Every box should be stitched using pairs of metal pins with an interval of two inches between each pair.

The flaps should uniform meet but should not overlap each other. The flap when turned by 45-60 should not crack.

Every box should be sealed with gum tape running along the top and lower opening.

CARRY STRAP:

Every box should be strapped with two parallel nylon carry straps (they should intersect.)

LABEL:

Every corrugated box should carry a large outer label clearly indicating that the product is for “**AIIMS, Raipur (C.G.)** Supply-Not for Sale”.

The Product label on the cartoon should be large, atleast 15 cms x 10 cms dimension. It should carry the correct technical name, strength or the product, date of manufacturing, date of expiry quantity packed and net weight of the box.

OTHERS:

NO box should contain mixed products or mixed batches of the same product.

II. SPECIFICATION FOR CORRUGATED BOXES HOLDING TABLETS/CAPSULES/PESSARIES

1. The total weight of the box should be approx of 7-8 Kgs.

III. SPECIFICATION FOR LARGE VOLUME BOTTLE i.e., ABOVE 100 ml AND BELOW 1 LIT.

1) All these bottles should be packed only in single row with partition between each and also with top and bottom pad of 3 ply.

IV. SPECIFICATION FOR IV FLUIDS

Each corrugated box may carry maximum of only 24 bottles of 500 ml in a single row or 50 bottles of 100 ml in 2 rows with individual sealed polythene cover and centre partition pad, top and bottom pads of 3 ply.

V. SPECIFICATION FOR LIQUID ORALS

100 bottles of 50 ml or 60 ml may be packed in a single corrugated in 2 rows with top, bottom and centre pad of 3 ply.

50 bottles of 100 ml – 120 ml may be packed in a similar manner in a single corrugated box.

If the bottles are not packed in individual carton, 3 ply partition should be provided between each bottle. The measuring device should be packed individually.

VI. SPECIFICATION FOR OINTMENT/CREAM/GELS PACKED IN TUBES

No corrugated box should weigh more than 7-8 Kg.

Every Ointment tube should be individually packed in cartoon and then packed in 20's in a grey board box] which may be packed in a corrugated box.

VII. SPECIFICATIONS FOR INJECTION (IN VIALS AND AMPOULES)

Vials may be packed in corrugated boxes weighing upto 15 Kgs. Ampoules should be packed in C.B weighing not more than 8 Kgs.

In the case of 10 ml Ampoules or 50 ampoules may be packed in a grey board box. Multiples of grey board boxes packed in CB. In case of ampoules larger than 10 ml only 25 ampoules may be packed in a grey board box with partition. If the vial is packed in individual cartoon, there is no necessity for grey board box packing. The individual cartoon may be packed as such in the CB with centre pad.

In case of ampoules every grey board box should carry 5 amps alongwith Cutters placed in a polythene bag.

Vials of eye and ear drops should be packed in a individual cartoon with a dispensing device. If the vial is of FFS/BFS technology, they should be packed in 50's in a grey board box.

VIII. SPECIFICATION FOR ORS

The sachets should be of Aluminium Foil laminated with glassin or heat sealable plastic film, Outer paper may contain label information.

50 sachets may be packed in grey board boxes and 10 grey board boxes in a C.B.

IX. LYSOL

Not more than four 5 liters cans may be packed in a single Box.

17. QUALITY TESTING

1. Sampling of supplies from each batch will be done at the point of supply or distribution/storage points for testing. (The samples would be sent to different empanelled laboratories for testing by the ordering authority after coding). The AIIMS, Raipur will deduct a sum of 1.5% from the amount of bill payable to supplier on account of handling and testing charges.
2. The Drugs shall have the active ingredients within the permissible level throughout the shelf life period of the drug. The samples may also be drawn periodically during the shelf life period. The supplies will be deemed to be completed only upon receipt of the quality certificates from the laboratories. Samples which do not meet quality requirements 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.
3. In the event of the samples of the Drugs and medicines supplied failing

quality tests or found to be not as per specification the ordering authority is at liberty to make alternative purchase of items of drugs and medicines for which the Purchase orders have been placed from any other sources or from 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 ordering authority has every right to recover the cost and impose penalty as mentioned in Clause 20.

4. The supplier shall furnish to the purchaser the evidence of bio-availability and/or bio-equivalence for certain critical drugs when asked for. If there is any problem in the field the B.M.R/B.P.R for the particular batch shall also be supplied when demanded.
5. The products should conform to the standards of IP/BP / USP as the case may be. In case the product is not included in the said compendium, the supplier, upon award of the contract, must provide the reference standards and testing protocols for quality control testing. For imported drugs respective countries pharmacopeia standards shall be acceptable (even if the product is official in IP).

18. PAYMENT PROVISIONS

1. No advance payment towards costs of drugs, medicines etc., will be made to the Bidder.
2. On receipt of the prescribed consolidated invoice duly stamped and signed by authorized signatory and analytical laboratory report regarding quality, the payment would be made in 30 days. (Annexure-XI & XII).
3. The in charge of pharmacy store AIIMS, Raipur will acknowledge the drugs received.
4. All bills/ Invoices should be raised in duplicate and in the case of excisable Drugs and Medicines; the bills should be drawn as per Central Excise Rules in the name of the authority as may be designated. The supplier will deliver following document at the time of delivery at AIIMS, Raipur.
 - a. In house test report of drug.
 - b. The challan / invoice copy pertaining to AIIMS, Raipur.
5. Payments for supply will be considered only after supply of 70% of items of Drugs ordered in the Purchase Order .However, the payment will be released only for the quantity in case of which the quality test report from approved test laboratories of State of Govt. of India has been received and found of standard quality.
6. If at any time during the period of contract, the price of Bided items is

reduced or brought down by any law or Act of the Central or State Government or by the Bidder himself, the Bidder shall be bound to inform ordering authority immediately about it. Ordering authority empowered to unilaterally effect such reduction as is necessary in rates in case the Bidder fails to notify or fails to agree for such reduction of rates.

7.

(a) In case of any enhancement in Excise Duty due to notification of the Government after the date of submission of Bids and during the Bid period, the quantum of additional excise duty so levied will be allowed to be charged extra as a separate item without any change in the basic of the price structure price of the Drugs approved under the Bid. For claiming the additional cost on account of the increase in Excise Duty, the Bidder should produce a letter from the concerned Excise authorities for having paid additional Excise Duty on the goods supplied to ordering authority and also must claim the same in the invoice separately.

Similarly if there is any reduction in the rate of essential drug, as notified by the Govt., after the date of submission of Bid, the quantum of the price to the extent of reduction of essential drug will be deducted without any change in the basic price of the price structure of the drugs approved under the Bid.

(b) In case of successful bidder has been enjoying excise duty exemption on any criteria of Turnover etc., such bidder will not be allowed to claim excise duty at later point of time, during the tenure of contract, when the excise duty is chargeable on goods manufactured

8.

- i. If the supplier requires an extension in time for completion of contractual supply, on account of occurrence of any hindrance he shall apply in writing for extension on occurrence of hindrance but not after the stipulated date of completion of supply.
- ii. The purchase Officer may extend the delivery period with or without liquidated damages in case they are satisfied that the delay in the supply of goods is on account of hindrances. Reasons shall be recorded.
- iii. **Extension in delivery period:-** In case of extension in the

delivery period with liquidated damages the recovery shall be made on the basis of following percentages of value of stores which the Bidder has failed to supply:-

- a) Delay upto one fourth period of the prescribed delivery period; 2.5%
- b) Delay exceeding one fourth but not exceeding half of the prescribed delivery period; 5%.
- c) Delay exceeding half but not exceeding three fourth of the prescribed delivery period; 7.5%.
- d) Delay exceeding three fourth of the prescribed delivery period. 10%.

Note:-Fraction of a day in reckoning period of delay in supplies shall be eliminated if it is less than half a day. The maximum amount of liquidated damages shall be 10%.

9. If, at any time during the continuance of this Agreement, the Supplier has, in the opinion of the Purchaser, delayed in making any supply ordered, by the reasons of any riots, mutinies, wars, fire, storm, tempest or other exceptional cause, on a specific request made by the Supplier, the time for effecting delivery may be extended by the Purchaser surely at his discretion for such period as may be considered reasonable by the Purchaser. No further representation from the Supplier will be entertained on this account.

19. DEDUCTION IN PAYMENTS:

1. If the supply is received in damaged conditions it shall not be accepted.
2. All the Bidder are required to supply the product with logogram and with prescribed packing specification. If there is any deviation in these Bid conditions a separate damages will be levied @ 2% irrespective of the ordering authority having actually suffered any damage/loss or not, without prejudice the rights of alternative purchase specified in Clause No.16.10.

20. QUALITY CONTROL DEDUCTION & OTHER PENALTIES

1. If the successful Bidder fails to execute the agreement and/or to deposit the required performance security within the time specified or withdraws his Bid after the intimation of the acceptance of his Bid has been sent to him or owing to any other reasons, he is unable to

- undertake the contract, his contract will be cancelled and the Earnest Money Deposit deposited by him along with his Bid, shall stand forfeited by the Bid Inviting Authority and he will also be liable for all damages sustained by the Bid Inviting Authority apart from **blacklisting/ debarring the supplier**. (As per guidelines for blacklisting/ debarring at annexure VIII including amendment).
2. If the samples drawn from supplies do not conform to statutory standards, the supplier will be liable for relevant action under the existing laws and the entire stock in such batch should be taken back by the supplier within a period of 30 days from the issue of letter from ordering authority the information of which may be communicated. The stock shall be taken back at the expense of the supplier. Ordering authority has the right to destroy such NOT OF STANDARD DRUGS IF THE SUPPLIER does not take back the goods within the stipulated time. Ordering authority will arrange to destroy the NOT OF STANDARD drugs within 90 days after the expiry of 30 days mentioned above, without further notice, and shall also collect demurrage charge calculated @ 2% per week on the value of the drugs rejected till such destruction.
 3. The supplier 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 ordering authority is entitled to deduct the cost of such batch of drugs from the any amount payable to the Bidder. On the basis of nature of failure, the product/supplier will be moved for Black Listing. (As per guidelines for blacklisting/ debarring at annexure VIII including amendment).
 4. For supply of drugs of NOT OF STANDARD QUALITY the respective Drugs Controller will be informed for initiating necessary action on the supplier and that the report of product shall be sent to the committee for appropriate action including blacklisting. (As per guidelines for blacklisting/ debarring at annexure VIII including amendment).
 5. The decision of the ordering authority or any Officer authorized by him as to the quality of the supplied drugs, medicines etc., shall be final and binding.
 6. Ordering 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.
 7. For infringement of the stipulations of the contract or for other

justifiable reasons, the contract may be terminated by the ordering authority, and the supplier shall be liable for all losses sustained by the ordering authority, in consequence of the termination which may be recovered personally from the supplier or from his properties, as per rules.

8. Non performance of any contract provisions shall be examine and may disqualify the firm to participate in the future Bids.
9. In the event of making ALTERNATIVE PURCHASE, as specified in Clause **14.10**, **Clause 16.10** and in **Clause 17.3** the penalty will be imposed on supplier apart from forfeiture of Security Deposit. The excess expenditure over and above contracted process incurred by the ordering authority in making such purchases from any other sources or from the open market or from any other Bidder who has quoted higher rates and other losses sustained in the process, shall be recovered from the performance security 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.
10. In all the above conditions, the decision of the Bid Inviting Authority, viz Director, AIIMS, Raipur would be final and binding; in case of any dispute regarding all cases under Bid procedure or in any other non-ordinary situation and would be acceptable to all.
11. All litigations related to the supplier for any defaults will be done by Bid Inviting Authority and his decision will be final and binding.

21. EMPANELMENT OF FIRMS

AIIMS, Raipur invites Applications from eligible firms for Empanelment for supply of Drugs & Medicines mentioned in Annexure- VI for one year. The empanelment would entitle a firm to participate in AIIMS, Raipur for limited tenders. Such situations may normally arise when the open tender for a Surgical & Sutures fails and there is an urgency to purchase it, or when the L-1 bidder has fail to supply, or the rate contract of an item ceases to exist for any reason. The Bidder has to submit an undertaking in the format given at Annexure-IX.

The empanelment can be renewed for the next one year term on payment of the empanelment fee as applicable at the time of renewal.

22. SAVING CLAUSE

No suit, prosecution or any legal proceedings shall lie against Bid Inviting Authority or any person for anything that is done in good faith or intended to

be done in pursuance of Bid.

23. JURISDICTION

In the event of any dispute arising out of the Bid or orders such dispute would be subject to the jurisdiction of the Courts of Raipur or Honorable High Court Bilaspur Only.

24. CORRECTION OF ARITHMETIC ERRORS:

Provided that a financial bid is substantially responsive, the procuring Entity will correct arithmetical errors during evaluation of Financial Bids on the following basis:

- i.** If there is a discrepancy between the unit price and the total price that is obtained by multiplying the unit price and quantity, the unit price shall prevail and the total price shall be corrected, unless in the opinion of the Procuring Entity there is an obvious misplacement of the decimal point in the unit price, in which case the total price as quoted shall govern and the unit price shall be corrected.
- ii.** If there is an error in a total corresponding to the addition or subtraction of subtotals, the subtotals shall prevail and the total shall be corrected; and.
- iii.** If there is a discrepancy between words and figures, the amount in words shall prevail, unless the amount expressed in words is related to an arithmetic error, in which case the amount in figures shall prevail subject to clause (a) and (b) above.

If the Bidder that submitted the lowest evaluated bid does not accept the correction of errors, its Bid shall be disqualified and its Bid Security shall be forfeited or its Bid Securing Declaration shall be executed.

25. PROCURING ENTITY'S RIGHT TO VARY QUANTITY:

- i.** At the time of award of contract, the quantity of Drugs, originally specified in the bidding documents may be increased or decreased. There will not be any minimum quantity guaranteed against bid quantity. The tender quantity is only indicative. Actual purchase can be more or less than the bid quantity based on actual

consumption in the hospitals during Rate Contract period. The supplier shall submit the supply commitment quantity” in Annexure VI at point no. 3 which will be used for the cases where the actual purchase quantity tends to increase substantially from the bid quantity.

- ii. If the procuring entity does not procure any subject matter of procurement or procures less than the quantity specified in the bidding documents due to change in circumstances, the bidder shall not be entitled for any claim or compensation except otherwise provided in the conditions of contract.
- iii. However a bidder is bound to supply up to quantity indicated in bid document, considering the total production capacity & capacity dedicated to AIIMS, Raipur. Moreover, the actual purchases beyond Bid quantity may be made keeping in view the supply commitment of bidder to corporation.

26. DIVIDING QUANTITIES AMONG MORE THAN ONE BIDDER AT (IN CASE OF PROCUREMENT OF GOODS):

As a general rule all the quantities of the subject matter of procurement shall be procured from the bidder, whose bid is accepted and declared successful L-1 bidder. However, when the quantity of drugs the subject matter of procurement is very large may not be in the capacity of the bidder, whose bid is accepted to deliver the entire quantity of such drugs or when it is considered that the drugs being of critical and vital nature, whose bid are accepted and the second lowest bidder in that order.

27. COMPLIANCE WITH THE CODE OF INTEGRITY AND NO CONFLICT OF INTEREST:

Any person participating in a procurement process shall-

Not offer any bribe, reward or gift or any material benefit either directly or indirectly in exchange for an unfair advantage in procurement process or to otherwise influence the procurement process.

Not misrepresent or omit misleads or attempts to mislead so as to obtain a financial or other benefit or avoid an obligation:

Not indulge in any collusion, Bid rigging or any-competitive behaviour to impair the transparency, fairness and progress of the procurement process:

Not misuse any information shared between the procuring Entity and the Bidders with an intent to gain unfair advantage in the procurement process:

Not indulge in any coercion including impairing or harming or threatening to do the same, directly or indirectly, to any part or to its property to influence the procurement process:

Not obstruct any investigation or audit of a procurement process:

Disclose conflict of interest, if any; and

Disclose any previous transgressions with any Entity in India or any other country during the last three years or any debarment by any other procuring entity.

Conflict of interest:

The Bidder participating in a bidding process must not have a Conflict of Interest

A Conflict of interest is considered to be a situation in which a party has interests that could improperly influence that party's performance of official duties or responsibilities, contractual obligations, or compliance with applicable laws and regulations

A Bidder may be considered to be in Conflict of interest with one or more parties in bidding process if, including but not limited to:

- (a) Have controlling partners/shareholders in common; **or**
- (b) Receive or have received any direct or indirect subsidy from any of them; **or**
- (c) Have the same legal representative for purposes of the Bid; **or**
- (d) Have a relationship with each other, directly or through common third parties, that puts them in a position to have access to information about or influence on the Bid of another Bidder, or influence the decisions of the Procuring Entity regarding the bidding process; **or**
- (e) The Bidder participates in more than one Bid in a bidding process. Participation by a Bidder in more than one Bid will result in the disqualification of all Bids in which the Bidder is involved. However, this does not limit the inclusion of the same subcontractor, not otherwise participating as a Bidder, in more than one Bid; **or**
- (f) The Bidder or any of its affiliates participated as a consultant in the

preparation of the design or technical specification of the Goods, Works or Services that are the subject of the Bid; **or**

(g) Bidder or any of its affiliates has been hired (or is proposed to be hired by the Procuring Entity as engineer-in-charge/ consultant for the contract.

28. FALL CLAUSE

The prices charged for the store supplies under the contract by successful bidder shall in no event exceeded the lowest price at which the successful bidder sells the stores of identical description to any other persons during the period of the contract, the bidder reduces the sales price chargeable under the contract, he shall forth with notify such reduction to the Director, AIIMS, Raipur and the price payable under the contract of the stores supplied after the date of coming into force of such reduction or sale shall stand correspondingly reduced.

**Director
AIIMS, Raipur**

**ANNEXURE-I
Ref. Clause No.8**

**Format of Affidavit
(On Non Judicial Stamp Paper of ₹ 10/-)**

I.....S/o.....Aged.....Yrs.....

residing at.....Proprietor/Partner/Director of
M/s.....do hereby solemnly affirm and declare that:

(a) My/Our above noted enterprises M/s..... has been issued acknowledgement of Entrepreneurial Memorandum Part-II by the Districts Industries Center.....The acknowledgement No. is.....dated.....and has issued for Manufacture of following items.

- (i)
- (ii)
- (iii)
- (iv)
- (v)

(b) My/Our above noted acknowledgement of Entrepreneurial Memorandum Part-II has not been cancelled or withdrawn by the Industries Department and that the enterprise is regularly manufacturing the above items.

(c) My/Our enterprise is having all the requisite plant and machinery and is fully equipped to manufacture the above noted items.

Place.....

**Signature of Proprietor/Director
Authorized Signatory with Rubber
Stamp and date**

VERIFICATION

I.....S/o.....Aged.....Yrs.....

residing

at.....Proprietor/Partner/Director of

M/s..... verify and confirm that the contents at (a), (b) & (c) above are true and correct to the best of my knowledge and nothing has been concealed therein. So help me God.

DEPONENT

**ANNEXURE-II
Ref. Clause No. 5 (m)**

ANNUAL TURN OVER STATEMENT

The Annual Turnover (*for drugs and medicines including Surgical and sutures Business*) of M/s._____ for the past three years are

given below and certified that the statement is true and correct.

S.No.	Years	Turnover in Lakhs (₹)
1	2010-11	
2	2011-12	
3	2012-13	
Total		₹ Lakhs
Average turnover per annual		₹ Lakhs

Date:

Seal:

**Signature of Auditor/
Chartered Accountant
(Name in Capital)**

**ANNEXURE-III
Ref. Clause No.14 (a)**

AGREEMENT

This Deed of Agreement is made on this _____day of _____2013 by M/s. _____ represented by its Proprietor/Managing partner/Managing Director having its Registered Office

at _____ and its Factory Premises
at _____

(hereinafter referred to as "Supplier" which term shall include its successors, representatives, heirs, executors and administrators unless excluded by the Contract) on one part and AIIMS, Raipur (C.G.) represented by its Director having is office at Tatibandh GE Road Raipur (hereinafter referred to as "The Purchaser" which term shall include its successors, representatives, executors assigns and administrator unless excluded by the Contract) on the other part.

Where as the Supplier has agreed to supply to the Purchaser, the Drugs and Medicines with specifications mentioned in the Schedule attached here to at the prices noted there in and in the manner and under the terms and conditions here in after mentioned and where as the Supplier has deposited with the Purchaser a sum of ₹ _____

(Rupees only) as Security Deposit for the due and faithful performance of this Agreement, to be forfeited in the event of the Supplier failing duly and faithfully to perform it. Now these presents witness that for carrying duly and faithfully to perform it. Now these presents witness that for carrying out the said Agreement in this behalf into execution the Supplier and the Purchaser do hereby mutually covenant, declare, contract and agree each of them with the other of them in the manner following, that is to say.

1. The term "Agreement", wherever used in this connection, shall mean and include the terms and conditions contained in the invitation to Bid floated for the rate contract cum supply for Drug & Medicines For AIIMS, Raipur (C.G.), the instruction to Bidders, the conditions of Bid, acceptance of Bid, particulars hereinafter defined and those general and special conditions that may be added from time to time.

2.

(a) The Agreement is for the supply by the Supplier to the Purchaser of the Drug and Medicines specified in the agreement on the terms and conditions set forth in the Agreement.

(b) The Agreement is for the supply by the Supplier to the Purchaser of the Drug and Medicines specified in the agreement on the terms and conditions set forth in the Agreement.

(c) This Agreement shall be deemed to have come into force with effect from the _____ and it shall remain in force upto _____.

(d) The Bid quantity noted against each item in the schedule attached hereto indicates only the probable total requirements of the Purchaser in respect of each item for the Agreement Period indicated in Clause.

(e) above. This quantity may increase or decrease at the discretion of the Purchaser. The Supplier shall make supplies of the Drugs and Medicines on the basis of the Purchaser Orders placed on him from time to time by the ordering Authorities of the purchaser specifying the quantities required to be supplied required to be supplied at the specific location, Hospital Store Segment of AIIMS, Raipur (C.G.).

TERMINATION OF CONTRACT ON BREACH OF CONDITION

1. (a) In case the Supplier fails or neglects or refuse to faithfully perform any of the Covenants on his part herein contained, it shall be lawful for the Purchaser to forfeit the amount deposited by the Supplier as Security Deposit and cancel the Contract.

(b) In case the Supplier fails, neglects, or refuse to observe, perform, fulfill and keep, all or any one or more or any part of any one of the Covenants, stipulation and provisions herein contained, it shall be lawful for the Purchaser on any such failure, neglect or refusal, to put an end to this Agreement and thereupon every article, cause and thing herein contained on the part of the Purchaser shall cease and be void, and in case of any damage, loss, expenses, difference in cost or other moneys from out of any moneys for the time being.

payable to the Supplier under this and/or any other Contract and in case such last mentioned moneys are insufficient to cover all such damages, losses, expenses, difference in cost and other moneys as aforesaid, it shall be lawful for the Purchaser to appropriate the Security Deposit made by the Supplier as herein before mentioned to reimburse all such damages, losses, expenses, difference in cost and other money as the Purchaser shall have sustained, incurred or been put to by reason of the Supplier having been guilty of any such failure, negligence or refusal as aforesaid or other breach in the performance of this Contract.

(c) If at any time during the course of the Contract, it is found that any information furnished by the Supplier to the Purchaser, either in his Bid or otherwise, is false, the Purchaser may put an end to the Contract/Agreement wholly or in part and thereupon the provisions of Clause (a) above shall apply.

2. The Purchaser reserves the right to terminate without assigning any reasons therefore the Contract/Agreement either wholly or in part without any notice to the Supplier. The Supplier will not be entitled for any compensation whatsoever in respect of such termination of

the Contract/Agreement by the Purchaser.

NOTICE ETC. IN WRITING

3. All Certificates or Notice or orders for time or for extra, varied or altered supplies which are to be the subject of extra or varied charges whether so described in the Agreement or not, shall be in writing, and unless in writing, shall not be valid, binding or be of any effect whatsoever.

SUPPLIERS NOT HAVE ANY INTEREST IN THE OFFICERS CONCERNED AND SUBORDINATES

4. The Supplier shall not be in any way interested in or concerned directly or indirectly with, any of the Officers, Subordinate or Servants of the Purchaser. In any trade, business or transactions nor shall the Supplier give or pay or promise to give or pay any such Officer, Subordinate or Servant directly or indirectly any money or fee or other consideration under designation of "Custom" or otherwise; nor shall the Supplier permit any person or persons whomsoever to interfere in the management or performance hereof under power of attorney or otherwise without the consent in writing the consent in writing of the Purchaser obtained in first hand.

BANKRUPTCY OF THE SUPPLIER

5. In case the Supplier at any time during the continuance of the Contract becomes bankrupt or insolvent or commits any act of bankruptcy or insolvency under the provisions of any law in that behalf for the time being in force, or should compound with his creditors, it shall be lawful for the Purchaser to put an end to the Agreement, and thereupon every article, clause and thing herein contained to be operative on the part of the Purchaser, shall cease and be void and the Purchaser shall have all the rights and remedies given to him under the preceding clauses.

SERVING OF NOTICE ON SUPPLIER

6. All notice or communication relating to or arising out of this Agreement or any of the terms thereof shall be considered duly served on or given to the Supplier if delivered to him or left at his

premises, place of business or abode.

7. And it is hereby agreed and declared between the parties hereto that in case any question of dispute arises touching the construction or wording of any of clause herein contained on the rights, duties, liabilities of the parties hereto or any other way, touching or arising out of the presents, the decision of the Director, AIIMS, Raipur (C.G.) in the matter shall be final and binding.
8. All disputes arising out of this agreement and all questions relating to the interpretation of this agreement shall be decide by the Govt. and the decision of the Govt. shall be final.

SUPPLIER

Director

Witness (Signature, Name & Address)

Witness

- 1.
- 2.

- 1.
- 2.

**ANNEXURE -IV
Ref. Clause No. 5(u)**

Check List

Section	Details of	Document Type	Yes/No
---------	------------	---------------	--------

	requirement		If Yes Page No.
A	EMD, Empanelment Fees.	FDR/BC/DD of EMD, tender fee Annexure-I Bank Guaranty	
B	Technical documents	Manufacturing License	
		Manufacturing License renewal /validity certificate	
		Non Conviction Certificate issued by the Drugs Controller	
		Good Manufacturing Practices Certificate	
		Import License, if imported.	
		Sale License, in the case of imported drugs	
		Copy of record of import to establish 3 years market standing, if imported.	
		Product Permissions by the Licensing Authority for each and every product quoted	
		Market Standing Certificate issued by the licensing Authority	
		Annexure-V Check List Of Details Regarding Products Quoted	
C	Other Documents	Documentary evidence for the constitution of the company / concern	
		The instruments such as power of attorney resolution of board etc	
		Copies of balance sheet & profit loss account for three years	
		Sales Tax clearance certificate	
		Excise Registration Certificate	
		Copy of PAN	
		Annual Turnover Statement	
		Annexure-VI Declaration and Undertaking	
		Annexure-IX Undertaking For Empanelment	

Annexure - V
Ref. Clause No. 5 (u)

Check list of details regarding products quoted

Product permission as per condition no. 5 (c) and Market Standing as per condition 5 (g)									
S.No.	Quoted Item/ Code NO.	Product permission enclosed on Page no.	Date of product permission/ Approval	Production permission of formulation Generic/ Branded	Specification as per Quoted Item Yes/No	As per MSC product Mfg & Mkd Since Last 3 year		Attested	Remark
						Page No.	Yes/ No.		

Annexure - VI
Ref. Clause No. 5 (o),(q),(r),(t),(v),(w)

Declaration & Undertaking

**(for Tender Enquiry No. ADMIN/Tender/DRUGS/1/2013 Dated : 02.11.2013)
(On Non-Judicial Stamp Paper of ₹ 500/- Attested by Notary Public)**

I Name.....S/o.....Age.....Prop./Partner/Director/Power of attorney holder of firm M/s.....situated at (Complete address of Mfg. unit).....bearing drug license on Form 25 & 28 **or form 10** bearing Number..... &.....respectively, issued on dated.....valid/Renewed up to.....do here by declare on oath as follows:-

1. That none of the quoted Drug and Medicines manufactured / imported by us since grant of above drug license have been found as of spurious or adulterated quality and no case in this regard is pending in any court.
2. That the quoted product is manufactured/imported by us, and none has been declared as “Not of standard quality” during last two years.
3. That we have following Commitment of quantity in our plant at above address:-**[Ref. Clause No. 25(i)]**.

S.No.	Quoted item & Name of Drugs	Monthly Capacity in all shifts in nos.	Annual Production Capacity	Monthly Supply Commitment ao AIIMS, Raipur in nos.	Monthly Supply Commitment ao AIIMS, Raipur in nos.	Estimated Bid Quantity as per Annexure VII
1.						
2.						
3.						

4. That concern/company/firm does not stand blacklisted/banned/debarred on any ground by Bid Inviting Authority or any Govt. Hospital State/Central/PSU **or its departments** on the date of bid submission. The concern/company/firm does not stand blacklisted/banned/debarred on the ground of **conviction by court of law or the products being found spurious or adulterated** by any other State /Central Government or **it's any agencies** (central Drugs procurement agencies).
5. That our Firm/Company and its Proprietor/Partner/Directors/ Power of attorney holders have not been convicted for contravention of any provisions of Drugs & Cosmetic Act 1940 and rules made there under since grant of license

6. That we have been granted product permission by the State Licensing Authority for manufacture of quoted products as per the details given below:-

S. No.	Name of the product	Specification IP/BP/USP/ Other	Date of product permission obtained from the Licensing Authority	Whether Endorsement is in Generic or Trade Name	Issuing Licensing Authority
1.					
2.					

7. That we have over three years' experience in the manufacture of the quoted product, or the quoted imported product has over 3 years market standing.

8. That we have approved qualified staff, machines & equipments along with capacity to manufacture above category of drugs and our unit have been issued G.M.P.* Certificate as per Schedule M by State Licensing Authority vide letter No.....dated.....valid upto.....

9. That we hereby confirm that we have deposited all the VAT/Sale Tax as on.....With the department No VAT/CST is due on M/s.....as on.....

10. That I will supply the Drug and Medicines per the designs given in Bid clause no 15 and as per the instructions given in this regard.

11. That I/We have carefully read all the conditions of Bid in Ref. no. Tender Enquiry No. ADMIN/Tender/Drugs/1/2013 dated: 02.11.2013 for Annual Rate Contract cum Supply, of Drugs and Medicines for **AIIMS, Raipur** and accept all conditions of Bid, including amendments if any.

12. I/We agree that the Bid Inviting Authority forfeiting the Earnest Money Deposit and or Security Deposit and blacklisting /Debarring/Banning me/ us for a period of 5 years or as deemed fit if, any information furnished by us proved to be false/fabricated at the time of inspection and not complying the conditions as per Schedule M of the said Act or at any time during the Bid process.

13. Our complete address for communication

.....

E-mail address: -

Phone No. /Mobile No.....

14. Bank detail for e banking:-

Name of account holder

Full name of Bank with Branch

A/c no. with full digits.....

IFSC code

Photograph of
Authorized/
nominating
person

15. Authorized/nominating person.

Name:

Designation:-.....

Complete address for communication:-

.....

.....

.....

E-mail address: -

Phone No. /Mobile No.....

(Name of Deponent & Signature)

Designation

Verification

I.....S/o.....(Designation).....Affirm on oath that the contents/information from para 1 to 16 as mentioned above, are true & correct to the best of my knowledge and nothing is hidden. I also declare on oath, that if any information furnished by me as above is found wrong, false, forged or fabricated; the Corporation will be at liberty to cancel the Bid for which I shall be solely responsible and the firm may be Debarred/Banned/ blacklisted / prosecuted for the same.

(Name of Deponent & Signature)

Witness :- (Name, Address & Signature)

1

2

*The GMP certificate must not be older than one year from the last date of Bid submission in case validity is not mentioned in the certificate.

Annexure - VII

Ref. Clause No. 9 (2, 3)

List of Groups/Drugs & their Indicators

Mian Code Indicators for Medicine Name

An	=	Anaesthetic
Al	=	Analgesic
Ar	=	Antiallergics and Medicines used in Anaphylaxis
Ad	=	Antidotes and Chelating agent
Ae	=	Antiepileptic
Ab	=	Antiinfective/ Antibiotics
Am	=	Antimalarial
Ac	=	Antineoplastic
Ap	=	Antiparkinsonism medicines
Cv	=	Cardiovascular medicine
Dm	=	Dermatological medicines
Da	=	Diagnostic agents
Di	=	Disinfectants and antiseptics
Du	=	Diuretics
En	=	ENT
Gi	=	Gastrointestinal medicines
Ho	=	Hormones, other endocrine medicines and contraceptives
Im	=	Immunologicals
Re	=	Medicines acting on the respiratory tract
Bl	=	Medicines affecting the blood
Cn	=	Medicines for nervous system disorders
Mr	=	Muscle relaxants
Op	=	Ophthalmological
Ox	=	Oxytocics and Antioxytocics
Va	=	Vaccine, Sera and immunomodulators
Vi	=	Vitamins and Minerals
Ms	=	Miscellaneous

Sub Code Indicators for Medicines/Consumables

T	=	Tablet/ Capsule
I	=	Injection
F	=	Fluids
L	=	Liquid (Syrup)
C	=	Consumable
S	=	Surgical
M	=	Other then above.

BOQ & Financial BID

Most Important & Essential

- Bidders are instructed to mention Code & Sub code No. of Group/ medicine/ consumable/ surgical on the top of package along with other specifications mentioned.
- The Rate Quoted in column No. "I" will decide L₁. For given Medicine/Surgical/Consumable.
- The total amount quoted in column no. "I" shall be Inclusive of transportation, insurance, packing and any incidental charges, Excise Duty, Customs duty & all other statutory duties of the govt. EXCEPT VAT and CST.

S.No.	Medicine Code	Name of Medicine (Drugs)	Dosage Forms	Strength	Require Unit/Year	Packing Unit	Rate of Packing Unit in ₹ Inclusive of transportation, insurance, Packing and any incidental charges, Excise Duty, Customs duty & all other statutory duties of the govt. <u>EXCEPT VAT & CST</u>	Total Rate of Required Unit/Year in ₹ Inclusive of transportation, insurance, Packing and any incidental charges, Excise Duty, Customs duty & all other statutory duties of the govt. <u>EXCEPT VAT & CST</u>
A	B	C	D	E	F	G	H	I
	An	Anaesthetic						
1	An-M/001	EMLA Cream	Cream	(2.5%/2.5%) 30g tube	1200	1x10		
2	An-M/002	Proparacaine hydrochloride	Eye Drops	0.50%	180	1x10		
3	An-F/001	Desflurane	Inhalation	500ml bottle	240	1x10		
4	An-F/002	Isoflurane	Inhalation	250 ml Bottle	360	1x10		
5	An-F/003	Sevoflurane	Inhalation	250ml PEN bottle	360	1x10		
6	An-I/001	Bupivacaine hydrochloride	Injection	0.25%, 20ml Vial	300	1x10		
7	An-I/002	Bupivacaine hydrochloride	Injection	0.5%, 20 ml Vial	300	1x10		

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8	An-I/003	Bupivacaine hydrochloride	Injection	0.5% to be mixed with 7.5% glucose, 4 ml Ampoule	600	1x10		
9	An-I/004	Ketamine hydrochloride	Injection	10mg/ml in vial	120	1x10		
10	An-I/005	Ketamine hydrochloride	Injection	50mg/ml in vial	120	1x10		
11	An-I/006	Lignocaine + Adrenaline	Injection	1%, with adrenaline 1:2000 00,30 ml Vial	120	1x10		
12	An-I/007	Lignocaine + Adrenaline	Injection	2% with adrenaline 1:2000 00,30 ml Vial	120	1x10		
13	An-I/008	Lignocaine Hydrochloride	Injection	2%, 30 ml Vial	2400	1x10		
14	An-I/009	Midazolam	Injection	1mg/ml , 5ml vial	3600	1x10		
15	An-I/010	Procainamide hydrochloride	Injection	100mg/ml, 10 ml vial	60	1x10		
16	An-I/011	Propofol	Injection	1% (Oil suspension) 20ml vial	1200	1x10		
17	An-I/012	Thiopentone sodium	Injection	0.5gm Powder , 50 ampoules pack	1200	1x10		
18	An-I/013	Thiopentone sodium	Injection	1gm Powder , 50 ampoules pack	1200	1x10		
19	An-I/014	Etomidate	Injection	2	300	1x10		

				mg/ml				
20	An-I/015	Lignocaine Hydrochloride	Spinal	5% + 7.5% Glucose, 2 ml ampoules	600	1x10		
21	An-T/001	Procainamide hydrochloride	Tablet	250mg	120	10x10		
22	An-M/003	Lignocaine Hydrochloride	Topical Forms	2%, 30 ml Vial	1200	1x10		
23	An-M/004	Lignocaine Hydrochloride	Topical Forms	5%, 30 ml Vial	1200	1x10		
	AI	Analgesic						
24	AI-T/001	Aspirin	Tablet A	75mg, Tablet (Enteric Coated)	2400	10x10		
25	AI-T/002	Aspirin	Tablet B	150mg, Tablet (Enteric Coated)	1200	10x10		
26	AI-T/003	Aspirin	Tablet C	325mg, Tablet (Enteric Coated)	1200	10x10		
27	AI-M/001	Diclofenac Gel	Gel	1% w/v, 25 gm tube	2400	1x10		
28	AI-I/001	Diclofenac sodium	Injection	25mg/ml PG surfactant free, 3ml ampoules	6000	1x10		
29	AI-T/004	Diclofenac sodium	Tablet	50mg	10800	10x10		
30	AI-I/002	Fentanyl	Injection	50µg/ml, 2 ml ampoules	1200	1x10		
31	AI-T/005	Ibuprofen	Tablet	200mg	10200	10x10		
32	AI-T/006	Indomethacin	Tablet	25mg	1200	10x10		
33	AI-T/007	Indomethacin	Tablet	50mg	1200	10x10		
34	AI-I/003	Ketorolac	Injection	30mg/ml, 1 ml ampoules	600	1x10		
35	AI-T/008	Mefenamic acid	Tablet	250 mg	6000	10x10		
36	AI-I/004	Morphine sulphate	Injection	10mg/ml, 1ml	2400	1x10		

				ampoule				
37	AI-T/009	Naproxen	Tablet	250 mg	600	10x10		
38	AI-M/002	Nepafenac	Eye Drops	0.1% ophthalmic Suspension	120	1x10		
39	AI-I/005	Paracetamol	Injection	150mg/ml, 2 ml ampoule	6000	1x10		
40	AI-L/001	Paracetamol	Syrup	125mg/5ml, 60ml bottle	2400	1x10		
41	AI-T/010	Paracetamol	Tablet	500mg	36000	10x10		
42	AI-T/011	Piroxicam	Tablet	20mg	3000	10x10		
43	AI-I/006	Tramadol	Injection	50mg/ml, 1ml ampoule	4200	1x10		
44	AI-T/012	Tramadol	Tablet	50mg	3600	10x10		
	Ar	Antiallergics						
45	Ar-I/001	Adrenaline	Injection	1mg/ml, 1ml ampoule	4200	1x10		
46	Ar-I/002	Alamine	Injection	infusion, 200ml infusion	600	1x10		
47	Ar-L/001	Cetirizine	Syrup	5mg/ml, 60 ml bottle	2400	1x10		
48	Ar-T/001	Cetirizine	Tablet	10mg	9000	10x10		
49	Ar-T/002	Chlorpheniramine maleate	Tablet	4mg	3600	10x10		
50	Ar-I/003	Dexamethasone	Injection	4mg/ml, 2 ml Vial	1200	1x10		
51	Ar-T/003	Dexamethasone	Tablet	0.5mg	5400	1x10		
52	Ar-L/002	Dexchlorpheniramine maleate	Syrup	0.5mg/ml	1200	1x10		
53	Ar-T/004	Montelukast	Tablet	10 mg	4800	10x10		
54	Ar-T/005	Montelukast	Tablet	4 mg	4800	10x10		
55	Ar-T/006	Montelukast	Tablet	5 mg	4800	10x10		
56	Ar-I/004	Pheniramine maleate	Injection	22.75mg/ml, 2ml vial	600	1x10		
57	Ar-I/005	Promethazine	Injection	25mg/ml, 2ml	4200	1x10		

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				ampou le				
58	Ar-L/003	Promethazine	Syrup	5mg/ml , 60 ml bottle	600	1x10		
59	Ar-T/007	Promethazine	Tablet	25mg	2400	10x10		
	Ad	Antidotes						
60	Ad-M/001	Atropine	Eye Drops	1%, 5 ml pack	60	1x10		
61	Ad-M/002	Atropine	Eye Ointment	1%, 5 ml pack	60	1x10		
62	Ad-I/001	Atropine	Injection	0.6mg/ ml, 2ml ampou le	3600	1x10		
63	Ad-I/002	Methylene blue	Injection	10mg/ ml, 10 ml ampou le	600	1x10		
64	Ad-M/003	Methylene blue	Lotion	10ml vial	60	1x10		
65	Ad-I/003	N-acetylcysteine	Injection	200mg/ ml (5ml)	1800	1x10		
66	Ad-I/004	Naloxone	Injection	0.4mg/ ml, 1 ml Amp	600	1x10		
67	Ad-I/005	Pralidoxime chloride	Injection	25mg/ ml, 20 ml Vial	180	1x10		
68	Ad-I/006	Sodium nitrite	Injection	30mg/ ml	60	1x10		
69	Ad-I/007	Sodium thiosulfate	Injection	250mg/ ml	60	1x10		
70	Ad-M/004	Activated Charcoal	Oral	Unit pack	3600	1x10		
71	Ad-T/001	Penicillamine	Tablet/Caps ule	250mg	120	10x10		
72	Ad-I/008	Desferrioxamine	Injection	500mg	180	1x10		
73	Ad-I/009	Dimercaprol	Injection	50mg/ ml, 2 ml Ampou le	60	1x10		
74	Ad-I/010	Flumazenil	Injection	0.1mg/ ml, 5 ml Ampou le	600	1x10		
75	Ad-T/002	Deferiprone	Tablet	500 mg, 50 caps pack	360	50x10		

76	Ad-T/003	Deferiprone	Tablet	250 mg, 50 caps pack	360	50x10		
	Ae	Antiepileptic						
77	Ae-I/001	Magnesium sulfate	Injection	500mg/ ml, 2 ml Ampoul e	3600	1x10		
78	Ae-I/002	Phenytoin sodium	Injection	50mg/ ml, 2 ml Ampoul e	3600	1x10		
79	Ae-I/003	Sodium Valproate	Injection	100mg/ ml, 5 ml vial	1200	1x10		
80	Ae-L/001	Carbamazepine	Syrup	100mg/ 5ml, 100 ml bottle	360	1x10		
81	Ae-L/002	Sodium Valproate	Syrup	200mg/ 5ml, 100 ml bottle	1200	1x10		
82	Ae-T/001	Carbamazepine	Tablet A	100mg	8400	10x10		
83	Ae-T/002	Carbamazepine	Tablet B	200mg	8400	10x10		
84	Ae-T/003	Sodium Valproate	Tablet A	200mg	9600	10x10		
85	Ae-T/004	Sodium Valproate	Tablet B	500mg	9600	10x10		
86	Ae-T/005	Phenytoin sodium	Tablet/Capsule A	50mg	2400	10x10		
87	Ae-T/006	Phenytoin sodium	Tablet/Capsule B	100mg	360	10x10		
88	Ae-I/004	Lorazepam	Injection	2mg/ml , 2ml ampoul e	4200	1x10		
89	Ae-T/007	Lorazepam	Tablet A	1mg	3600	10x10		
90	Ae-T/008	Lorazepam	Tablet B	2mg	600	10x10		
	Ab	Antiinfective						
91	Ab-T/001	Amoxicillin	Capsule A	250mg	3600	10x10		
92	Ab-T/002	Amoxicillin	Capsule B	500mg	3600	10x10		
93	Ab-T/003	Cephalexin	Capsule A	250mg	3000	10x10		
94	Ab-T/004	Cephalexin	Capsule B	500mg	3000	10x10		
95	Ab-T/005	Chloramphenicol	Capsule	250mg	120	10x10		
96	Ab-T/006	Clofazimine	Capsule A	50mg	1200	10x10		
97	Ab-T/007	Clofazimine	Capsule B	100mg	1200	10x10		
98	Ab-T/008	Cloxacillin	Capsule A	500mg	4200	10x10		

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99	Ab-T/009	Cloxacillin	Capsule B	1gm	1200	10x10		
100	Ab-T/010	Rifampicin	Capsule A	50mg	1560	10x10		
101	Ab-T/011	Rifampicin	Capsule B	150mg	120	10x10		
102	Ab-T/012	Rifampicin	Capsule C	300mg	120	10x10		
103	Ab-T/013	Rifampicin	Capsule D	450mg	120	10x10		
104	Ab-T/014	Fluconazole	Capsule/Tab let A	50mg	1800	10x10		
105	Ab-T/015	Fluconazole	Capsule/Tab let B	100mg	1200	10x10		
106	Ab-T/016	Fluconazole	Capsule/Tab let C	150mg	1200	10x10		
107	Ab-T/017	Fluconazole	Capsule/Tab let D	200mg	1200	10x10		
108	Ab-T/018	Griseofulvin	Capsule/Tab let A	150mg	600	10x10		
109	Ab-T/019	Griseofulvin	Capsule/Tab let B	250mg	600	10x10		
110	Ab-M/001	Acyclovir	Cream	5%, 15gm tube	240	1x10		
111	Ab-M/002	Permethrin	Cream	5%, 30g tube	600	1x10		
112	Ab-M/003	Silver sulfadiazine	Cream	1%, 50g, 50g tube	600	1x10		
113	Ab-M/004	Ciprofloxacin	Ear Drops	0.003, 5ml vial	1200	1x10		
114	Ab-M/005	Chloramphenico l	Eye Drops	0.004,p ack of 50 applica ps	600	1x10		
115	Ab-M/006	Ciprofloxacin	Eye Drops	0.003, 5ml vial	360	1x10		
116	Ab-M/007	Miconazole	Eye Drops	1%, 10ml vial	600	1x10		
117	Ab-M/008	Moxifloxacin	Eye Drops	0.5%, 5ml vial	240	1x10		
118	Ab-M/009	Natamycin	Eye Drops	5%, 5ml vial	120	1x10		
119	Ab-M/010	Tobramycin	Eye Drops	0.003, 5ml vial	1200	1x10		
120	Ab-M/011	Chloramphenico l	Eye Ointment	1%, pack of 50 applica ps	1200	1x10		
121	Ab-M/012	Chloramphenico l Hydrocortisone	Eye Ointment	30g tube	120	1x10		

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122	Ab-M/013	Ciprofloxacin	Eye Ointment	0.003, 30g tube	120	1x10		
123	Ab-M/014	Clotrimazole	Gel	0.02, 5ml vial	1200	1x10		
124	Ab-I/001	Amikacin	Injection A	250mg/2ml, vial	8400	1x10		
125	Ab-I/002	Amikacin	Injection B	500mg/2ml, vial	2400	1x10		
126	Ab-I/003	Amoxicillin + Clavulanic acid	Injection A	(500mg +100mg), 600mg	4200	1x10		
127	Ab-I/004	Amoxicillin + Clavulanic acid	Injection B	1.2gm, vial	1200	1x10		
128	Ab-I/005	Amphotericin B	Injection	50mg, vial	1800	1x10		
129	Ab-I/006	Ampicillin	Injection	500mg, vial	4800	1x10		
130	Ab-I/007	Azithromycin	Injection	500mg	1200	1x10		
131	Ab-I/008	Benzathine penicillin	Injection A	6 Lac units	600	1x10		
132	Ab-I/009	Benzathine penicillin	Injection B	12 Lac units	600	1x10		
133	Ab-I/010	Cefotaxime	Injection A	125mg, vial	6600	1x10		
134	Ab-I/011	Cefotaxime	Injection B	250mg, vial	2400	1x10		
135	Ab-I/012	Cefotaxime	Injection C	500mg, vial	2400	1x10		
136	Ab-I/013	Ceftazidime	Injection A	250mg, vial	3600	1x10		
137	Ab-I/014	Ceftazidime	Injection B	1g vial	1200	1x10		
138	Ab-I/015	Ceftriaxone	Injection A	250mg, vial	8400	1x10		
139	Ab-I/016	Ceftriaxone	Injection B	1gm	8400	1x10		
140	Ab-I/017	Cefuroxime	Injection A	250mg	4200	1x10		
141	Ab-I/018	Cefuroxime	Injection B	500mg	4200	1x10		
142	Ab-I/019	Cefuroxime	Injection C	1gm	3600	1x10		
143	Ab-I/020	Ciprofloxacin	Injection	200mg/100ml	5400	1x10		
144	Ab-I/021	Clindamycin	Injection	150mg/ml, 2 ml ampoule	600	1x10		
145	Ab-I/022	Gentamicin	Injection A	10mg/ml	6600	1x10		
146	Ab-I/023	Gentamicin	Injection B	40mg/ml	3600	1x10		
147	Ab-I/024	Linezolid	Injection	vial	3000	1x10		

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148	Ab-I/025	Metronidazole	Injection	500mg/ 100ml	7200	1x10		
149	Ab-I/026	Ofloxacin	Injection	IV Infusion, 200mg/ ml	3600	1x10		
150	Ab-I/027	Pentamidine	Injection	200mg/ vial	180	1x10		
151	Ab-I/028	Sodium stibogluconate	Injection	100mg/ ml	120	1x10		
152	Ab-I/029	Streptomycin	Injection A	0.75gm / vial	600	1x10		
153	Ab-I/030	Streptomycin	Injection B	1gm	600	1x10		
154	Ab-I/031	Vancomycin hydrochloride	Injection A	500mg	6000	1x10		
155	Ab-I/032	Vancomycin hydrochloride	Injection B	1gm	600	1x10		
156	Ab-I/033	Acyclovir	Injection A	250mg	600	1x10		
157	Ab-I/034	Acyclovir	Injection B	500mg	600	1x10		
158	Ab-I/035	Benzyl penicillin	Injection (Powder)	10 Lac units (600mg)	3000	1x10		
159	Ab-I/036	Cefazolin	Injection (Powder)	1gm (as sodium salt) in vial	4200	1x10		
160	Ab-I/037	Cloxacillin	Injection (Powder)	500mg in vial	7800	1x10		
161	Ab-I/038	Chloramphenicol	Injection (sodium succinate)	1gm	600	1x10		
162	Ab-M/0015	Permethrin	Lotion A	1%, 50ml bottle	300	1x10		
163	Ab-M/0016	Permethrin	Lotion B/ Cream	5%, 30g bottle	120	1x10		
164	Ab-M/0017	Miconazole	Ointment	15g tube	240	1x10		
165	Ab-M/0018	Mupirocin	Ointment	15g tube	1200	1x10		
166	Ab-M/0019	Neomycin + Bacitracin	Ointment	5mg + 500 IU /gm	1200	1x10		
167	Ab-M/0020	Chloramphenicol	Oral Suspension (Palmitate)	150mg/ 5ml, 60ml bottle	300	1x10		
168	Ab-M/0021	Clotrimazole	Pessary A	100mg	120	10x10		
169	Ab-M/0022	Clotrimazole	Pessary B	200mg	120	10x10		
170	Ab-M/0023	Nystatin	Pessary	100000	120	10x10		

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				IU				
171	Ab-M/0024	Clotrimazole	Powder	30g vial	1200	1x10		
172	Ab-I/039	Imipenem + Cilastatin	Powder for injection	250mg (as monohydrate) + 250 mg(as sodium salt), 30ml bottle	1200	1x10		
173	Ab-I/040	Imipenem + Cilastatin	Powder for injection	500mg (as monohydrate) + 500mg (as sodium salt), 30ml bottle	120	1x10		
174	Ab-M/025	Cloxacillin	Powder for oral solution	125mg/5ml, 30ml bottle	1800	1x10		
175	Ab-M/026	Amoxicillin	Powder for suspension	125mg/5ml, 30ml bottle	3000	1x10		
176	Ab-M/027	Amoxicillin + Clavulanic acid	Powder for suspension	228.5mg/5ml, 30ml bottle	4800	1x10		
177	Ab-M/028	Ampicillin	Powder for suspension	125mg/5ml, 30ml bottle	1200	1x10		
178	Ab-M/029	Acyclovir	Suspension	400mg/5ml, 30ml	240	1x10		
179	Ab-M/030	Albendazole	Suspension	200mg/5ml, 10ml bottle	600	1x10		
180	Ab-L/001	Azithromycin	Syrup	100mg/5ml, 15 ml bottle	1200	1x10		
181	Ab-L/002	Cephalexin	Syrup	125mg/5ml/30 ml	1200	1x10		
182	Ab-L/003	Erythromycin	Syrup	125mg/	240	1x10		

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		estolate		5ml/30 ml				
183	Ab-L/004	Metronidazole	Syrup	100mg/5ml/30 ml	1800	1x10		
184	Ab-L/005	Rifampicin	Syrup	100mg/5ml/30 ml	120	1x10		
185	Ab-T/020	Acyclovir	Tablet A	200mg	2400	10x10		
186	Ab-T/021	Acyclovir	Tablet B	400mg	2400	10x10		
187	Ab-T/022	Albendazole	Tablet	400mg	2400	1x10		
188	Ab-T/023	Amoxicillin + Clavulanic acid	Tablet	(500mg + 125mg), 625mg, 6 tab pack	30000	1 x 10		
189	Ab-T/024	Ampicillin	Tablet A/	250mg	3000	10x10		
190	Ab-T/025	Ampicillin	Tablet A/Capsules IP 500 mg	500mg, 10x10 cap blister	1200	10x10		
191	Ab-T/026	Azithromycin	Tablet A	100mg, 6 tab pack	6000	1x10		
192	Ab-T/027	Azithromycin	Tablet B	250mg, 6 tab pack	600	1x10		
193	Ab-T/028	Azithromycin	Tablet C	500mg, 6 tab pack	600	1x10		
194	Ab-T/029	Cefixime	Tablet A/Capsule	100mg	12000	10x10		
195	Ab-T/030	Cefixime	Tablet B/Capsule	200mg	12000	10x10		
196	Ab-T/031	Cefuroxime	Tablet A	250mg	6000	10x10		
197	Ab-T/032	Cefuroxime	Tablet B	500mg	6000	10x10		
198	Ab-T/033	Ciprofloxacin	Tablet A	250mg	18000	10x10		
199	Ab-T/034	Ciprofloxacin	Tablet B	500mg	18000	10x10		
200	Ab-T/035	Clindamycin	Tablet A	150mg	3600	10x10		
201	Ab-T/036	Clindamycin	Tablet B	300mg	3600	10x10		
202	Ab-T/037	Diethylcarbamazine	Tablet	50mg	1800	10x10		
203	Ab-T/038	Diloxanide Furoate	Tablet	500mg	240	10x10		
204	Ab-T/039	Doxycycline	Tablet/capsules	100mg	8400	10x10		
205	Ab-T/040	Erythromycin estolate	Tablet A	250mg	1200	10x10		
206	Ab-T/041	Erythromycin estolate	Tablet B	500mg	2400	10x10		

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207	Ab-T/042	Levofloxacin	Tablet	500mg	9600	10x10		
208	Ab-T/043	Linezolid	Tablet	600mg	6000	10x10		
209	Ab-F/001	Linezolid	Infusion	600mg, 300ml infusion bottle	6000	1x10		
210	Ab-T/044	Metronidazole	Tablet A	200mg	12000	10x10		
211	Ab-T/045	Metronidazole	Tablet B	400mg	6000	10x10		
212	Ab-T/046	Nitrofurantoin	Tablet	100mg	3600	10x10		
213	Ab-T/047	Nystatin	Tablet	500000 IU	1200	10x10		
214	Ab-T/048	Ofloxacin	Tablet A	100mg	8400	10x10		
215	Ab-T/049	Ofloxacin	Tablet B	200mg	6000	10x10		
216	Ab-T/050	Praziquental	Tablet	600mg	360	10x10		
217	Ab-T/051	Tinidazole	Tablet	500mg	4800	10x10		
218	Ab-I/041	Capsfungin	Injection	50mg Vial	180	1x10		
219	Ab-I/042	Capsfungin	Injection	70mg vial	120	1x10		
220	Ab-I/043	Cefoparazone + Sulbactum	Injection	(1gm+0 .5gm) powder for injection 20ml vial	2400	1x10		
221	Ab-F/002	Colistin sulphet	syrup	25mg in 30ml bottle	1200	1x10		
222	Ab-I/044	Meropenam	Injection	1gm in Vial	2400	1x10		
223	Ab-T/052	Nelfinavir	Tab	250mg 100 tab/stri p	600	1x10		
224	Ab-T/053	Oseltamivir	capsule	30mg	600	10x10		
225	Ab-T/054	Oseltamivir	capsule	45mg	1200	10x10		
226	Ab-T/055	Oseltamivir	capsule	75mg	2400	10x10		
227	Ab-T/056	Oseltamivir	Dry powder for oral suspension	12mg/ ml	600	1x10		
228	Ab-T/057	Oseltamivir	syrup	100ml bottle	120	1x10		
229	Ab-T/058	Piperacillin + Tazobactum	Injection	(3+1.5) gm in vial	3600	1x10		
230	Ab-T/059	Piperazine	tablet	500mg	300	10x10		
	Am	Antimalarial						
231	Am-I/001	Artesunate	Injection	60mg, 2ml ampoul	3600	1x10		

				e				
232	Am-I/002	Chloroquinine	Injection	40mg/ ml	1200	1x10		
233	Am-I/003	Quinine	Injection	300mg/ ml	1200	1x10		
234	Am-L/001	Chloroquinine	Syrup	50mg/5 ml	600	1x10		
235	Am-T/001	Artemether + Lumefantrine	Tablet	20mg + 120mg	4800	10x10		
236	Am-T/002	Artesunate	Tablet	50mg	3000	10x10		
237	Am-T/003	Chloroquinine	Tablet	150mg base	8400	10x10		
238	Am-T/004	Mefloquine	Tablet	250mg (base)	1200	10x10		
239	Am-T/005	Primaquine	Tablet A	7.5mg	1200	10x10		
240	Am-T/006	Primaquine	Tablet B	15mg	600	10x10		
241	Am-T/007	Quinine	Tablet	300mg	600	10x10		
242	Am-T/008	Sulfadoxine + Pyrimethamine	Tablet	500mg + 25mg	3000	10x10		
	Ac	Antineoplastic						
243	Ac-T/001	Hydroxyurea	Capsule	500mg	240	10x10		
244	Ac-T/002	Etoposide	Capsules	100mg, 4cap.Pa ck	120	10x10		
245	Ac-T/003	Procarbazine	Capsules	50mg	120	10x10		
246	Ac-I/001	5-F-L-uorouracil	Injection	250 mg- 5 ml Vial	120	1x10		
247	Ac-I/002	Actinomycin D	Injection	0.5mg, ampoul e/vial	120	1x10		
248	Ac-I/003	Alpha Interferon	Injection	3 million IU	60	1x10		
249	Ac-I/004	Bleomycin	Injection	15mg, vial	120	1x10		
250	Ac-I/005	Carboplatin	Injection A	150mg, vial	120	1x10		
251	Ac-I/006	Carboplatin	Injection B	450mg vial	120	1x10		
252	Ac-I/007	Cisplatin	Injection A	10mg/1 0ml vial	120	1x10		
253	Ac-I/008	Cisplatin	Injection B	50mg/v ial	120	1x10		
254	Ac-I/009	Cyclophosphami de	Injection	500 mg vial	120	1x10		
255	Ac-I/010	Cytosine arabioside	Injection A	100mg/ vial	120	1x10		
256	Ac-I/011	Cytosine arabioside	Injection B	500mg/ vial	120	1x10		

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257	Ac-I/012	Cytosine arabinoside	Injection C	1000mg/vial	120	1x10		
258	Ac-I/013	Dacarbazine	Injection	500mg/vial	120	1x10		
259	Ac-I/014	Daunorubicin	Injection	20mg vial	120	1x10		
260	Ac-I/015	Doxorubicin	Injection	10mg/5 ml vial	120	1x10		
261	Ac-I/016	Folinic acid	Injection	3mg/ml , apm./vial	240	1x10		
262	Ac-I/017	Gemcitabine hydrochloride	Injection A	200mg, vial	120	1x10		
263	Ac-I/018	Gemcitabine hydrochloride	Injection B	1gm, vial	120	1x10		
264	Ac-I/019	Ifosfamide	Injection	1gm/2 ml vial	120	1x10		
265	Ac-I/020	L-Asparaginase	Injection	5000 KU, vial	600	1x10		
266	Ac-I/021	Mercaptopurine	Injection	100mg/ml, vial	120	1x10		
267	Ac-T/004	Mercaptopurine	TAB	50mg	120	10x10		
268	Ac-I/022	Mesna	Injection	200mg, vial	60	1x10		
269	Ac-I/023	Methotrexate	Injection	50mg/ml, 2 ml glass vial	120	1x10		
270	Ac-I/024	Mitomycin-C	Injection	10mg, vial	120	1x10		
271	Ac-I/025	Oxaliplatin	Injection	50mg, vial	120	1x10		
272	Ac-I/026	Paclitaxel	Injection	30mg/5 ml	120	1x10		
273	Ac-I/027	Vinblastine sulphate	Injection	10mg, vial	120	1x10		
274	Ac-I/028	Vincristine	Injection	1mg/ml , vial	120	1x10		
275	Ac-T/005	Azathioprine	Tablet	50mg	120	10x10		
276	Ac-T/006	Busulphan	Tablet	2mg	120	10x10		
277	Ac-T/007	Chlorambucil	Tablet	2mg	120	10x10		
278	Ac-I/029	Cyclophosphamide	injection	200mg, 30ml vial	120	1x10		
279	Ac-I/030	Cyclophosphamide	Injection	500mg/vial	120	1x10		
280	Ac-T/008	Flutamide	Tablet	250mg	120	10x10		
281	Ac-T/009	Imatinib	Tablet A	100mg	120	10x10		
282	Ac-T/010	Imatinib	Tablet B	400mg	120	10x10		
283	Ac-T/011	Melphalan	Tablet A	2mg, 10 tab.	120	10x10		

				Pack				
284	Ac-T/012	Melphalan	Tablet B	5mg	120	10x10		
285	Ac-T/013	Mercaptopurine	Tablet	50mg	120	10x10		
286	Ac-T/014	Methotrexate	Tablet A	2.5mg	600	10x10		
287	Ac-T/015	Methotrexate	Tablet B	5mg	120	10x10		
288	Ac-T/016	Raloxifene	Tablet	60mg	120	10x10		
289	Ac-T/017	Tamoxifen	Tablet A	10mg	120	10x10		
290	Ac-T/018	Tamoxifen	Tablet B	20mg	120	10x10		
	Ap	Antiparkinsonism medicines						
291	Ap-T/001	Bromocriptine	Tablet A	1.25mg	120	10x10		
292	Ap-T/002	Bromocriptine	Tablet B	2.5mg	300	10x10		
293	Ap-T/003	Levodopa + Carbidopa	Tablet A	250mg + 25mg	6000	10x10		
294	Ap-T/004	Levodopa + Carbidopa	Tablet B	100mg + 10mg	6000	10x10		
295	Ap-T/005	Trihexyphenidyl	Tablet	2mg	3600	10x10		
	Cv	Cardiovascular medicine						
296	Cv-M/001	Digoxin	Elixir	0.05mg /ml	240	1x10		
297	Cv-I/001	Adenosine	Injection	3mg/ml , 2ml ampoul e	180	1x10		
298	Cv-I/002	Amiodarone	Injection	50mg/ ml	180	1x10		
299	Cv-I/003	Arginine Vasopression	Injection	40 IU/ml 1ml ampoul e	180	1x10		
300	Cv-I/004	Digoxin	Injection	0.25mg /ml	120	1x10		
301	Cv-I/005	Dobutamine	Injection	50mg/ ml, 5ml ampoul e	2400	1x10		
302	Cv-I/006	Dopamine hydrochloride	Injection	40mg/ ml, 5ml vial	3600	1x10		
303	Cv-I/007	Esmolol	Injection	10mg/ ml	120	1x10		
304	Cv-I/008	Isoprenaline	Injection	2mg/ml , 1 ml Ampoul e	120	1x10		
305	Cv-I/009	Labetolol	Injection	10mg/ ml	1800	1x10		
306	Cv-I/010	Mephentermine	Injection	30mg/ ml	120	1x10		

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				ml, 10ml Vial				
307	Cv-I/011	Metoprolol	Injection	1mg/ml , 5ml ampoul e	600	1x10		
308	Cv-I/012	Nitroglycerine	Injection	5mg/ml	300	1x10		
309	Cv-I/013	Sodium nitroprusside	Injection	50mg/5 ml	360	1x10		
310	Cv-I/014	Streptokinase	Injection A	750000 IU, per vial	60	1x10		
311	Cv-I/015	Streptokinase	Injection B	150000 0 IU, per vial	480	1x10		
312	Cv-I/016	tenecteplase	Injection	30 mg, Vial	60	1x10		
313	Cv-I/017	Verapamil	Injection	2.5mg/ ml	180	1x10		
314	Cv-T/001	Nifedipine	SR tab	10mg	360	10x10		
315	Cv-T/002	Nifedipine	SR tab	20mg	600	10x10		
316	Cv-T/003	Nitroglycerine	Sublingual tablet	0.5mg	1200	10x10		
317	Cv-T/004	Amiodarone	Tablet A	100mg	240	10x10		
318	Cv-T/005	Amiodarone	Tablet B	200mg	600	10x10		
319	Cv-T/006	Amlodipine	Tablet A	2.5mg	3000	10x10		
320	Cv-T/007	Amlodipine	Tablet B	5mg	6000	10x10		
321	Cv-T/008	Atenolol	Tablet	50mg	2400	14x10		
322	Cv-T/009	Atorvastatin	Tablet	10mg	1200	10x10		
323	Cv-T/010	Clopidogrel	Tablet	75mg	1200	10x10		
324	Cv-T/011	Digoxin	Tablet	0.25mg	1200	10x10		
325	Cv-T/012	Diltiazem	Tablet A	30mg	600	10x10		
326	Cv-T/013	Diltiazem	Tablet B	60mg	1200	10x10		
327	Cv-T/014	Enalapril	Tablet A	2.5mg	1800	10x10		
328	Cv-T/015	Enalapril	Tablet B	5mg	6000	10x10		
329	Cv-T/016	Isosorbide dinitrate	Tablet A	5mg	1200	10x10		
330	Cv-T/017	Isosorbide dinitrate	Tablet	10mg	3000	10x10		
331	Cv-T/018	Isosorbide mononitrate	Tablet	5mg	6000	10x10		
332	Cv-T/019	Isosorbide mononitrate	Tablet	10mg	6000	10x10		
333	Cv-T/020	Losartan	Tablet A	25mg	1200	10x10		
334	Cv-T/021	Losartan	Tablet B	50mg	1200	10x10		
335	Cv-T/022	Metoprolol	Tablet A	25mg	2400	10x10		
336	Cv-T/023	Metoprolol	Tablet B	50mg	2400	10x10		
337	Cv-T/024	Nifedipine	Tablet A	5mg,	2400	10x10		
338	Cv-T/025	Nifedipine	Tablet B	10mg	1200	10x10		

339	Cv-T/026	Propranolol	Tablet A	10mg	3000	10x10		
340	Cv-T/027	Propranolol	Tablet B	40mg	3000	10x10		
341	Cv-T/028	Spiroinolactone	Tablet	25mg	600	10x10		
342	Cv-T/029	Verapamil	Tablet A	40mg	1200	10x10		
343	Cv-T/030	Verapamil	Tablet B	80mg	3000	10x10		
344	Cv-T/031	Methyldopa	Tablet	250mg	1200	10x10		
345	Cv-T/032	Clonidine	tablet	0.1 mg	6000	10x10		
346	Cv-I/018	Noradrenaline	Injection	2 mg per ml, 2 ml ampoul e	12000	1x10		
347	Cv-I/019	Terlipressin	Injection	1mg/10 ml - 10ml vial	240	1x10		
	Dm	Dermatological Medicines						
		Dermatological Miscellaneous						
348	Dm-M/001	Calamine	Lotion	8%	480	1x10		
349	Dm-M/002	Dithranol	Ointment	0.1 to 2%, 20 gm Tube	120	1x10		
350	Dm-M/003	Gentian violet	Paint	0.50%	1800	1x10		
351	Dm-M/004	Gentian violet	Paint	1%	120	1x10		
352	Dm-M/005	Salicylic acid	Solution	5%	600	1x10		
353	Dm-M/006	Coal Tar	Solution	0.05	180	1x10		
354	Dm-M/007	Glycerine + Magnesium sulfate + Acriflavin	Solution	10%	120	1x10		
355	Dm-M/008	Glycerine solution	Solution , cream	10%	600	1x10		
356	Dm-M/009	Glycerine solution	Solution , cream	15%	600	1x10		
		Topical antibiotics						
358	Dm-M/010	Framycetin	Cream	0.005	600	1x10		
359	Dm-M/011	Fusidic acid	Ointment	2%	1200	1x10		
360	Dm-M/012	Fusidic acid	cream	10g	1200	1x10		
361	Dm-M/013	Fusidic acid	gel	15g	1200	1x10		
362	Dm-M/014	Fusidic acid + Hydrocortisone	cream	2%	1200	1x10		
363	Dm-M/015	Fusidic acid + Beclomethasone	cream	FA 2%	1200	1x10		
364	Dm-M/016	Fusidic Acid + (2%)	cream	10g	600	1x10		

		Betamethasone (1.2mg) 10.1%					
365	Dm-M/017		gel	0.75% or 0.8% or 1%	600	1x10	
		Metronidazole					
366	Dm-M/018	Nadifloxacin	cream	1%	3600	1x10	
		Antifungals Topicals				1x10	
367	Dm-M/019	Clotrimazole	cream,	1%	1200	1x10	
368	Dm-M/020	Clotrimazole	powder	1%	1200	1x10	
369	Dm-M/021	Clotrimazole	lotion	1%	1200	1x10	
370	Dm-M/022	Clotrimazole	gel	1%	1200	1x10	
371	Dm-M/023	Clotrimazole 1% + Beclomethasone 0.25%	cream,	10g	360	1x10	
372	Dm-M/024	Clotrimazole 1% + Beclomethasone 0.25%	cream,	20g	360	1x10	
373	Dm-M/025	Clotrimazole 1% + Selenium Sulfide 2.5%	lotion	60ml	600	1x10	
374	Dm-M/026	Ketoconazole 2%	Solution	60ml	600	1x10	
375	Dm-M/027	Ketoconazole 2%	Shampoo	100ml	600	1x10	
376	Dm-M/028	Ketoconazole 2%	cream,	20g	360	1x10	
377	Dm-M/029	Miconazole 2%	cream,	15g	1200	1x10	
378	Dm-M/030	Miconazole 2%	gel	20g	1200	1x10	
379	Dm-M/031	Sertaconazole 2%	cream,	10g	1200	1x10	
380	Dm-M/032	Sertaconazole 2%	cream,	15g	1200	1x10	
381	Dm-M/033	Sertaconazole 2%	lotion	15ml	1200	1x10	
382	Dm-M/034	Terbinafine	cream,	10g	1200	1x10	
383	Dm-M/035	Terbinafine	cream,	15g	1200	1x10	
384	Dm-M/036	Oxiconazole 1%	cream,	30g	600	1x10	
385	Dm-M/037	Oxiconazole 1%	lotion	30g	600	1x10	
386	Dm-M/038	Zinc Pyrithrone + ciclopirox	Solution	100ml	600	1x10	
		Antiparasitic					
387	Dm-M/039		lotion	1% 30ml	1200	1x10	
		Permethrin					
388	Dm-M/040		lotion	5% 30ml	1200	1x10	
		Permethrin					
389	Dm-M/041	Permethrin	cream	5% 30g	1200	1x10	
390	Dm-M/042	Lindane 1%	lotion	50ml	1200	1x10	
391	Dm-M/043	Lindane 1%	lotion	60ml	1200	1x10	

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392	Dm-M/044	Lindane 1%	lotion	100ml	1200	1x10		
		Topical Corticosteroids						
393	Dm-M/045	Beclomethasone 0.025%	cream	15g	600	1x10		
394	Dm-M/046	Beclomethasone 0.025%	lotion	15ml	600	1x10		
395	Dm-M/047	Beclomethasone 0.05% , 0.1%	cream	15g	600	1x10		
396	Dm-M/048	Beclomethasone 0.05% , 0.1%	gel	15g	600	1x10		
397	Dm-M/049	Beclomethasone 0.05% , 0.1%	gel	20g	360	1x10		
398	Dm-M/050	Beclomethasone 0.05% , 0.1%	lotion	20ml	360	1x10		
399	Dm-M/051	Beclomethasone 0.05% , 0.1%	lotion	30ml	360	1x10		
400	Dm-M/052	Beclomethasone 0.05% , 0.1%	lotion	50ml	360	1x10		
401	Dm-M/053	Betamethasone 0.05% +2n Sulphate 2.5%	cream	15g	600	1x10		
402	Dm-M/054	Clobetasol 0.05%	cream	10g	1200	1x10		
403	Dm-M/055	Clobetasol 0.05%	cream	15g	1200	1x10		
404	Dm-M/056	Clobetasol 0.05%	cream	30g	1200	1x10		
405	Dm-M/057	Clobetarol 0.05 + Gentamycin 0.1% + Miconazole 2%	cream	10g	600	1x10		
406	Dm-M/058	Clobetasone 0.05% + Gentamycin 0.1% + Miconazole 2%	cream	15g	600	1x10		
407	Dm-M/059	Clobetasone 0.05%	cream	10g	600	1x10		
408	Dm-M/060	Clobetasone 0.05%	cream	15g	600	1x10		
409	Dm-M/061	Fluocinolone 0.025%	cream	15g	360	1x10		
410	Dm-M/062	Fluocinolone 0.001%	Shampoo	100ml	360	1x10		
411	Dm-M/063	Fluocinolone 0.025%	lotion	15ml	360	1x10		
412	Dm-M/064	Fluticasone 0.05%	cream	10g	360	1x10		
413	Dm-M/065	Fluticasone 0.05%	cream	20g	600	1x10		

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414	Dm-M/066	Fluocinolone 0.05% + Mupirocin 2%	Ointment	5g	360	1x10		
415	Dm-M/067	Fluocinolone 0.05% + Mupirocin 2%	Ointment	10g	360	1x10		
416	Dm-M/068	Mometasone 0.1%	cream	15g	1200	1x10		
417	Dm-M/069	Mometasone 0.1%	Ointment	15g	1200	1x10		
418	Dm-M/070	Mometasone 0.1%	solution	15ml	1200	1x10		
419	Dm-M/071	Mometasone 0.1%	lotion	15ml	1200	1x10		
420	Dm-M/072	Triamcinolone 0.025%	Ointment	15g	360	1x10		
421	Dm-M/073	Triamcinolone 0.05%	Ointment	15g	360	1x10		
422	Dm-M/074	Triamcinolone 0.1%	Ointment	15g	360	1x10		
423	Dm-M/075	Desonide 0.05%	cream	10g	600	1x10		
424	Dm-M/076	Desonide 0.05%	lotion	30ml	600	1x10		
		Other Topicals						
425	Dm-M/077	Salicylic acid 3%	Ointment	25g	600	1x10		
426	Dm-M/078	Salicylic acid 6%	Ointment	25g	600	1x10		
427	Dm-M/079	Salicylic acid 2%	lotion	60ml	600	1x10		
428	Dm-M/080	Tretinoin 0.025%	cream	20	1200	1x10		
429	Dm-M/081	Tretinoin 0.05%	cream	20	1200	1x10		
430	Dm-M/082	Urea 10%	cream	50g	360	1x10		
431	Dm-M/083	Urea 20%	cream	50g	360	1x10		
432	Dm-M/084	Urea 3%	lotion	150ml	360	1x10		
433	Dm-M/085	Urea 10%	lotion	150ml	360	1x10		
434	Dm-M/086	Lactic Acid 12%	cream	30g	360	1x10		
435	Dm-M/087	Lactic Acid 12%	cream	75g	360	1x10		
436	Dm-M/088	Lactic Acid 12%	cream	100g	360	1x10		
437	Dm-M/089	Lactic Acid 12%	Ointment	30g	360	1x10		
438	Dm-M/090	Lactic Acid 12%	lotion	100ml	360	1x10		
439	Dm-M/091	iron Oxide 8.5%	cream	20g	360	1x10		
440	Dm-M/092	iron Oxide 8.5%	lotion	20ml	360	1x10		
441	Dm-M/093	Calamine 8% + Liquid Paraffin 10%	lotion	120ml	2400	1x10		
442	Dm-M/094	Liquid paraffin	Daily	100ml	2400	1x10		
443	Dm-M/095	Liquid paraffin	Preparation	100ml	2400	1x10		
444	Dm-M/096	Tacrolimus 0.03%	Ointment	10g	1200	1x10		
445	Dm-M/097	Tacrolimus 0.1%	Ointment	10g	1200	1x10		
446	Dm-M/098	Pimecrolimus	Ointment	10	600	1x10		

	Da	Diagnostic agents						
447	Da-M/001	Fluorescein	Eye Drops	1%, 5ml vial with sterilized dropper or squeeze vial	60	1x10		
448	Da-I/001	Calcium ipodate	Injection	3gm	0	1x10		
449	Da-I/002	Meglumine iothalamate	Injection	60% w/v (Iodine 280mg/ml), 50 ml vial	600	1x10		
450	Da-I/003	Sodium Iothalamate	Injection	70% w/v (Iodine = 420 mg/ml), 50 ml vial	600	1x10		
451	Da-I/004	Sodium meglumine diatrizoate	Injection	60% w/v, 50,100 ml vial (Iodine =292mg/ml), 76% w/v (Iodone = 370mg/ml)	600	1x10		
452	Da-M/002	Iopromide	Solution	300-370mg/ml, 50,100 ml vial	300	1x10		
453	Da-M/003	Propyliodone	Oily, suspension		0	1x10		
454	Da-I/005	Ioversol	Injection, solution	300-375 mg/ml, 50,100 ml vial	300	1x10		
455	Da-M/004	Meglumine iotroxate	Solution	5-8 gm iodine in 100 -	600	1x10		

				250ml 50,100 ml vial				
456	Da-I/006	Iohexol	injection,sol ution	300- 350 mg/ml 20,50,1 00 ml vial	600	1x10		
457	Da-M/005	Barium sulfate	Suspension	100% w/v, 1 Ltr	600	1x10		
458	Da-M/006	Barium sulfate	Suspension	250% w/v, 1 Ltr	240	1x10		
459	Da-M/007	Barium sulfate	Suspension	100% w/v, 250% w/v	120	1x10		
	Di	Disinfectants and antiseptics						
458	Di-M/001	Potassium permanganate	Crystals for solution	value in gms.	6000	1x10		
459	Di-M/002	Zinc oxide	Dusting Powder	5%	1200	1x10		
460	Di-M/003	Zinc oxide	Dusting Powder	25%/2 5gm	1200	1x10		
461	Di-M/004	Povidone Iodine	Eye Drops	0.006	120	1x10		
462	Di-M/005	Povidone Iodine	Eye Ointment	0.006	60	1x10		
463	Di-I/001	Hypochlorite solution	Injection	10%	120	1x10		
464	Di-M/006	Povidone Iodine	Mouth gargles	50ml	1200	1x10		
465	Di-M/007	Povidone Iodine	Mouth gargles	100ml	120	1x10		
466	Di-M/008	Povidone Iodine	Ointment	0.5%, 15gm tube	1800	1x10		
467	Di-M/009	Bleaching powder	Powder (Contains not less than 30% w/w of available chlorine)	5kg bags	600	1x10		
468	Di-M/010	Acridflavin + Glycerine	Solution	1ltr can	60	1x10		
469	Di-M/011	Cetrimide	Solution	0.2	240	1x10		

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470	Di-M/012	Chlorhexidine	Solution	0.5%, 10ml/1 00ml bottle	1200	1x10		
471	Di-M/013	Ethyl alcohol	Solution	0.7	120	1x10		
472	Di-M/014	Glutaraldehyde	Solution	0.02	60	1x10		
473	Di-M/015	Hydrogen peroxide	Solution	0.06	1200	1x10		
474	Di-M/016	Povidone Iodine	Solution A	5%, 500ml bottle	1200	1x10		
475	Di-M/017	Povidone Iodine	Solution B	7.50%	120	1x10		
476	Di-M/018	Povidone Iodine	Solution C	10%, 100ml/ 500ml/ 1litre	120	1x10		
477	Di-M/019	Formaldehyde	Solution (Dilute 34 ml with water to produce 100mg)	10%	1200	1x10		
478	Di-T/001	Formalin	Tablet		7200	10x10		
479	Di-M/020	Formalin	solution	10%	120	1x10		
480	Di-F/001	Benzoine compund	Tincture	200ml bottle	120	1x10		
481	Di-M/021	Chlorhexidine	solution	5%	600	1x10		
482	Di-M/022	Isopropyl Alcohol	solution	70%	600	1x10		
483	Di-M/023	surgical Spirit	solution	94%	600	1x10		
	Du	Diuretics						
484	Du-I/001	Acetazolamide	Injection	500mg vial	600	1x10		
485	Du-I/002	Furosemide	Injection	10mg/ ml, 2ml ampoul e	2400	1x10		
486	Du-I/003	Mannitol	Injection A	10%	3000	1x10		
487	Du-I/004	Mannitol	Injection B	20%, 100 ml Bottle	120	1x10		
488	Du-T/001	Acetazolamide	Tablet	250mg	3000	10x10		
489	Du-T/002	Furosemide	Tablet	40mg	3000	10x10		
490	Du-T/003	Hydrochlorothia zide	Tablet A	25mg	600	10x10		
491	Du-T/004	Hydrochlorothia zide	Tablet B	50mg	120	10x10		
492	Du-T/005	Metolazone	Tablet	2.5mg	240	10x10		
493	Du-T/006	Metolazone	Tablet	5mg	600	10x10		
494	Du-I/005	Milrinone	Injection	10mg/1	120	1x10		

				0ml, 10ml vial				
495	Du-T/007	Torsemide	Tablet	10mg	1200	10x10		
496	Du-T/008	Torsemide	Tablet	5mg	2400	10x10		
	En	ENT						
497	En-M/001	Chloramphenico 15%/Beclometha sone.025%/Clot rimazole1%	Ear drops	10ml bottle	600	1x10		
498	En-M/002	Ciprofloxacin+D examethasone	Ear drops	10ml bottle	600	1x10		
499	En-I/001	Dimenhydranate	injection	50mg	1200	1x10		
500	En-M/003	Xylometazoline Nasal Drop	Nasal Solution 0.1%	0.1% - 10ml vial.	600	1x10		
501	En-M/004	Xylometazoline Nasal Drop	Nasal Solution 0.5%	0.05% - 10ml vial	600	1x10		
502	En-M/005	Fluticasone propionate	Nasal spray	50mcg/ puff, 7ml bottle	600	1x10		
503	En-M/006	fucidic acid	ointment		120			
504	En-M/007	Mometasone	Nasal spray	50mcg, 120 metere d dose	240	1x10		
505	En-L/001	Ambroxol	Syrup	60mg/5 0ml	600	1x10		
506	En-L/002	Lactulose	Syrup	50mg	2400	1x10		
507	En-L/003	Lactulose	Syrup B	100mg	2400	1x10		
508	En-T/001	Ambroxol	Tablet	30	1200	10x10		
509	En-T/002	Ambroxol	Tablet	60	120	10x10		
510	En-T/003	Cinnarzine	Tablet	25mg	2400	10x10		
511	En-T/004	Dimenhydranate	Tablet	50mg	1200	10x10		
512	En-T/005	Itraconazole	Tablet	100mg	2400	10x10		
513	En-T/006	Prochloroperazi ne	Tablet	5mg	1200	10x10		
514	En-M/008	Ichthamol Glycerine	Topical solution	5mg	240	1x10		
515	En-M/009	Beclomethasone ointment	ointment	5gm	1200	1x10		
516	En-T/007	Betahistine Hydrochloride	tablet	8mg	360	10x10		
517	En-T/008	Betahistine Hydrochloride	tablet	16mg	360	10x10		
518	En-M/010	xylocain Jelly	lubricant	2%	600	1x10		
519	En-M/011	Wax dissolving ear drops	ear drops		600	1x10		

	Gi	Gastrointestinal medicines						
520	Gi-T/001	Omeprazole	Capsule A	20mg	14400	10x10		
521	Gi-T/002	Omeprazole	Capsule B	40mg	120	10x10		
522	Gi-M/001	Ispaghula	Granules	100g	1200	1x10		
523	Gi-I/001	Dicylomine hydrochloride	Injection	10mg/ml	1200	1x10		
524	Gi-I/002	Hyoscine butyl bromide	Injection	20mg/ml	120	1x10		
525	Gi-I/003	Metoclopramide	Injection	5mg/ml , 2ml ampoule	2400	1x10		
526	Gi-I/004	Ondansetron	Injection	2mg/ml , 2ml/4 ml ampoule	5400	1x10		
527	Gi-I/005	Pantoprazole	Injection	40mg	10200	1x10		
528	Gi-I/006	Ranitidine	Injection	50mg/2 ml	8400	1x10		
529	Gi-L/001	Antacid (Alumunium hydroxide + Magnesium hydroxide)	Liquid/gel	170ml bottle	4200	1x10		
530	Gi-T/003	Antacid (Alumunium hydroxide + Magnesium hydroxide)	Tab	50 pack strips	4200	1x10		
531	Gi-M/002	Oral Rehydration Salts (ORS)	Powder for solution	35 gms pack.	6000	1x10		
532	Gi-M/003	Bisacodyl	Suppository	5mg, 10mg 1x5 pkt	600	(1x5 pkt) 1x10		
533	Gi-L/002	Domperidone	Syrup	1mg/ml , 30ml	2400	1x10		
534	Gi-L/003	Lactulose	Syrup	3.1- 3.7gm/ 5ml	1800	1x10		
535	Gi-L/004	Metoclopramide	Syrup	5mg/5 ml	600	1x10		
536	Gi-L/005	Ondansetron	Syrup	2mg/5 ml, 30ml	1200	1x10		
537	Gi-T/004	Antacid (Alumunium hydroxide +	Tablet	tablet	5400	10x10		

		Magnesium hydroxide)						
538	Gi-T/005	5-Amino salicylic acid	Tablet	400mg	120	10x10		
539	Gi-T/006	Bisacodyl	Tablet	5mg	1200	10x10		
540	Gi-T/007	Dicylomine hydrochloride	Tablet	10mg	3600	10x10		
541	Gi-T/008	Domperidone	Tablet	10mg	4800	10x10		
542	Gi-T/009	Hyoscine butyl bromide	Tablet	10mg	600	10x10		
543	Gi-T/010	Metoclopramide	Tablet	10mg	4800	10x10		
544	Gi-T/011	Ondansetron	Tablet A	4mg	4800	10x10		
545	Gi-T/012	Ondansetron	Tablet B	8mg	120	10x10		
546	Gi-T/013	Ranitidine	Tablet A	150mg	10800	10x10		
547	Gi-T/014	Ranitidine	Tablet B	300mg	120	10x10		
548	Gi-T/015	Sucralfate	Tablet	1gm	3000	10x10		
549	Gi-T/016	Sulfasalazine	Tablet	500mg	600	10x10		
550	Gi-L/006	Liquid paraffin	liquid	500ml bottle	6600	1x10		
	Ho	Hormones, other endocrine medicines and contraceptives						
551	Ho-T/001	Danazol	Capsule	50mg	600	10x10		
552	Ho-T/002	Danazol	Capsule	100mg	120	10x10		
553	Ho-T/003	Testosterone	Capsule	40mg (As undecanoate)	60	10x10		
554	Ho-M/001	Betamethasone	Cream/Ointment	0.0005, 15g tube	120	1x10		
555	Ho-M/002	Leuprolide hydrochloride 3.75	Depot	tablet	120	1x10		
556	Ho-M/003	Prednisolone acetate	Eye Drops	0.01, 5ml	180	1x10		
557	Ho-M/004	Prednisolone Sodium Phosphate	Eye Drops	0.5%, 5ml	180	1x10		
558	Ho-I/001	Betamethasone	Injection	4mg/ml	660	1x10		
559	Ho-I/002	Glucagon	Injection	1mg/ml, 1 ml Vial	120	1x10		
560	Ho-I/003	Medroxyprogesterone acetate	Injection	Depot preparation	600	1x10		
561	Ho-I/004	Methyl prednisolone	Injection	40mg/ml, vial	1800	1x10		
562	Ho-I/005	Nandrolone decanoate	Injection	50mg	720	1x10		

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563	Ho-I/006	NPH/Lente Insulin	Injection	40 IU/ml	1200	1x10		
564	Ho-I/007	Premix Insulin	Injection	40 IU/ml (30:70), Vial	1200	1x10		
565	Ho-I/008	Regular insulin (Soluble)	Injection	40 IU/ml	3000	1x10		
566	Ho-I/009	Testosterone	Injection	25mg/ml (As appropriate)	120	1x10		
567	Ho-M/005	Hormone (Levonorgestrel) releasing IUD	IUD	medicated IUD	60	1x10		
568	Ho-M/006	IUD containing Copper	IUD	cu380A	60	1x10		
569	Ho-M/007	Iodine solution	Solution	8mg/5 ml	60	1x10		
570	Ho-T/004	Carbimazole	Tablet A	5mg	300	10x10		
571	Ho-T/005	Carbimazole	Tablet B	10mg	120	10x10		
572	Ho-T/006	Clomiphene citrate	Tablet A	50mg	120	10x10		
573	Ho-T/007	Clomiphene citrate	Tablet B	100mg	120	10x10		
574	Ho-T/008	Ethinyl estradiol + Norethisterone	Tablet	0.035mg + 1mg	60	10x10		
575	Ho-T/009	Ethinylestradiol	Tablet A	0.01mg	120	10x10		
576	Ho-T/010	Ethinylestradiol	Tablet B	0.05mg	120	10x10		
577	Ho-T/011	Ethinylestradiol + Levonorgestrel	Tablet	0.03mg + 0.15mg	120	10x10		
578	Ho-T/012	Glibenclamide	Tablet A	2.5mg	120	10x10		
579	Ho-T/013	Glibenclamide	Tablet B	5mg	120	10x10		
580	Ho-T/014	Iopanoic acid	Tablet	500mg	120	10x10		
581	Ho-T/015	Levothyroxine	Tablet A	25µg	7200	10x10		
582	Ho-T/016	Levothyroxine	Tablet A	50µg	7200	10x10		
583	Ho-T/017	Levothyroxine	Tablet B	100µg	120	10x10		
584	Ho-T/018	Medroxyprogesterone acetate	Tablet A	5mg.	1200	10x10		
585	Ho-T/019	Medroxyprogesterone acetate	Tablet B	10mg	120	10x10		
586	Ho-T/020	Metformin	Tablet	500mg	12000	10x10		
587	Ho-T/021	Norethisterone acetate	Tablet	5mg	1200	10x10		
588	Ho-T/022	Prednisolone	Tablet A	5mg,	4800	10x10		
589	Ho-T/023	Prednisolone	Tablet B	10mg	120	10x10		
590	Ho-T/024	Propylthiouracil	Tablet A	50mg	3000	10x10		
591	Ho-T/025	Propylthiouracil	Tablet B	100mg	120	10x10		
592	Ho-T/026	Tamsulosin/Fin	Tablet	5mg+0.	1200	10x10		

		asteride		4mg				
593	Ho-T/027	Calcitriol	Capsule	0.25mcg	1800	10x10		
594	Ho-I/010	HCG	Injection	5000 Unit	120	1x10		
595	Ho-I/011	Hydrocortosone acetate	Injection	100mg	1200	1x10		
596	Ho-M/008	Mometasone	Nasal spray	50mcg/puff, 7ml bottle	600	1x10		
597	Ho-I/012	Octreotide	Injection	100mcg/ml, 1ml amp	240	1x10		
598	Ho-I/013	Octreotide	Injection	50mcg/ml, 1ml amp	240	1x10		
599	Ho-I/014	Carboprost	Injection	0.25mg/ml, 2ml ampoule	240	1x10		
600	Ho-I/015	Isoxsuprine	Injection	5mg/ml	120	1x10		
601	Ho-I/016	Methyl ergometrine	Injection	0.2mg/ml, 1 ampoule	1200	1x10		
602	Ho-I/017	Oxytocin	Injection A	5 IU/ml	2400	1x10		
603	Ho-I/018	Oxytocin	Injection B	10 IU/ml, 1 ampoule	120	1x10		
604	Ho-T/028	Isoxsuprine	Tablet A	20mg	360	10x10		
605	Ho-T/029	Isoxsuprine	Tablet B	40mg	120	10x10		
606	Ho-T/030	Methyl ergometrine	Tablet	0.125mg	120	10x10		
607	Ho-T/031	Mifepristone	Tablet	200mg	360	10x10		
608	Ho-T/032	Misoprostol	Tablet	100µg	1200	10x10		
609	Ho-I/020	Valethamide Bromide	injection	8mg/ml amp.	120	1x10		
	Im	Immunologicals						
610	Im-I/001	Rabies Immunoglobulin	Injection	150 IU/ml	180	1x10		
611	Im-I/002	Tetanus toxoid	Injection	5ml ampoule	2400	1x10		
	Re	Medicines acting on the respiratory tract						

612	Re-M/001	Budesonide	Nebulizer suspension	0.25mg/ml	2400	1x10		
613	Re-T/001	Caffeine citrate	Tablets	30mg	300	10x10		
614	Re-M/002	Beclomethasone	Inhalation A	50µg, per dose	120	1x10		
615	Re-M/003	Beclomethasone	Inhalation B	250µg per dose	120	1x10		
616	Re-M/004	Ipratropium bromide	Inhalation	20µg/dose, 200 MDI	7800	1x10		
617	Re-M/005	Salmeterol + Fluticasone	Inhalation	25µg + 125µg/dose	1800	1x10		
618	Re-I/001	Aminophylline	Injection	25mg/ml, 10ml ampoule	300	1x10		
619	Re-I/002	Hydrocortisone sodium succinate	Injection	100mg, Vial	3600	1x10		
620	Re-I/003	Terbutaline	Injection	0.5mg/ml, 1ml ampoule	240	1x10		
621	Re-M/006	Ipratropium bromide	Nebulizer solution	250µg/ml, 15 ml vial	1200	1x10		
622	Re-M/007	Salbutamol	Respiratory solution	5mg/ml	4800	1x10		
623	Re-L/001	Codeine phosphate	Syrup	15mg/5 ml	600	1x10		
624	Re-L/002	Salbutamol	Syrup	2mg/5 ml	600	1x10		
625	Re-T/002	Codeine phosphate	Tablet	10mg	120	10x10		
626	Re-T/003	Dextromethorphan	Tablet	30mg	1200	10x10		
627	Re-T/004	Etiophylline + Theophylline	Tablet	77mg + 23mg	2400	10x10		
628	Re-T/005	Salbutamol	Tablet A	2mg	600	10x10		
629	Re-T/006	Salbutamol	Tablet B	4mg	120	10x10		
630	Re-T/007	Terbutaline	Tablet	2.5mg	600	10x10		
	BI	Medicines affecting the blood						

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631	BI-I/001	Albumin	Injection A	5% in amp/vial	600	1x10		
632	BI-I/002	Albumin	Injection B	20% in 50ml	120	1x10		
633	BI-I/003	Cryoprecipitate	Injection	Amp.	120	1x10		
634	BI-I/004	Dextran 40	Injection	0.1	600	1x10		
635	BI-I/005	Dextran 70	Injection	0.06	600	1x10		
636	BI-I/006	Enoxaparin	Injection A	40mg in amp./vial	600	1x10		
637	BI-I/007	Enoxaparin	Injection B	60mg, Vial / PFS	120	1x10		
638	BI-I/008	Filgastrim	Injection	1 ml vial	120	1x10		
639	BI-I/009	Heparin sodium	Injection A	1000 IU/ml, 1ml ampoule	1800	1x10		
640	BI-I/010	Heparin sodium	Injection B	5000 IU/ml, 1ml ampoule	120	1x10		
641	BI-I/011	Hydroxyethyl starch	Injection	6%, 500ml	1800	1x10		
642	BI-I/012	Iron Dextran	Injection	50mg iron/ml	1200	1x10		
643	BI-I/013	Polygeline	Injection	0.035	120	1x10		
644	BI-I/014	Tranexamic acid	Injection	500mg/5ml, 5ml ampoule	1800	1x10		
645	BI-I/015	Factor IX Complex (Coagulation Factors II, VII, IX, X)	Injection (Dried)	600 IU	120	1x10		
646	BI-I/016	Factor VIII concentrate	Injection (Dried)	250 IU	120	1x10		
647	BI-I/017	Factor VIII concentrate	Injection (Dried)	500 IU	120	1x10		
648	BI-I/018	Factor VIII concentrate	Injection (Dried)	1000 IU	120	1x10		
649	BI-M/001	Cotrimoxazole	Suspension	40mg + 200mg/5ml	600	1x10		
650	BI-L/001	Ferrous sulfate/fumarate	Syrup	20mg element	1200	1x10		

				al iron/ml				
651	BI-T/001	Cotrimoxazole	Tablet	80mg + 400mg	1800	10x10		
652	BI-T/002	Cotrimoxazole	Tablet	160mg + 800mg	120	10x10		
653	BI-T/003	Ferrous sulfate/fumarate	Tablet	Tablets equival ent to 60 mg element al iron	10200	10x10		
654	BI-T/004	Folic acid	Tablet	5mg	6000	10x10		
655	BI-T/005	Warfarin sodium	Tablet	5mg	14400	10x10		
656	BI-T/006	Tranexamic acid	Tablet	500mg	3000	10x10		
657	BI-I/019	Antithrombin III (Human)	Injection	amp.	120	1x10		
658	BI-T/007	Ethamsylate	Tablets A	250mg,	1200	10x10		
659	BI-T/008	Ethamsylate	Tablets B	500mg	1200	10x10		
660	BI-I/020	Ethamsylate	Injection	250mg, 2ml	1200	1x10		
661	BI-F/001	Haemacel	infusion	500ml bottle	1200	1x10		
662	BI-I/021	Hemocoagulase	injection	2ml amp	600	1x10		
663	BI-I/022	Hemodialysis fluid with dextrose	infusion	5000ml	360	1x10		
664	BI-I/023	Low molecular weight heparin	injection	Amp.	2400	1x10		
665	BI-I/024	Tenectaplast	Injection	30mg	300	1x10		
666	BI-M/002	Thrombophobe ointment	ointment	20g	2400	1x10		
	Cn	Medicines for nervous system disorders						
667	Cn-T/001	Fluoxetine	Capsule	20mg	2400	10x10		
668	Cn-M/001	Haloperidol	Depot Preparation	50mg/ ml	120	1x10		
669	Cn-I/001	Chlorpromazine	Injection	25mg/ ml	60	1x10		
670	Cn-I/002	Diazepam	Injection	5mg/ml , 2ml ampoul e	6000	1x10		
671	Cn-I/003	Haloperidol	Injection	5mg/ml , 1ml ampoul e	2400	1x10		

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672	Cn-I/004	Phenobarbitone	Injection	200mg/ ml, 1ml ampoul e	1200	1x10		
673	Cn-M/002	Diazepam	Suppository	5mg	120	1x10		
674	Cn-L/001	Diazepam	Syrup	2mg	1200	1x10		
675	Cn-L/002	Phenobarbitone	Syrup	20mg/5 ml, 60ml bottle	240	1x10		
676	Cn-T/002	Alprazolam	Tablet	0.25mg	6600	10x10		
677	Cn-T/003	Alprazolam	Tablet	0.50mg	120	10x10		
678	Cn-T/004	Amitriptyline	Tablet	25mg	240	10x10		
679	Cn-T/005	Chlorpromazine	Tablet	100mg	3000	10x10		
680	Cn-T/006	Diazepam	Tablet	5mg	3600	10x10		
681	Cn-T/007	Dihydroergotam ine	Tablet	1mg	600	10x10		
682	Cn-T/008	Haloperidol	Tablet	5mg	2400	10x10		
683	Cn-T/009	Imipramine	Tablet	25mg	600	10x10		
684	Cn-T/010	Lithium carbonate	Tablet	300mg	3600	10x10		
685	Cn-T/011	Olanzapine	Tablet	5mg	3000	10x10		
686	Cn-T/012	Phenobarbitone	Tablet A	30mg	600	10x10		
687	Cn-T/013	Phenobarbitone	Tablet B	60mg	120	10x10		
688	Cn-T/014	Risperdione	Tablet	2mg	120	10x10		
689	Cn-T/015	Clozapine	Tablet	100mg	120	10x10		
690	Cn-L/003	Chloral hydrate	Syp	60ml	240	1x10		
691	Cn-T/016	Clobazam	tab	5mg	2400	10x10		
692	Cn-T/017	Levetiracetam	tab	500mg	600	10x10		
693	Cn-F/001	Levetiracetam	solution	100mg/ ml 300ml bottle	600	1x10		
	Mr	Muscle relaxants						
694	Mr-I/001	Atracurium besylate	Injection	10mg/ ml	120	1x10		
695	Mr-I/002	Neostigmine	Injection	0.5mg/ ml	60	1x10		
696	Mr-I/003	Succinyl choline	Injection	50mg/ ml, 1ml Amp (10 ampoul e)	600	1x10		
697	Mr-I/004	Vecuronium	Injection	2mg/ml - 2ml vial	1200	1x10		
698	Mr-T/001	Neostigmine	Tablet	15mg	120	10x10		
699	Mr-I/005	Pancuronium	injection	2mg/ml	600	1x10		

				2ml amp.				
700	Mr-I/006	Rocuronium bromide	injection	5mg/ml 5ml aml.	120	1x10		
701	Mr-M/001	Succinyl choline	Powder for suspension	200g pack	600	1x10		
702	Mr-M/002	Surfactants		6.38 % w/w x 50g	360	1x10		
	Op	Ophthalmological						
703	Op-M/001	Betaxolol hydrochloride	Eye Drops A	0.25%, 5ml	120	1x10		
704	Op-M/002	Betaxolol hydrochloride	Eye Drops B	0.5%, 5ml	120	1x10		
705	Op-M/003	Carboxymethyl cellulose	Eye Drops	0.05	180	1x10		
706	Op-M/004	Homatropine	Eye Drops	2%, 5ml	120	1x10		
707	Op-M/005	Phenylephrine	Eye Drops	4%/5% ,5ml	120	1x10		
708	Op-M/006	Pilocarpine	Eye Drops	2%,	240	1x10		
709	Op-M/007	Pilocarpine	Eye Drops	4%	240	1x10		
710	Op-M/008	Timolol	Eye Drops A	0.25%	120	1x10		
711	Op-M/009	Timolol	Eye Drops B	0.50%	120	1x10		
712	Op-M/010	Tropicamide	Eye Drops	0.5%, 5ml drops	120	1x10		
713	Op-M/011	Tropicamide + Phenylephrine	Eye Drops	0.8% + 5%, 5ml	120	1x10		
714	Op-M/012	Tropicamide	Eye Drops	0.5%, 5ml drops	120	1x10		
715	Op-I/001	Methyl cellulose	Injection	0.02	120	1x10		
716	Op-I/002	Phenylephrine	Injection	4%, 5ml	120	1x10		
	Va	Vaccine, Sera and immunomodulators						
717	Va-T/001	Cyclosporine	Capsule A	10mg,	60	10x10		
718	Va-T/002	Cyclosporine	Capsule B	20mg	120	10x10		
719	Va-T/003	Cyclosporine	Capsule C	50mg,	120	10x10		
720	Va-T/004	Cyclosporine	Capsule D	100mg	120	10x10		
721	Va-I/001	Cyclosporine	Concentrate for Injection	100mg/ ml, apm./vi al	120	1x10		
722	Va-I/002	Anti-D immunoglobulin (Human)	Injection	300µg	180	1x10		
723	Va-I/003	Anti-gas gangrene serum	Injection		360	1x10		

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724	Va-I/004	Anti-T-etanus human immunoglobulin	Injection A	250 IU	600	1x10		
725	Va-I/005	Anti-T-etanus human immunoglobulin	Injection B	500 IU	120	1x10		
726	Va-I/006	BCG Vaccine	Injection	vial	120	1x10		
727	Va-I/007	Diphtheria Antitoxin	Injection	10000 IU, vial	240	1x10		
728	Va-I/008	DPT Vaccine	Injection		600	1x10		
729	Va-I/009	Hepatitis B Vaccine	Injection	20mcg, 1ml Vial	240	1x10		
730	Va-I/010	Measles Vaccine	Injection	Vial	120	1x10		
731	Va-I/011	Rabies Vaccine	Injection		360	1x10		
732	Va-I/012	Tuberculin Purified Protein Derivative	Injection	1 TU	1200	1x10		
733	Va-I/013	Tuberculin Purified Protein Derivative	Injection	5 TU	120	1x10		
734	Va-I/014	Anti-snake venom	Injection (Polyvalent solution/Lyophilized Polyvalent Serum)	10ml	2400	1x10		
735	Va-M/001	Oral Poliomyelitis Vaccine	Solution	0.5ml vial	120	1x10		
736	Va-M/002	Anti-snake venom	Specific	10ml ampoule	120	1x10		
737	Va-I/015	Td Vaccine	injection	0.5ml ampoule	1200	1x10		
	Vi	Vitamins and Minerals						
738	Vi-T/001	Vitamin D (Ergocalciferol)	Capsule A	0.25mg	6000	10x10		
739	Vi-T/002	Vitamin D (Ergocalciferol)	Capsule B	1mg	120	10x10		
740	Vi-I/001	Calcium gluconate	Injection	100mg/ml in 10ml ampoule	8400	1x10		
741	Vi-I/002	Cyanocobolamin	Injection	1mg/ml	1200	1x10		
742	Vi-I/003	Phytomenadione	Injection	10mg/ml, 5ml ampoule	3000	1x10		
743	Vi-I/004	Vitamin A	Injection	50000I	120	1x10		

				U/ml				
744	Vi-I/005	Vitamin K1	Injection	1mg/ml , 1ml ampoul e	4200	1x10		
745	Vi-L/001	Ascorbic acid	Syrup	20mg/5 ml	1200	1x10		
746	Vi-T/003	Ascorbic acid	Tablet A	100mg	2400	10x10		
747	Vi-T/004	Ascorbic acid	Tablet B	500mg	120	10x10		
748	Vi-T/005	Calcium carbonate	Tablet A	250mg,	21600	10x10		
749	Vi-T/006	Calcium carbonate	Tablet B	500mg	120	10x10		
750	Vi-T/007	Multivitamin	Tablet/cap		46800	10x10		
751	Vi-T/008	Nicotinamide	Tablet	50mg	600	10x10		
752	Vi-T/009	Pyridoxine	Tablet	25mg	2400	10x10		
753	Vi-T/010	Riboflavin	Tablet	5mg	120	10x10		
754	Vi-T/011	Thiamine	Tablet	100mg	1800	10x10		
755	Vi-T/012	Vitamin C	Tablet A	100mg	7800	10x10		
756	Vi-T/013	Vitamin C	Tablet B	500mg	120	10x10		
757	Vi-T/014	Vitamin A	Tablet/Caps ule A	5000 IU	3600	10x10		
758	Vi-T/015	Vitamin A	Tablet/Caps ule B	50000I U	120	10x10		
759	Vi-T/016	Vitamin A	Tablet/Caps ule C	10000I U	120	10x10		
	Ms	Miscellaneous						
760	Ms-M/001	Cyclopentolate hydrochloride	Eye Drops	0.01	120	1x10		
761	Ms-M/002	Dinoprostone	Gel A	0.5gm	120	1x10		
762	Ms-M/003	Dinoprostone	Gel B	3gm	120	1x10		
763	Ms-M/004	Oxygen	Inhalation	Cylinde r	60	1x10		
764	Ms-I/001	Alprostadil	Injection	500µg	60	1x10		
765	Ms-I/002	Calcium chloride	Injection	Amp.	2400	1x10		
766	Ms-I/003	Glucose	Injection A	5%	12000	1x10		
767	Ms-I/004	Glucose	Injection B	10%	120	1x10		
768	Ms-I/005	Glucose	Injection C	15%	120	1x10		
769	Ms-I/006	Glucose	Injection D	25%	120	1x10		
770	Ms-I/007	Glucose	Injection E	50%	120	1x10		
771	Ms-I/008	Glucose with sodium chloride	Injection	5% + 0.9%	18000	1x10		
772	Ms-I/009	Glycopyrrolate	Injection	0.2mg/ ml, 1ml ampoul e	600	1x10		
773	Ms-I/010	Hypertonic	Injection	0.03	600	1x10		

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		saline						
774	Ms-I/011	N/2 saline	Injection	300ml bottle	14400	1x10		
775	Ms-I/012	N/5 saline	Injection	300ml bottle	4200	1x10		
776	Ms-I/013	Normal saline	Injection	0.9%, 100ml/ 500ml bottle	30000	1x10		
777	Ms-I/014	Potassium chloride	Injection	11.2% solution , 20 ml ampoul e	7800	1x10		
778	Ms-I/015	Protamin sulfate	Injection	10mg/ ml	120	1x10		
779	Ms-I/016	Pyridostigmine	Injection	1mg/ml	120	1x10		
780	Ms-I/017	Ringer lactate	Injection	300ml bottle	18000	1x10		
781	Ms-I/018	Sodium bicarbonate	Injection	10 ml Amp	4200	1x10		
782	Ms-I/019	Tham	Injection	Amp.	600	1x10		
783	Ms-I/020	Water for injection	Injection A	2ml, 2ml ampoul e	36000	1x10		
784	Ms-I/021	Water for injection	Injection B	5ml ampoul e	120	1x10		
785	Ms-I/022	Water for injection	Injection C	10ml ampoul e	120	1x10		
786	Ms-M/005	Glucose	Oral Powder	100g pack	120	1x10		
787	Ms-L/001	Glycerol	syrup		120	1x10		
788	Ms-T/001	Allopurinol	Tablet	100mg	1800	10x10		
789	Ms-T/002	Colchicin	Tablet	0.5mg	240	10x10		
790	Ms-T/003	Hydroxychloroq uine	Tablet	200mg	1200	10x10		
791	Ms-T/004	Leflunomide	Tablet A	10mg	1200	10x10		
792	Ms-T/005	Leflunomide	Tablet B	20mg	120	10x10		
793	Ms-T/006	Potassium iodide	Tablet	60mg	120	10x10		
794	Ms-T/007	Pyridostigmine	Tablet	60mg	120	10x10		
795	Ms-M/006	Enema phosphate	enema	100ml	1200	1x10		
796	Ms-I/023	Intraperitoneal dialysis solution	injection	500ml bottle	360	1x10		
797	Ms-M/007	Isolyte M	infusion	500ml/ bottle	3600	1x10		
798	Ms-M/008	Isolyte-P	infusion	500ml/ bottle	2400	1x10		

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799	Ms-M/009	Phosphorus	oral suspension	100ml bottle	360	1x10		
800	Ms-M/010	Placentrex	lotion	60ml lotion	300	1x10		
801	Ms-M/011	Polystyrene sulfonate	Powder for solution	15g	240	1x10		
802	Ms-T/008	Sildenafil	Tablets	25mg	600	10x10		

Note:-

The above quantity mentioned for this supply cum rate contract is indicative and may vary as per the actual requirement of hospitals.
The bidder should quote rate for the above mentioned packing unit only

General Requirement :

The manufacturer should ensure Stability of the formulations and its ingredients in the packing supplied.

2. The blister packing of tablets/Capsules should have Aluminium foil back.
3. Strip packing should be of Aluminium / Alu- Alu foils.
4. Aluminium foil strips refer to thickness not less than 40 microns.
5. The rigid PVC used in blister packing should be of not less than 250 microns.
6. Small tablets packed in blister should be packed to facilitate easy removal of a tablet without breaking/ crushing.

Generic Name of drug should be printed in clearly legible bold letters

Annexure -VIII
Ref. Clause No. 14, 17 & 20

ALL INDIA INSTITUTE OF MEDICAL SCIENCES, RAIPUR (C.G.)

GUIDELINES FOR BLACKLISTING/DEBARRING OF
PRODUCT OR SUPPLIER/COMPANY

- 1. ON SUBMISSION OF FALSE, FORGED OR FABRICATED DOCUMENTS OR CONCEALING OF FACTS:**

The Bidder who submits false, forged or fabricated documents or conceals facts with intent to win over the Bid or procure purchase order; EMD of such Bidder firm will be forfeited and firm will be liable for blacklisting for a period of not Less than 2 years. The firm will also be liable for Legal action depending on the facts & circumstances of the case.

2. ON ACCOUNT OF FAILURE TO ENTER INTO AGREEMENT OR WITHDRAWAL AFTER AGREEMENT OR REFUSAL / FAILURE TO SUPPLY:

- 2.1 The successful Bidder fails to execute the agreement after being declared as L-1, L-2 or L-3 etc. to perform the obligations under the Bid conditions, EMD of such Bidder firm will be forfeited and firm will be liable for blacklisting for a period of not less than 2 years or the period specified in Bid document.
- 2.2 The successful Bidder after entering into an agreement withdraw or fail to honour commitments as per Bid conditions, EMD of such Bidder firm will be forfeited and firm will be liable for blacklisting for a period of not Less than 2 years

3. ON ACCOUNT OF NON-SUPPLY:

- 3.1 The supplier shall start to supply according to Bid condition from the date of purchase order and shall complete the supplies within 45/60 days as mentioned in Purchase Order or as stated in Bid condition.
- 3.2 AIIMS, RAIPUR will be at liberty to accept or reject the supply made belatedly as per the terms and conditions of the Bid documents. In the event of acceptance of delayed supply the liquidated damages shall be imposed at the rate stipulated in conditions of the Bid document.
- 3.3 If the supplier fails to execute the purchase order and informs AIIMS, Raipur about its inability to execute the order and non-compliance of the purchase order due to act of vis-majeure, then the Director, AIIMS Raipur will issue appropriate order on merits of case.
- 3.4 If the supplier fails to execute atleast 50% of the quantity mentioned in single purchase order and such failure in supply continues for three purchase orders, then supplier firm will be liable for blacklisting for a period of not Less than 2 years. As a result such supplier will be ineligible to participate in any of the Bids for

particular item(s) of drugs / medicines for a period of not less than 2 years or the period specified in Bid document.

4. ON ACCOUNT OF QUALITY FAILURE OF DRUGS & MEDICINES:

- 4.1 The drugs supplied by the suppliers to the **AIIMS, Raipur** are quarantined and samples of each and every batch of drugs /medicines are drawn on random basis and forwarded to Quality Control Wing of the State Govt./Central Govt or Authorized by Govt. Organization. The samples are then sorted; common batches pooled, coded and are sent to the empanelled laboratories for quality control test as per the QC Policy of AIIMS, Raipur.
- 4.2 Samples of all sterile surgicals & sutures items falling in the categories of drugs will also be drawn as per above policy and all of them will be subjected essentially for sterility testing.
- 4.3 If such samples pass quality test in all respects, AIIMS, Raipur will instruct to issue items of drugs to various hospitals / institutions.
- 4.4 If the sample fails in quality test and report is received certifying that sample is not of standard quality, the drugs of the batch will not qualified for issue and supplier shall be informed to take back stocks of such batch, which failed the quality test and other consequences would follow as per the conditions in the Bid documents.
- 4.5 If two batches of a particular item supplied under a Bid tenure by the supplier are declared as Not of Standard Quality by an empanelled lab or Govt. Lab in test for assay and such failures are further confirmed by another empanelled lab / Govt. Lab, then the particular item of the drug shall be liable for blacklisting for a period of not Less than 2 years.
- 4.6 If three batches of a particular item supplied under a Bid tenure by the supplier are declared as Not of Standard Quality during its entire shelf life by an empanelled lab or Govt. Lab in test for assay and / or in any other parameter(s) and if such failures are further confirmed by another empanelled lab or Govt. Lab during its entire shelf life, the particular item of the drug shall be liable for blacklisting for a period of not Less than 2 years.
- 4.7 In case three products of a company/supplier are blacklisted for supply made during Bid duration the Supplier / Company shall be liable for blacklisting for a period of not Less than 2 years.

- 4.8 In case, any sample (even one batch) is declared as Spurious or Adulterated by an empanelled lab or Govt. Lab and if such failure is further confirmed by another empanelled lab / Govt. Lab during its entire shelf life, the Supplier / Company shall be liable for blacklisting for a period of not less than 3 years.
- 4.9 If any statutory sample of **AIIMS, Raipur** supply drug is drawn by Drugs Control Officer on suo-moto basis or on complaint and if it fails in quality parameters, the report is conclusive till it is challenged by supplier / company. If it is challenged then the report of Director, C.D.L., Kolkatta shall be conclusive and action as contemplated in foregoing paragraphs will be initiated in the matter of blacklisting of product or company. However if failure is of such nature wherein Drugs Controller of State grants prosecution sanction under Drugs & Cosmetics Act, 1940, then even failure of such one batch shall be considered adequate for blacklisting the product for not less than 2 years and in case of involvement of three different products the **Supplier / Company** as a whole shall be liable for blacklisting for a period of not Less than 3years.

5. PROCEDURE IN THE EVENT OF QUALITY FAILURE WILL INVOLVE THE FOLLOWING STEPS:

- 5.1 On receipt of adverse quality test report from empanelled lab or Govt. Lab of a quarantined stock, instructions will be issued immediately through e-mail to the **AIIMS, Raipur** not to release such stock and entries be made by QC Cell at headquarter.
- 5.2 **AIIMS, Raipur** will take appropriate measures immediately to segregate such stock and label all cartons as "NOSQ Drugs-Not for release" and shift it from quarantine area to Non-Release / Rejected Drugs Area (which is under lock & key) till its lifting by the supplier.
- 5.3 Immediately on receipt of NOSQ report, the second sample should be sent to another empanelled lab / Govt. Lab by the QC Cell.
- 5.4 The supplier shall be informed immediately about the test results and instructions be issued to lift the entire stock at supplier's expenses of such batch no. drug which is declared as "NOSQ" by the empanelled lab / Govt. Lab. However, in case of serious quality failure i.e. if drug is declared or adjudged spurious, adulterated or grossly substandard, **AIIMS, Raipur** will be directed to contact the District Drugs Control officer for drawing statutory sample of such batch as per Act. The DDW Incharge has to keep adequate quantity

of such drug for statutory sampling by Drugs Control officer.

- 5.5** In case of drug declared as **Not of Standard Quality** on subsequent sampling after the batch was released the procedure given in sub-para 5.2 will be followed in respect of stock available with the Drug store house. In respect of stock already issued and **AIIMS, Raipur** will take immediate steps to RETRIEVE the unused stock of such drugs from all such institutions and D.D.C.s by all possible mode and means and he/she will ensure that no such NOSQ drug is further distributed to the patients and ensure effective recall.
- 5.6** On receipt of test report from empanelled lab / Govt. Lab, show cause notice will be issued immediately to the concerned supplier calling for explanation within 3 days from the date of receipt of notice in respect of quality failure of concerned batches of drug. The supplier will be required to submit the batch manufacturing record, batch analysis report, raw material purchase record & raw material test reports etc. Opportunity for personal hearing, if desired by supplier, may also be accorded.
- 5.7** On confirmation of the test result by the second laboratory, the case will be referred to the disciplinary committee of AIIMS, Raipur for further action..
- 5.8** In case when the second report is contradictory to the first report, the statutory sample will be sent to Govt. Lab, whose report will be final and if the sample has been tested by the Govt. Lab at any stage, its report will be conclusive & final unless challenged as per provisions of Drugs & Cosmetics Act, 1940.

6. EXAMINATIONS OF ISSUES BY DISCIPLINARY COMMITTEE OF AIIMS, RAIPUR:

- 6.1** Each & every case of submission of false documents, failure to execute agreement, non-supply or quality failure, etc. will be referred to disciplinary committee of AIIMS, Raipur for examination on a case to case basis for making appropriate technical recommendation to Director for further appropriate action.
- 6.2** The recommendations of disciplinary committee will be placed before the Director, AIIMS, Raipur who shall take appropriate action which may deem fit in the light of facts & circumstances of the case

by way imposing penalty or debarring or Blacklisting of the particular product or supplier/ company.

- 6.3 If, the quality failure is of such nature that a particular product has been blacklisted according to the procedure stated above, the supplier will not be eligible for participating in any of the Bids for the particular item floated by AIIMS, Raipur for the specified period. For such purpose period of blacklisting will be counted from date of issue of order and it will deemed to be over on completion of the period and as such no fresh orders will normally be required for re-eligibility purpose. Similarly if the supplier /company is blacklisted the supplier will not be eligible for participating in any of the Bids for any of the items during blacklisted period.

7. POWER OF REVIEW:

Subsequent to the action taken on the basis of available facts if some new facts & evidences such as reversal of test results findings by Appellate Laboratories etc. are brought to the notice of the corporation, the Director of AIIMS, Raipur will have the right to review the earlier action. He may seek advice from the disciplinary committee in such matters.

8. Savings:

The blacklisting of particular product or supplier / firm will be done without prejudice to other penalty which may be imposed as per the conditions of Bid documents and also to other actions which may be initiated under Drugs and Cosmetics Act 1940 or any other law of land. AIIMS, Raipur will display names of such blacklisted products and companies on its website and also circulate the same among all stakeholders viz. Director of remaining 6 AIIMS, including respective State Drug Controllers where the supplier / company is located.

9. JURISDICTION

In the event of any dispute arising out of the orders and implementation thereof, such dispute shall be subject to the jurisdiction of the Courts of Raipur City only or Hon'ble Chhattisgarh High Court, Bench at Bilaspur.

EXPLANATIONS:

- i. Increase in the cost of raw materials, power cut, Labour strike, insolvency, closure of the factory would not be considered as act of vis-majeure.

- ii. The meaning of 'Spurious drugs' or 'Adulterated Drugs' will be construed in strict sense under the provisions of Drugs & Cosmetics Act, 1940. For the purpose of blacklisting a drug will be considered 'Spurious' if empanelled lab / Govt. Lab so declare the product or it is found containing either no drug or very poor drug contents on testing or it is purported to be manufactured of whom it is not truly a product or which is likely to cause grievous hurt within the meaning of Sec. 320 Of IPC. Similarly for the purpose of blacklisting a drug will be considered 'Adulterated' if empanelled lab / Govt. Lab so declare the product or it is found containing any poisonous, deleterious, harmful or toxic substances or which is likely to cause grievous hurt.
- iii. Purchase Orders, if any, already issued before taking any blacklisting action or replacement orders given in past will not be affected in view of action taken as per above guidelines but all strict quality checks shall be observed for each supply of products.
- iv. The action proposed as above is not in conflict to any express conditions laid down in corresponding Bid and in case of any overlapping, the Bid condition will prevail.

ANNEXURE-IX
Ref. Clause No. 1(d), 5(y), 21

UNDERTAKING FOR EMPANELMENT

I Name.....S/o.....Age.....Prop./Partner/Director/
Power of attorney holder of firm M/s.....situated at (Complete address
of Mfg. unit).....bearing drug license on Form 25 & 28 or form 10 bearing
Number..... &.....respectively, issued on
dated.....valid/Renewed up to.....do here by declare on oath as

follows:-

- 1 That I have applied for empanelment for supply of Drugs & Medicines for the items I have quoted in the tender as enlisted in Annexure –VI.
- 2 That I/We have carefully read all the conditions of Bid in Tender Enquiry No. ADMIN/Tender/DRUGS/1/2013 Dated 02.11.2013 for supply cum rate contract and empanelment for supply of Drug and Medicines for **AIIMS, Raipur** and accept all conditions of Bid, including amendments if any.
- 3 That I will be considered empanelled for the items which are declared technically responsive.

That I have deposited the required fees for empanelment.

Date

**Name & Signature
with Seal**

Annexure-X
Ref. Clause No.18.2

Supplier Consolidated Invoice

Name of Supplier:		
Complete Address:		
E-mail ID:		
DL NO.:	TIN No.:	Invoice No.:
		Date:

Purchaser: Director Address: AIIMS, Raipur Tatibandh, GE Road Raipur Raipur, Phone No. 0771-2573222	Purchase Order No.: Date:
--	--

Name of Item/Description:

S. No.	Name of Medicine	Medicine Code	Invoice/Challan No.	Date	Packing Size	Batch No	Mfg. Date	Exp. Date	Quantity supplied in No. (Batch wise)	Unit-Rate in ₹		Total Amount to be paid in ₹	
										inclusive of transportation, insurance, Packing and any incidental charges, Excise Duty, Customs duty, VAT, CST , & all other statutory duties of the govt.	inclusive of transportation, insurance, Packing and any incidental charges, Excise Duty, VAT, CST Customs duty & all other statutory duties of the govt.	In Figure	In word
A	B	C	D	E	F	G	H	I	J	In Figure	In word	In Figure	In word
1													
2													
3													
Grand Total													

Authorised Signatory

**Annexure-XI
Ref. Clause No. 18.2**

Analytical Report Regarding Quality

Name of Supplier:-	
Address:-	
PO No:-	Date:-
Drug Name:-	
Details of in house test report:-	

S.No.	Name of Lab.	Test report No.	Date	Batch No.	Qty. Supplied	Result
1.						
2.						
3.						
4.						
5.						
6.						
7.						
8.						
9.						
10.						
11.						
12.						

**Authorised
Signatory**

**Annexure-XII Ref.
Clause No. 12**

Security form (Bank guarantee)

To,

Director, AIIMS Raipur

WHEREAS (Name of Supplier)

Hereinafter called "the Supplier" has undertaken, in pursuance of

Contract (Letter of Acceptance) No.....dated.....2013 to

Sign of Bidder

supply..... (Description of Goods) hereinafter called "the Contract".

AND WHEREAS it has been stipulated by you in the said Contract that the Supplier shall furnish you a bank Guarantee from a Scheduled Bank for the sum specified therein as security for compliance with the Supplier's performance obligations in accordance with the Contract.

AND WHEREAS we have agreed to give the supplier a Guarantee:

THEREFORE WE hereby affirm that we are Guarantors and responsible to you, on behalf of the Supplier, up to a total of.....(Amount of the Guarantee in Words and Figures) and we undertake to pay you, upon your first written demand declaring the Supplier to be in default under the said Contract and/or any other contract or for set off any other dues pending against the supplier, without cavil or argument, any sum or sums within the limit of(Amount of Guarantee) as aforesaid, without your needing to prove or to show grounds or reasons for your demand or the sum specified therein.

This guarantee is valid until the.....day of.....2015.....

Signatures and Seal of Guarantors

Date.....

Address:.....

.....

.....

.....

Note:- The validity of bank guarantee should be for 24 months from the date of issuance of Bank Guarantee.