

Issued to M/s. _____

Sl.No

NOT TRANSFERABLE

Ref.No: 011/M(P)/ARV-DRUG/TNMSC/2022, Dt.05.07.2022

TAMILNADU MEDICAL SERVICES CORPORATION LIMITED
(A Government of Tamilnadu Undertaking)
No. 417, Pantheon Road, Egmore, Chennai – 600008.
Phone: 044-28191890, 28190259
Website: www.tnmsc.tn.gov.in, www.tenders.tn.gov.in
Email: mngRPurDrug.tnmsc@tn.gov.in, gmdrug.tnmsc@tn.gov.in

TENDER FOR THE SUPPLY OF ARV DRUGS

TO

TAMILNADU STATE AIDS CONTROL SOCIETY

**UNDER RATE CONTRACT SYSTEM FOR ONE YEAR FROM THE DATE OF
ACCEPTANCE**

LAST DATE FOR RECEIPT OF TENDER: 19.07.2022 upto 11.00 A.M.



***TamilNadu Medical Services Corporation Limited
No. 417, Pantheon Road,
Egmore, Chennai – 600008.***

Telephones : 044-28190259 / 28191890

Email : mngripurdrug.tnmsc@tn.gov.in, gmdrug.tnmsc@tn.gov.in

**TENDER FOR THE SUPPLY OF ARV DRUGS TO TAMILNADU STATE AIDS
CONTROL SOCIETY UNDER RATE CONTRACT SYSTEM FOR ONE YEAR FROM
THE DATE OF ACCEPTANCE**

Tender Reference	:	011/M(P)/ARV-DRUG/TNMSC/2022 Dt.05.07.2022
Date of commencement of Sale of tender document	:	05.07.2022
Last date for sale of Tender document	:	18.07.2022
Last date and time for Receipt of tender	:	19.07.2022 upto 11.00 A.M.
Time and date of opening of tender	:	19.07.2022 at 11.30 A.M.
Place of opening of Tender	:	Tamilnadu Medical Services Corporation Ltd., No. 417, Pantheon Road, Egmore, Chennai - 600 008.
Address for Communication	:	Tamilnadu Medical Services Corporation Ltd., No. 417, Pantheon Road, Egmore, Chennai - 600 008.
Cost of the Tender Document	:	Rs.5,725/- (Inclusive of GST)

Alternatively, the tender document can be downloaded from the websites of www.tnmsc.tn.gov.in and www.tenders.tn.gov.in on free of cost.

TAMILNADU MEDICAL SERVICES CORPORATION LIMITED

TENDER FOR THE SUPPLY OF ARV DRUGS TO TAMILNADU STATE AIDS CONTROL SOCIETY UNDER RATE CONTRACT SYSTEM FOR ONE YEAR FROM THE DATE OF ACCEPTANCE

The TNMSC Ltd., a Govt. of Tamilnadu undertaking is involved in procuring quality Drugs and Medicines directly from the manufacturers and importers through open tenders system and ensures timely Supply of Drugs and Medicines to about 11,500 medical institutions all over Tamilnadu.

Tender Inviting Authority - The General Manager (Drugs), Tamilnadu Medical Services Corporation Limited, No.417, Pantheon Road, Egmore, Chennai-600008, (hereinafter referred as **Tender Inviting Authority** unless the context otherwise requires).

Tender Accepting Authority - Tamilnadu Medical Services Corporation Limited (hereinafter referred as **TNMSC Ltd.**, unless the context otherwise requires)

Tender Inviting Authority invites **Tender for the Supply of ARV Drugs to Tamilnadu State AIDS Control Society under rate contract system for one year from the date of acceptance.**

1. LAST DATE FOR RECEIPT OF TENDERS.

- (a) Sealed Tenders [in two separate covers {Technical bid (Cover "A") and Price Bid (Cover "B")}] will be received till **11.00 A.M. on 19.07.2022** by the Tender Inviting Authority, Tamilnadu Medical Services Corporation Limited (TNMSC Ltd.), No.417, Pantheon Road, Egmore, Chennai - 600008, for the Supply of ARV Drugs

to Tamil Nadu Medical Services Corporation Limited under rate contract system for one year from the date of acceptance. The Name and address of the Firm / Company quoting the Tender should be clearly indicated in the outer cover as per the constitution of the Firm / Company particulars.

- (b) The bid shall be valid for a period of 120 days from the date of opening of Cover-A (Technical Bid) and prior to the expiry of the bid validity, the Tender Inviting Authority may request the Tenderers to extend the bid validity for further period as deemed fit. However, TNMSC Ltd., reserves the right to place purchase orders at the quoted rate till such period of validity of the tender and the tenderer(s) are bound to accept the orders at the rates quoted / accepted and within the production capacity indicated in the tender, irrespective of execution of agreement / finalization of price.

2. ELIGIBILITY CRITERIA

- (a) Tenderer shall be a manufacturer having valid Own Manufacturing License / Direct importer holding valid Import License and Loan Licensees having Valid Loan License. Distributors / Suppliers / Agents are not eligible to participate in the Tenders.
- (b) (i) **Average Annual turnover for Non Domestic Enterprise in last 3 years i.e., 2018-2019, 2019-2020 and 2020-2021 shall not be less than Rs.4.00 Crores and the turnover should be a minimum of Rs.4.00 Crores**

in any one of the financial years (ie., 2018-2019, 2019-2020 and 2020-2021).

(ii) Average Annual turnover for Domestic Enterprise **in last 3 years i.e., 2018-2019, 2019-2020 and 2020-2021** shall not be less than *Rs.2.00 Crores* and the turnover should be a minimum of *Rs.2.00 Crores* ***in any one of the financial years (ie., 2018-2019, 2019-2020 and 2020-2021).***

(c) (i) Tenderer should at least have 3 years Market Standing as a manufacturer / loan licensees for each drug quoted in the tender as manufacturer / loan licensees. In case of an Importer, their principal manufacturer should have 3 years market standing in India and the Importer should have 3 years market standing in the pharmaceutical field. And also, the importer should have due authorization for quoting drugs from the principal manufacturer along with relevant import licenses. In case of any new drug / drugs out of patent period, should possess relevant market standing as applicable.

(a) In case of product(s) with similar formulation but with varied strengths, Market Standing of 3 years for any strength of similar formulation shall be considered for quoted product(s) as equivalent, subject to possession of Manufacturing License for the quoted product(s) for a period of not less than 3 years.

- (ii) Tenderer should have obtained permission to manufacture the drugs(s) quoted as per specification in the tender from the competent authority. The imported product(s) should have valid import license by the competent authority.
- (iii) In both cases (i) and (ii) above, the permission provided by Drug Controller General of India (DCGI) shall be provided as applicable.

(d) **FOR PRODUCT(S)**

- (i) Tender should not be submitted by the firm / company / loan licensee for the Product(s) for which the firm / Company / loan licensee has been blacklisted / banned / debarred by TNMSC Ltd., on any grounds.
- (ii) Tender should not be submitted for the product(s) for which the firm / company / loan licensee has been blacklisted by any other State Government / Central Government / its Drug procurement agencies due to quality failure and/or fraudulent/illegal practices of the drugs supplied.

(e) **FOR FIRM/COMPANY**

- (i) The Company / Firm / loan licensee which has been blacklisted by TNMSC Ltd., due to quality failure of the drugs / non-performance of tender conditions / any other grounds should not participate in the tender during the period of blacklisting.

- (ii) The Company / Firm / loan licensee which has been blacklisted by any other State Government / Central Government / its Drug procurement agencies due to quality failure of the drugs and / or fraudulent / illegal practices involved in the tender should not participate in the tender during the period of blacklisting.
- (f)
 - (i) The Tenderer should give a notarized affidavit that they have not been black listed due to quality failure of the drug(s) and / or fraudulent / illegal practices for the quoted product(s) / firm in the tender by any other State Government / Central Government / its Drug procurement agencies or by TNMSC Ltd., and also not blacklisted by TNMSC Ltd., due to non-performance of tender conditions and thereby eligible to participate in the present tender. (Notarized affidavit as per **Annexure-IV**).
 - (ii) During the validity of the tender if the firm / Company / loan licensee is blacklisted by any other State Government / Central Government / its Drug procurement agencies on the grounds stated above in Para (d), (e) and (f) / convicted by any Court of law in India, shall be intimated to TNMSC Ltd., by the corresponding firm / company / loan licensee.

In case of Loan licensee,
 - (a) The Loan licensee and / or its manufacturing facility if blacklisted for particular product(s) by any other State Government / Central

Government / its Drug procurement agencies on the grounds stated above / convicted by any Court of law in India, shall be intimated to TNMSC Ltd.,

(b) The Loan licensee and / or its manufacturing facility if blacklisted in whole by any other State Government / Central Government / its Drug procurement agencies on the grounds stated above / convicted by any Court of law in India, shall be intimated to TNMSC Ltd.,

(c) Tenderer should quote at least for **60%** of the tendered quantity of each drug exclusively earmarked for TNMSC Ltd., in this tender irrespective of any other tenders that may be floated by TNMSC Ltd., for any drug in which the same firm / company / loan licensee become eligible/selected. However after evaluation of the price bid, the eligible tenderers are permitted only for upward revision of the production capacity / earmarked quantity of any drug(s) approved. On any circumstances downward revision of the quoted quantity and/or production capacity is not permitted. TNMSC Ltd., have the prerogative either to accept or to reject the revision given by the tenderer among the drugs.

3. GENERAL CONDITIONS

(i) A complete set of tender documents may be purchased by any interested eligible person from the office of Tender Inviting Authority between **10.00 AM to 5.00 PM**

on or before 18.07.2022 on all working days either in person or by post by making an application in writing and upon payment of a non-refundable fee as indicated in the advertisement in the form of Demand draft drawn in favor of Tamilnadu Medical Services Corporation Limited payable at Chennai. Tender Inviting Authority will not be responsible in any way for postal delay.

- (ii) Alternatively, the tender document can be downloaded from the websites www.tnmsc.tn.gov.in and www.tenders.tn.gov.in at free of cost.
- (iii) The complete set of tender (Cover A and Cover B) should be submitted on or before **11.00 A.M. on 19.07.2022**.
- (iv) All tenders must be accompanied with Earnest Money Deposit as specified against each drug in **Annexure-IX** of the Tender document UNLESS EXEMPTED under clause 7.2 here under.
- (v) Tenders will be opened in the presence of Tenderers / **authorized representatives** of tenderers who choose to attend on the specified date and time or as per Covid-19 restrictions in place at the time of opening.
- (vi) (a) At any time prior to the date of submission of Tender, Tender Inviting Authority may, for any reason, whether on own initiative or in response to a clarification requested by a prospective Tenderer, may modify the condition in Tender documents by an amendment. All the prospective Tenderers

who have purchased the tender document will be notified of the amendment in writing and that will be binding on them. In order to provide reasonable time to take the amendment into account in preparing their bid, Tender Inviting Authority may at discretion, extend the date and time for submission of tenders.

- (b) Any person who has downloaded the tender document should watch for amendment, if any, on the website www.tnmsc.tn.gov.in and www.tenders.tn.gov.in for which TNMSC Ltd., will not issue any separate communication to them.
- (vii) Interested eligible Tenderers may obtain further information in this regard from the office of the Tender Inviting Authority on all working days between 10.00 AM and 5.00 PM.

4. TECHNICAL BID - COVER "A"

- 4.1. The Tenderer should furnish the following documents in a separate cover hereafter called **"Cover A"**. **(All the documents submitted should be signed and sealed by the Tenderer in each page and photo copies of the documents should be attested by the Tenderer and also be notarized in each page)**

- (a) Earnest Money Deposit, IF NOT EXEMPTED under clause 7.2, shall be paid to TNMSC Ltd., as indicated against each drug in **Annexure-IX** of the tender document in the form of **Bankers Cheque or Demand Draft**

and/or irrevocable Bank Guarantee favoring Tamilnadu Medical Services Corporation Ltd., payable at Chennai and the Tenderer should also enclose split up details of EMD for each drug. EMD in any other form like ***cheque/cash/postal order*** etc. will not be accepted and the tender will be rejected.

- (b) Documentary evidence for the constitution of the Company/Firm such as Memorandum and Articles of Association, Partnership deed, Permanent Registration Number, GST registration certificate etc., with details of the Name, Address, Telephone Number, Fax Number, e-mail address of the firm and of the Managing Director / Partners / Proprietor. Also the list of present Directors in the Board of the Company / Partners duly certified by a Company Secretary of the Company / Practicing Company Secretary / Chartered Accountant to be furnished.
- (c) The Tenderer should furnish attested photocopy of Manufacturing Licence / Loan License for the product, duly approved by the Licensing Authority for each and every product quoted as per specification in the tender. The licence must have been duly renewed up to date and the drugs quoted shall be clearly highlighted in the licence. Original documents should be produced when demanded for verification.

- (d) Attested photocopy of import license (in Form 10 with Form 41) in accordance with Rule 21 of the Drugs and Cosmetics Act 1940 and Rules 1945 (if the product is imported) should be furnished. In case of new drug, import permission in Form-45 / Form-45A should be furnished in accordance with Rule 122-A of the Drugs and Cosmetics Act 1940 and Rules 1945. The licence must have been renewed up to date. A copy of a valid licence for the sale of Drugs imported by the firms issued by the State Licensing Authority shall also be enclosed. Original documents should be produced as and when demanded for verification.
- (e) The instruments such as power of attorney, resolution of board etc., authorizing an officer (employee) of the Tenderer should be enclosed with the tender duly signed by the Authorized signatory of the Company/Firm and such authorized officer (employee) of the Tenderer should sign the tender documents.
- (f) Authorization letter nominating **a representative (With the photograph) of the Tenderer to submit the tender and to transact with TNMSC Ltd till price bid opening if different from (e) above. Such nominated representative shall not be permitted to represent more than two companies, bidding for the same product. In case of companies bidding for different products, the nominated representative should not represent more than 10 companies. Further transactions shall be**

restricted with the officer (employee) as stated in (e) above. TNMSC Ltd shall then provide all communications directly to the tenderer only through postal / Courier services and e-mails to the designated address / mail id.

- (g) **Market Standing Certificate** issued by the Licensing Authority as a Manufacturer for each product quoted **for last 3 years (i.e., 2018-2019, 2019-2020 and 2020-2021)** (Certificate obtained specifically for TNMSC Ltd., tender / in General) should be enclosed with list of products. In case of direct importer, evidence of import of the said products for the last three years such as bill of landing, bill of entry for last three years and certificate of analysis are to be submitted (irrespective of the Importer). In case of supply of Serum or Vaccines, Market standing certificate issued by Licensing Authority **for one year (i.e.2019-20 or 2020-21)** should be enclosed.
- (h) **Performance statement of manufacturing / importing unit** to establish market standing **for last 3 years (i.e., 2018-2019, 2019-2020 and 2020-2021)** as per format in **Annexure-VII** along with the Attested scanned copies of the BMR for the quoted products to establish 3 years Market standing in the CD in Cover A (In case of Serum or Vaccines one Year) shall be furnished. However as and when required by TNMSC

Ltd., the said documents should be submitted as attested true copy by the tenderers.

- (i) **Non-conviction Certificate / Conviction Status Certificate** issued by an appropriate authority of the State certifying that the Firm / Company / Loan licensee has not been convicted / convicted [along with list of product(s)] and such firm should not submit tender for products which are convicted in any Court of Law for 3 years from the date of conviction.
- (j) **Current Good manufacturing practices Certificate (cGMP) as per revised Schedule-‘M’ or WHO-GMP** (for manufacturers / manufacturing units of loan licensee) issued by the Licensing Authority. The Tenderer shall also furnish a notarized affidavit in the format given in Annexure-III declaring that the Tenderer complies with the requirements of cGMP (as per revised Schedule-‘M’) or WHO-GMP. In case of Imported drugs, labels and product literature of all quoted product(s) must be submitted with WHO-GMP or certificate which is at par with WHO-GMP issued by the authorities of exporting countries like U.S. FDA etc., or COPP certificate of their Principal Manufacturing Company or firm.
- (k) **Annual turnover statement for last 3 years i.e., 2018-2019, 2019-2020 and 2020-2021** should be furnished in the format given in Annexure-VIII duly certified by the Chartered Accountant.

- (l) Printed Annual reports including the **Balance Sheet and Profit and Loss Account for last 3 years i.e., 2018-2019, 2019-2020 and 2020-2021** duly certified by the Chartered Accountant.
- (m) **GST returns from 01.01.2021 to 31.03.2021 (as applicable)** along with GST registration copy of the tenderer.
- (n)
 - (i) Undertaking (as in the proforma given in **Annexure-I**) for embossment of logo on tablets, capsule shell, on labels of vials, Ampoules, bottles, tubes etc. as the case may be, and for supply of tablets / capsules on the strips as per the conditions specified under Clause 14 herein, duly notarized by the Notary Public.
 - (ii) Undertaking (as in the proforma given in **Annexure-II**) for affixing the logo on the Secondary/Primary packing for the imported drugs along with Brand/Trade names.
- (o) The details containing the name and address of the manufacturing premises / importing unit where the drugs quoted are actually manufactured / imported should be given as per the format in **Annexure-XIII** along with exact address of the Registered / Corporate office and address to which communications are to be sent.

- (p) Documents, if any, to show that the manufacturing unit/importer has been recognized by any other Indian / International Standard Organizations as applicable etc (optional).
- (q) Details of technical personnel employed in the manufacturing and testing of drugs (Employee Name, Qualification, and Experience) as endorsed in license.
- (r) List of drugs quoted (The name & Drug code of the Drugs quoted, yearly production capacity of individual drug earmarked exclusively for the tender of TNMSC Ltd., and the amount of EMD for each drug alone should be furnished and **the rate of those drugs should not be indicated in this list**), as shown in the **Annexure-XIV** should be given in duplicate.
- (s) A Checklist (**Annexure-XVIII**) indicating the above documents submitted with the tender document and their respective page number shall be enclosed with the tender document. The documents should be serially arranged as per this **Annexure-XVIII** and should be securely tied or bound. If a company/firm has two or more separate manufacturing units at different sites / States, which are not separate entities, then the company will be allowed to submit only one tender for all units but necessary document regarding separate manufacturing units will have to be

submitted as a separate set with the same tender. But a bidder will be allowed to submit only one offer for one product.

(t) The tender document should be signed only by the authorized official of the Tenderer in all pages with office seal as indicated in (e) above.

(u) All the documents enclosed with the tender document should also be signed by the authorized official of the Tenderer as indicated in (e) above.

4.2. The above documents should be sealed in a separate Cover Super scribed as **"TECHNICAL BID - COVER "A" – TENDER FOR THE SUPPLY OF ARV DRUGS TO TAMILNADU STATE AIDS CONTROL SOCIETY UNDER RATE CONTRACT SYSTEM FOR ONE YEAR FROM THE DATE OF ACCEPTANCE DUE ON 19.07.2022 AT 11.00 A.M.". AND ADDRESSED TO "THE TENDER INVITING AUTHORITY, TAMILNADU MEDICAL SERVICES CORPORATION LIMITED, NO.417, PANTHENON ROAD, EGMORE, CHENNAI – 600008"**.

5. PRICE BID - COVER "B"

5.1. Cover "B" contains the Price Bid of the Tenderer.

(i) Bid should be typewritten and every correction and interlineations in the bid should be attested with full signature by the Tenderer, failing which the product(s) will be treated as ineligible. Corrections done with correction fluid should also be duly attested.

- (ii) Each page of the price bid should be duly signed by the Tenderer affixing the official seal.
- (iii) The Tenderer shall fill in the rate in the **Annexure-XIX** for the drugs quoted and also in the Pendrive (supplied with tender document) and such filled in **Annexure-XIX** along with the Pendrive (Soft Copy) should be submitted.
- (iv) The Tenderers who have downloaded Tender document, shall also download and copy the **ARVDRUG2022.XLS** (Excel) file in the Pendrive. Such Pendrive may be used for quoting the rate in soft copy.
- (v) In determining the lowest evaluated price, (the rate quoted per unit or landed price in **Annexure-XIX**) the evaluation shall include Basic Price Inclusive of Incidental Charge + Packing & Forwarding Charges + Freight and Insurance Charges + Customs Duty (if applicable) + GST.
- (vi) The rate quoted in column 8 of **Annexure-XIX** should be for a unit and for the given specification. **The rates quoted in paisa are to be in 2 digits.** The Tenderer is not permitted to change/alter specification or unit size given in the **Annexure-XIX**. **Request for any changes in the rates quoted will not be entertained by TNMSC Ltd., on any circumstance other than the Tax revision made by any Government. Tax portion should be mentioned in the form of percentage only (%).**

- (vii) The details of rates and manufacturing capacity earmarked for TNMSC Ltd., (for each drug individually) given in **Annexure-XIX** should be entered clearly given along with the tender. **The production capacity earmarked for TNMSC Ltd., as indicted in Annexure-XIV in Serial No 7 (Column 8) or up-scaled revised quantity of the Tenderers alone will be considered for placement of purchase orders.**
- (viii) The bidder shall necessarily quote the excise duty or customs duty applicable and when the drug is excisable or imported as the case may be. The relevant chapters and tariff applicable should be indicated for each drug (or) as per the relevant Acts / rules in force at the time of price bid opening.
- (ix) The bidder shall specifically mention “ **EXEMPTED** “ when the drug is excisable but exempted for the time being, based on turn over or for any other grounds by the notification issued by the Government of India (**Also refer clause 17.6**).
- (x) The bidder once quoted the excise rate is not permitted to change the rate/amount unless such change is supported by the notification issued by the Government of India or by the order of the court, after submission of Tender (or) as per the relevant Acts / rules in force at the time of price bid opening.
- 5.2. The Tenderers shall submit duly signed **Annexure-XIX** in a sealed cover Superscribed as “**PRICE BID COVER “B” – TENDER FOR THE SUPPLY OF ARV DRUGS TO TAMILNADU STATE AIDS CONTROL SOCIETY UNDER**

RATE CONTRACT SYSTEM FOR ONE YEAR FROM THE DATE OF ACCEPTANCE". The "Cover-B" should also be addressed to the "TENDER INVITING AUTHORITY, TAMILNADU MEDICAL SERVICES CORPORATION LIMITED, NO.417, PANTHEON ROAD, EGMORE, CHENNAI - 600008".

5. 3. Two separate sealed covers {Technical bid (Cover "A") {Refer Clause No.4.2} and Price Bid (Cover "B")} {Refer clause 5.(2) } shall be placed in a cover which shall be sealed and Superscribed as **"TENDER FOR THE SUPPLY OF ARV DRUGS TO TAMILNADU STATE AIDS CONTROL SOCIETY UNDER RATE CONTRACT SYSTEM FOR ONE YEAR FROM THE DATE OF ACCEPTANCE DUE ON 19.07.2022 AT 11.00 A.M."** and addressed to the **TENDER INVITING AUTHORITY, TAMILNADU MEDICAL SERVICES CORPORATION LIMITED, NO. 417, PANTHEON ROAD, EGMORE, CHENNAI-8**, which shall be submitted within the date and time as specified in Clause 1(a).

5.4. If the last date for submission of Tender is declared as holiday in the state of Tamilnadu including local holiday if any, the tenders may be submitted on the next working day up to 11.00 A.M.

6. OPENING OF COVER "A" AND COVER "B" OF TENDER

6.1 Only authorized official as indicated in Clause 4.1.(e) or (f) only are entitled to be present at the time of opening of Technical Bid - Cover "A" of the tender submitted by them. **No other persons will be permitted.**

6.2 Tenderers, who are found eligible on satisfying the criteria for technical evaluation and inspection will only be invited and the authorized official as indicated in 4.1(e) or (f) alone will be allowed to be present at the time of opening of Price Bid - Cover "B" of the tender.

7. EARNEST MONEY DEPOSIT

7.1 (a) The Earnest Money Deposit referred to under Clause 4.1(a) UNLESS EXEMPTED under Clause 7.2, shall be for the amount as indicated against each drug in Annexure-IX of the tender documents. In case a Tenderer is quoting for more than one drug, the Earnest Money Deposit payable by such Tenderer shall be the aggregate total of the Earnest Money Deposit for all the drugs quoted by such bidder. The bidders are required to furnish the breakup of the Earnest Money Deposit for the drugs quoted along with the Bankers Cheque or Demand Draft or irrevocable Bank Guarantee in the format favoring Tamilnadu Medical Services Corporation Limited, Chennai. **However, whenever the total EMD payable is less than Rs.50,000/- the EMD to be paid only by way of Bankers Cheque / Demand Draft, failing which the tender will be rejected. In other case, a minimum of Rs.50,000/- to be paid only by Bankers Cheque / Demand Draft and the balance may be paid by irrevocable Bank Guarantee/Demand Draft/Bankers Cheque. EMD furnished in the form of a Bank Guarantee should remain valid for a**

minimum period of 60 days beyond the validity period of tender (i.e., 120 days + 60 days = 180 days from the date of opening of Cover “A”). The format of Bank Guarantee for EMD is enclosed in the Annexure-XVII. This should be enclosed with the tender in Cover “A”. For the matter of clarity, if the due date for receiving the tenders is extended, the validity period of the tender will automatically stand extended and it is the responsibility of tenderers to ensure that the EMD is valid at the time of Cover-A opening. Earnest Money Deposit in the form of Cheque/Cash/Postal order will not be accepted. Earnest Money Deposit will not earn any interest.

- (b) In case the EMD submitted by the bidder is not sufficient to meet the EMD requirement of all the drugs quoted, the available EMD will be adjusted for the drug(s) in the ascending order of the drug codes of the drugs quoted by the Tenderer, till the EMD is exhausted. Further, the tender of such bidder for the remaining drugs, out of the quoted drugs, will be treated as non-responsive for want of EMD. Any part value of EMD remaining unadjusted will be treated as an excess value furnished.

7.2 EXEMPTION FROM PAYMENT OF EARNEST MONEY DEPOSIT TO DOMESTIC ENTERPRISE

- (i) Domestic Enterprises – Micro and Small Enterprise (MSME) as defined in the Micro, Small, Medium Enterprise Development Act, 2006 (Central Act

27 of 2006), which manufacturers or produces goods within the State and filed Part-II of the Entrepreneurs Memorandum in the District Industries Centres or filed Udyog Aadhaar portal are exempted from payment of EMD.

- (ii) The Domestic Enterprise will be required to furnish a notarized undertaking (as per **Annexure-VI**) to the effect that in the event of non-fulfillment or non-observance of any of the condition stipulated in the **tender / contract**, the Domestic Enterprise shall pay a penalty, equivalent to the **applicable penalty / Earnest Money Deposit of the drug(s) quoted stipulated in the tender** to offset the loss incurred by the Tender Inviting Authority consequent on such breach condition of **tender / contract**.

- 7.3. (i) The tender submitted without EMD will be summarily rejected.
- (ii) The tenders with insufficient Earnest Money Deposit will be processed in accordance to clause 7.1 (b) above.
- (iii) The Earnest Money Deposit will be refunded to the lowest bidders within 30 days from the date of signing the contract agreement and on the deposit of Performance Security.

- (iv) The Earnest Money Deposit will be refunded to the matched bidders within 30 days from the date of signing the contract agreement.
- (v) The Earnest Money Deposit (EMD) of the other bidders would be returned after the expiry of validity of the tender.
- (vi) The Earnest Money Deposit (EMD) will be forfeited, if the Tenderer withdraws his bid **either fully or partially during the validity of the tender / contract period.**
- (vii) The Earnest Money Deposit (EMD) will be forfeited, in case of the lowest / matched bidder, fails to execute the contract agreement and / or deposit the Performance Security within the stipulated time besides other actions.
- (viii) The Tenderer, whose manufacturing unit found to be not complying with the cGMP (but furnished an affidavit in **Annexure-III**) during inspection, will be levied with a fine of Rs.50,000/- or the expenditure incurred by the TNMSC Ltd., in such inspection, whichever is higher. This fine amount shall be deducted from the EMD deposited by the bidder or any other amount payable to them in any nature. The amount will be deducted without any notice. In case of deficit, legal action will be taken against the tenderer for recovery.

8. OTHER CONDITIONS

- 8.1. (i) The details of the required ARV Drugs, etc., are shown in **Annexure-IX**. ***The tender quantity mentioned herein is not a fixed procurement quantity and it is only a tentative requirement and may be increased or decreased*** by the TNMSC Ltd., at its discretion, depending on the actual need. **Hence the Rule 14 (9) (with respect to the limitation on quantity to be ordered) of the TamilNadu Transparency in Tenders Rules 2000 is not applicable being a rate contract.** Though the tentative quantity is indicated in the agreement, the TNMSC Ltd., will confirm the actual requirement then and there only through purchase order(s). The tenderers shall supply the drugs only on the basis of the purchase order issued by the TNMSC Ltd. Any supply without a valid purchase order will not be accepted by TNMSC Ltd., for payment and the TNMSC Ltd., shall not be responsible for any loss on this account.
- (ii) However, once the purchase order/orders is/are issued by the TNMSC Ltd., the tenderer should not renege from the commitment of supplying the quantity mentioned in the agreement / undertaking.
- (iii) The rates quoted shall not be varied with the order quantity or the destination during the full contract period.

- 8.2. Tender has been called for in the **generic name of drugs**. The Tenderers should quote the rates for the generic products only. The composition and strength of each product should be as per specifications given in **Annexure-IX**. Any variation, if found, will result in rejection of the tender/drug. However the imported/combination drugs are allowed to be quoted in trade / brand name.
- 8.3. Rates (inclusive of Customs duty, Transportation, Insurance, any Incidental charges, but Exclusive of GST) should be quoted for each of the required drugs, medicines etc., separately on door delivery basis according to the unit ordered as in the format in **Annexure-IX**. Tender for the supply of drugs, medicines, etc. with cross conditions like “AT CURRENT MARKET RATES” shall not be accepted. Handling, clearing, transport charges etc., will not be paid separately. The delivery should be made as stipulated in the purchase order placed with Tenderers (or) as per the relevant Acts / rules in force at the time of price bid opening.
- 8.4. The price quoted by the tenderers shall not, in any case exceed the controlled price, if any, fixed by the Central/State Government, the Maximum Retail Price (MRP) and the selling price of the tenderer with other organizations. Tender Inviting Authority at its discretion, may exercise, the right to revise the price at any stage so as to conform to the controlled price or MRP or the lowest selling price of the tenderer as the case may be. This discretion will be exercised without prejudice to any other action that may be taken against the Tenderer.

- 8.5. To ensure sustained supply without any interruption, the Tender Inviting Authority reserves the right to split orders for supplying the requirements among more than one Tenderers, pursuant to the provisions laid under the Tamilnadu Transparency in Tenders Act 1998 and Rules made thereunder subject to the production capacity available with other tenderers.
- 8.6. The rates quoted and accepted will be binding on all the Tenderer(s) for the full contract period of one year and any increase in the price on any account/reasons will not be entertained till the completion of this contract period. Accordingly this clause will be applicable for all the orders placed by TNMSC Ltd during the contract period.
- 8.7. No Tenderer shall be allowed at any time and on any ground, whatsoever it may be, to claim revision or modification in the rates quoted by them. Representation to make correction in the tender documents on the ground of clerical error, typographical error, etc., committed by the Tenderers in the Bids shall not be entertained after submission of the tenders. Cross Conditions such as "SUBJECT TO AVAILABILITY", "SUPPLIES WILL BE MADE AS AND WHEN SUPPLIES ARE RECEIVED" etc., will not be entertained under any circumstances and the tenders of those who have mentioned such conditions shall be treated as incomplete and accordingly the Tender will be summarily rejected.

- 8.8. For the drug formulation like Injections, Liquid orals, Tablets and Capsules, rates should be quoted only for the composition stated in the tender. Blood products should be supplied along with HIV and Hepatitis-B screening certificate, failing which the drugs will not be accepted. A copy of these Certificates duly notarized should be sent with every consignment and every invoice.
- 8.9. Supplies should be made directly by the tenderer and not through any other Agency / Dealer / Distributors.
- 8.10. The Tenderer shall allow inspection of the factory at any time during the validity of the tender / currency of the contract by a team of Experts/Officials nominated by the Tender Inviting Authority for the purpose. The Tenderer shall extend necessary cooperation to such team in inspection of the manufacturing process, quality control measures adopted etc., in the manufacture of the drugs quoted. If Company/Firm does not allow for any such inspection, their tenders will be rejected during the currency of the contract. Non compliance to any of the manufacturing process / Quality control measures brought out during such inspection render the tenderers ineligible / disqualified.
- 8.11. The Tenderer should not influence the Inspection team in any manner including providing conveyance, accommodation, food etc., any effort may result in rejection of the tender without prejudice to other conditions.

9. ACCEPTANCE OF TENDER

- 9.1. Tenders will be evaluated in accordance to the provisions of the Tamilnadu Transparency in Tenders Act, 1998 and Rules made thereunder and the criteria mentioned herein. Rate per unit inclusive of various taxes and charges (landed price) as mentioned in **Annexure-XIX** shall be worked out for determining the L1 rate (Lowest rate) (or) as per the relevant Acts / rules in force at the time of price bid opening.
- 9.2. TNMSC Ltd., reserves the right to accept or reject the tender for the supply of all or any one or more drugs of the drugs tendered for in a tender without assigning any reason.
- 9.3. TNMSC Ltd., or its authorized representative(s) has the right to inspect the manufacturing (own manufacturer / loan licensee's manufacturing facility) premises of Tenderers, before, accepting the rate quoted by them or before releasing any purchase order(s) or at any point of time during the continuance of tender and also has the right to reject the tender or terminate/cancel the purchase orders issued and/or not to place further order, based on adverse reports brought out during such inspections by any statutory authorities besides blacklisting for a period of 5 years.
- 9.4. The acceptance of the tenders will be communicated to the lowest / matched Tenderers in writing.

10. PERFORMANCE SECURITY

On being informed about the acceptance of the tender and at the time of signing the Agreement, the lowest/matched Tenderer shall pay the Performance Security as indicated below in the form of ***Demand Draft or irrevocable Bank Guarantee*** in favor of Tamilnadu Medical Services Corporation Limited. In case the Performance Security is paid in form of Bank Guarantee, the bank guarantee shall be valid for a period of 2 years from the date of communication of the acceptance letter from the Tender inviting Authority. The format of Bank Guarantee is at **Annexure-XII**. Failure to deposit the performance security will attract Clause No. 20.1 (a).

(a)	Total value of contract upto Rs.1.00 Crore	3% of the contract value (subject to a minimum of Rs. 5,000/-) at the time of entering into contract.
(b)	Total Value of contract Exceeding Rs.1.00 Crore	Rs.3.00 Lakhs Plus @ 2% of the contract value over and above Rs.1.00 Crore at the time of entering into contract.

11. AGREEMENT

11.1. The lowest/matched Tenderer shall execute an agreement on a non-judicial stamp paper of value of Rs.100/- (stamp duty to be paid by the Tenderer) within 15 days from the date of the intimation from TNMSC Ltd., The Specimen form of agreement is available in **Annexure-XI** and available in the TNMSC Ltd., Website (www.tnmsc.tn.gov.in) also.

- 11.2. The Tenderer shall not, at any time, assign, sub-let or make over the contract or the benefit thereof or any part thereof to any person or persons what so ever.
- 11.3 All notices or communications relating to and arising out of this agreement or any of the terms thereof shall be considered duly served on or given to the Tenderer if delivered to him or left at the premises, places of business or abode as provided by the tenderer.
- 11.4 If the lowest/matched Tenderer fails to execute the agreement and/or to deposit the required Performance Security within the time specified or withdraws the tender, after the intimation of the acceptance of the tender or owing to any other reasons to undertake the contract, the contract will be cancelled and the Earnest Money Deposit deposited by the tenderer along with the tender shall stand forfeited by the TNMSC Ltd., and the firm will also be liable to make for the damages/losses suffered by TNMSC Ltd., apart from blacklisting and other penal actions.

12. METHODOLOGY FOR PLACING ORDERS

For the above purpose the following procedures will be adopted

- (a) After the conclusion of Price Bid opening (Cover B), the lowest offer of the Tenderer is considered for negotiation and rate arrived after negotiation is declared as the lowest rate and that tenderer is the lowest evaluated for the drug (s) for which the tender has been invited.

- (b) The Tenderer, who has been declared as lowest tenderer for certain drug(s), shall execute necessary agreement for the supply of the tendered quantity of such drug(s) as specified in the Tender Document. On depositing the required amount as Performance Security and on execution of the agreement, such Tenderer(s) is eligible for the placement of Purchase Orders.
- (c) If two or more Tenderers are declared as lowest suppliers for the same drug(s), such Tenderers shall execute necessary agreement as specified in the Tender Document. On depositing the required amount as Performance Security and on execution of the agreement, such Tenderers are eligible for the placement of Purchase Orders.
- (d) TNMSC Ltd., will inform the lowest rate to other Tenderers who had qualified for Price Bid (Cover B) opening, inviting their consent to match with the lowest rate for the drug(s). The Tenderers who agree to match lowest rate, will be considered as Matched lowest tenderer.
- (e) The Tenderer, who agrees to match the lowest rate, shall furnish the revised breakup details of Price (Lowest Rate) in Format in **Annexure-XIX**.
- (f) Subject to para (e) above, while TNMSC Ltd., has chosen to place Purchase Orders with the Matched lowest supplier and if there are more than one such Matched lowest supplier, then the Purchase Orders for the requirement of drug(s) will be placed based on the production capacity provided by the suppliers and in

accordance with Transparency in Tenders Acts and Rules or as per relevant Acts / Rules in Force.

- (g) The Matched lowest supplier, on placement of Purchase Order, will be deemed as lowest rate supplier for the purpose of the tender and all provisions of the tender documents applicable to L1 rate Tenderer will apply mutatis mutandis to the Matched L1 supplier also.
- (h)
 - (a) In the case of purchase of drug(s) where the total quantity earmarked by the lowest / matched tenderers, is less than the total quantity required, the TNMSC Ltd., may, after placing orders with the lowest evaluated / matched Tenderers for the entire quantity earmarked by such Tenderer subject to the tenderer's ability to supply, require all the other eligible Tenderers who participated in the tender and offered a price higher than that of the lowest evaluated price, to submit revised price and quantity they would be willing to supply. TNMSC Ltd may place order for the remaining required quantity at the revised offered / quoted rates in strict ascending orders with due priority to the tenderer who offered the lowest rate.
 - (b) In addition, provision under Rule 31A of The TamilNadu Transparency in Tenders Act 1998, 2000 Tender Rules and made thereunder or any relevant Acts / Rules in force will be invoked wherever found necessary.

- (c) All the supplier(s) shall be provided with **e-purchase order(s) to the registered mail-id provided in the technical bid**. Intimation regarding placement of purchase orders shall also be provided as **SMS** to the registered mobile number provided in the technical bid for the purpose. The email-id and the mobile number provided shall be of the bidder's origin and not any other person.
- (i) The supplier, on receipt of the e-purchase order deems that the purchase quantity exceeds the production capacity declared in the tender document/agreement and the delay would occur in executing such order, shall inform the TNMSC Ltd., immediately without loss of time (within 7 days from the date of purchase order), failing which the supplier is estopped from disputing the imposition of liquidated damages, fine for the delayed supply etc.
- (j) If the lowest supplier has failed to supply the required drug(s) within the stipulated time or within the extended time, as the case may be, TNMSC Ltd., will cancel such purchase orders and on cancellation, TNMSC Ltd., will place Purchase Orders with the alternate source at the risk and cost of the defaulted supplier.
- (k) If the supplier fails to supply the drug(s) for any of the three Purchase Orders placed for the same drug(s), at any point of time, either fully or partly, within the stipulated time, TNMSC Ltd., is at liberty to place Purchase Orders either with other Tenderers (in ascending order, viz., L2,L3 and so on) at the price offered

by them or with alternate sources and in such cases the defaulted supplier is liable to indemnify TNMSC Ltd, WITH OUT ANY PROTEST OR DEMUR, for the difference in cost incurred by TNMSC Ltd., and the TNMSC Ltd., is entitled to recover the difference in cost from any amount due/payable to the defaulted supplier.

- (l) Notwithstanding anything contained in para (k) above, the supplier, after committing the default in supply either partly or fully, can inform the TNMSC Ltd., about his willingness to execute the Purchase Order during the tender period. The TNMSC Ltd., at discretion, may consider the willingness of the supplier on merit. However, such supplies will be subjected to the levy of Liquidated Damages and other penalties as stipulated in the tender document/ agreement and purchase order.
- (m) The supplier shall start supply of the Drug/Medicines required by TNMSC Ltd., at the destination mentioned in the schedule, within the period stipulated in the Purchase Order.
- (n) The Drug/Medicines supplied in excess of the ordered quantity shall not be accepted and the supplier shall take back the excess at their cost. TNMSC Ltd., will not be responsible for the loss to the supplier and will not entertain any demand/claim for excess supply.

- (o) The supplier shall supply the Drug(s) at the specified destination along with original excise invoice, Test reports of finished products for every batch, delivery Challan and other relevant documents at the destinations. Any supply without the above documents will not be accepted and the said supply will be accepted only on the date of submission of the required document (s).

However, the test reports for the raw materials used in the product to be furnished as and when called by TNMSC Ltd.,

- (p) The supplier shall take utmost care in supplying the quality Drug/Medicines and ensure that the batch number mentioned in the packages of the Drug/Medicines tally with the batch number mentioned in the Invoice(s) produced to TNMSC Ltd., for payment. Also the supplier shall ensure the quantity relevant to the Batch Number of the Drug/Medicines is mentioned in the invoice. Any variation will be viewed seriously and the payment for the supply will be released only after confirmation of the batch number by the supplier. At the discretion of the TNMSC Ltd., the variations in batch numbers in Invoices Vs. actual supply will be accepted. However, any abnormal variation may lead to Blacklisting of the product (s) at the discretion of TNMSC Ltd.,
- (q) It is the duty of the supplier to supply Drug/Medicines at the destinations mentioned in the Purchase Order and supply shall conform to the conditions mentioned in the provisions of tender documents, viz., logo, nomenclature in Tamil, etc.,

- (r) Subject to para (q) above, TNMSC Ltd., will process the invoices submitted by the supplier and the payments against supply will be made, within 30 days from the date the Drug(s) supplied has been declared of STANDARD QUALITY, by the Empanelled laboratory.
- (s) Subject to the conditions mentioned in the Purchase Order, Tender Document, Agreement executed by the supplier and here under, the Supplier is entitled for the payment against supply. In case of any discrepancy in levy of LD, Penalty, Unexecuted Fine, Short Passing of Bills shall be intimated **within 15 days** from the date of receipt of payment, failing which TNMSC Ltd., will not entertain any claim thereafter.

13. SUPPLY CONDITIONS

- 13.1. e-purchase orders along with the place of supply (destinations) will be issued to the Tenderer(s) at the discretion of the TNMSC Ltd., preferably once in a month.
- 13.2. Within 3 days from the receipt of e-purchase orders the Tenderer should upload on the website of the TNMSC Ltd., and/or inform through fax the confirmation for the receipt of the purchase order.
- 13.3. The Tenderer should also upload the details of supply schedule as specified in Annexure, on the TNMSC Ltd., website within 7 days from the receipt of the e-purchase order.

- 13.4 (a) The supplier shall supply at least **50%** of the ordered quantity within **45 days** from the date of purchase order and the balance quantity within **60 days** from the date of purchase order at the destinations mentioned in the purchase order. If the above day happened to be a holiday for TNMSC Ltd., the supply should be completed by 5.00 PM on the next working day. If the Tenderer fails to execute the supply within the stipulated time **(45 / 60 days)**, the TNMSC Ltd., without any notice/information is at its liberty to make alternative arrangement for purchase of the drugs of ARV Drugs for which the Purchase orders have been placed, from any other sources or in the open market or from any other Tenderer who might have quoted higher rates, at the risk and the cost of the defaulted supplier and in such cases the TNMSC Ltd., has every right to recover the cost and impose penalty as mentioned in Clause 19.
- (b) The supplier may continue the supply of the unexecuted quantity after **60th day upto 5 PM of 80th day**, subject to levy of appropriate Liquidated Damages as specified in clause 18.1 of the tender conditions.
- 13.5. The supplier is entitled to receive a bonus payment **@ 0.25% / 0.50% / 0.75%** flat on the order value PROVIDED the entire order is completed within **45th days / 35th days / 25th days respectively**. However, such bonus shall be paid to the extent of the least of difference in rate between L2 and L1 or the actual amount arrived then and there.

- 13.6. Supplier shall complete the earliest pending purchase order before commencing the supply of subsequent purchase orders.
- 13.7. All supplies will be scheduled for the period from the date of acceptance till the completion of the tender in installments, as may be stipulated in the Purchase Order, subject to various conditions mentioned here under. The supplied medicines and Drugs (covered in SCHEDULE "P" of Drugs and Cosmetics Act) should have the prescribed potency throughout the shelf life period as prescribed in the Drugs and Cosmetics Act 1940 and rules there under and in relevant Pharmacopoeias. All other drugs of ARV Drugs should have shelf life period of minimum 24 / 18 / 12 months from the date of manufacture as prescribed in official compendiums. Each batch of product (s) supplied should have ingredients at the lower limit of 95% at the entry level to the TNMSC Ltd., warehouses and the upper limits should be as prescribed in the official Pharmacopoeias throughout its shelf life. Failure to comply with this condition may lead to rejection of drugs at discretion of TNMSC Ltd.,
- 13.8. The Tenderer must submit an Analysis report from a Government approved Laboratory for every batch of drug along with invoice. In case of failure on part of the supplier to furnish such report, the batch of drugs will be returned back to the suppliers and he is bound to replenish the same with Govt. approved lab test report. The ARV Drugs supplied by the successful Tenderer shall be of the best

quality and shall comply with the specifications, stipulations and conditions specified in the tender.

13.9 Tenderer should supply the product, **within 30 days (Category “A”) and 40 days (Category “B”)** from the date of manufacture of that product. In case, the product is received after 30 / 40 days from date of manufacture and the product is not consumed before its expiry date, the supplier should replace the short expiry/expired quantity with fresh stock of longer shelf life, otherwise the expired product will be returned to the supplier and the value equal to the cost of expired quantity will be recovered from any dues payable.

13.10. The order **stands cancelled at the end of 80th day** from the issue of the purchase order after levying penalty on the value of unexecuted order as specified under Clause 18.2. Further, the Tenderer shall also be liable to pay other penalties as specified under Clauses 19. However if such default occurs for 3 or more purchase orders placed during the tender period, penal action like blacklisting from participating in present and future tenders of TNMSC Ltd., may be enforced by the TNMSC Ltd.,

13.11. It shall be the responsibility of the Tenderer for any shortages/damage at the time of receipt in Warehouse. TNMSC Ltd., is not responsible for the stock of drug received, for which no order is placed.

13.12. If at any time the Tenderer has, in the opinion of the TNMSC Ltd., delayed the supply of drugs due to one or more reasons related to Force Majeure events such as riots, mutinies, wars, fire, storm, tempest or other exceptional events at the manufacturing premises, the time for supplying the drugs may be extended by the TNMSC Ltd., at discretion for such period as may be considered reasonable. However such extension shall be considered only if a specific written request is made by the Tenderer within 10 days from the date of occurrence of such event with necessary documentary evidence. The exceptional events does not include the Scarcity of raw material, Increase in the cost of raw material, Electricity failure, breakdown of machineries, Labour disputes/Strikes, Insolvency, and Closure of the Factory/Manufacturing unit on any grounds etc.

13.13. The Tenderer shall take back drugs, which were supplied **beyond 30 / 40 days from the date of manufacturing** which are not utilized by the TNMSC Ltd., within the shelf life period based on mutual agreement.

13.14. The supplier shall not be liable to pay LD/penalty and forfeiture of performance security for the delay in executing the contract on account of the extension of supply period granted on the ground of force majeure events.

14. LOGOGRAMS

Logogram means, wherever the context occurs, the design as specified in **Annexure-I and Annexure-II. The name of the drug shall be mentioned in Tamil and English only.**

- 14.1. Tenders for the supply for ARV Drugs etc., shall be considered only if the Tenderer gives an undertaking that the product(s) will be prepared as per the specifications such as strength, minimum size and packed with appropriate size of the strips/blisters and with the logogram of proportionate size either printed or embossed on tablets and capsules, bottles etc., as per the design enclosed as per **Annexure-I and Annexure-II**.
- 14.2 All tablets and capsules have to be supplied in standard packing of 10 X 10 in strip or blister packing with printed logogram of proportionate size 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 unless exempted by TNMSC Ltd.,
- 14.3 Vials, Ampoules and Bottles containing the drugs tendered for should also carry the printed logogram of proportionate size unless exempted by TNMSC Ltd.,
- 14.4 Failure to supply Drugs etc., with the printed logogram of proportionate size will be treated as breach of the terms of agreement / violation of tender conditions and action will be taken to blacklist the product and/or fine will be deducted from the amount payable as per condition in Clause 18.4. However if such failure continues despite notice, will be viewed as a serious lapse. Tenderers who are not willing to agree to conditions above will be summarily rejected.

15. PACKING

- 15.1. The ARV Drugs shall be supplied in the package specified in **Annexure-IX** and **Annexure-X** and the package shall carry the logograms of proportionate size specified in **Annexure-II**. Affixing of labels in smaller size will be treated as violation of tender conditions and fine will be deducted from the amount payable as per condition in Clause 18.4.
- 15.2. (a) 2D bar coding as per GS1 standard should be done on tertiary packing of the supplies as per the specifications given in **Annexure-XV**.
- (b) 2D bar coding should be done on **secondary packing** for fixed variables as per the specifications given in **Annexure-XV (a)**.
- 15.3. The minimum size of each tablet should be 6.4 mm in diameter and the minimum size of the blister packing and strip packing should be 70mm x 30mm and 45mm x 110mm respectively. Failure to comply with this shall lead to non-acceptance of the goods besides imposition of penalties as per clause 18.4.
- 15.4. The packing in each carton shall be strictly as per the specification mentioned in **Annexure-X**. The outer carton should be of **white board** with a **minimum of 300 GSM with laminated packing** for the strips, blisters, ointments, creams etc. and for ampoules and vials should be with **white board** of **450 GSM**. Failure to comply with this shall lead to non-acceptance of the goods besides imposition of penalties

as per clause 18.4. **However in case of poor / damaged packing, necessary replacement should be provided for damaged goods.**

- 15.5. The caps of bottle preparations should not carry the name of the supplier.
- 15.6. The labels in the case of Injectable preparations should clearly indicate whether the preparations are meant for Intravenous (IV), Intra Muscular (IM), Intra Dermal (ID), Subcutaneous (SC) administration etc.,
- 15.7. The capsule shell should have the name of the drug, in addition to the logo.
- 15.8. It should be ensured that only first-hand fresh packaging material of uniform size, including bottle and vial, is used for packing.
- 15.9. All primary packing containers should be strictly conforming to the specification included in the relevant pharmacopoeia.
- 15.10. Packing should be able to prevent damage or deterioration during transit.
- 15.11. In the event of drugs of drugs supplied found to be **not as per specifications in respect of their packing and logogram**, the TNMSC Ltd., is at liberty to make alternative purchase of the drugs of ARV Drugs for which the Purchase orders have been placed from any other sources or in the open market or from any other Tenderer who might have quoted higher rates, at the risk and the cost of the supplier. In such cases the TNMSC Ltd., has every right to recover the cost and impose penalty as mentioned in Clause 18.4.

16. QUALITY TESTING

- 16.1. Samples of supplies in each batch will be chosen at the point of supply or distribution/storage points for testing. The samples will be sent to different laboratories including Government Drugs Testing Laboratory/Kings Institute for testing as decided by the TNMSC Ltd.,
- 16.2. The Drugs shall have the active ingredients at the prescribed level as indicated in official compendiums throughout the shelf life period of the drug. The samples will be drawn periodically throughout the shelf life period and if found “Not of Standard Quality”, the cost of entire batch paid will be recovered whether consumed fully/partially. Also action will be initiated for blacklisting as per clause.19 irrespective of the period of supply. The supplies will be deemed to be completed only upon receipt of the quality certificates from the laboratories. Samples which do not meet quality requirement shall render the relevant batches liable to be rejected. If the sample is declared to be “Not of Standard Quality” or spurious or adulterated or misbranded, such batch/batches will be deemed to be rejected goods.
- 16.3. In the event of the samples of ARV Drugs supplied fails in quality tests or found to be not as per specifications, the TNMSC Ltd., is at liberty to make alternative purchase of the drugs of ARV Drugs for which the Purchase orders have been placed from any other sources or in the open market or from any other Tenderer who might have quoted higher rates, at the risk and the cost of the supplier and

in such cases the TNMSC Ltd., has every right to recover the cost and impose penalty as mentioned in Clause 19.

- 16.4. The supplier shall furnish to the TNMSC Ltd., the Evidence of bio-availability and/or bio-equivalence reports for certain critical drugs upon demand.
- 16.5. The supplier shall furnish evidence of the basis for expiration dating and other stability data concerning the commercial final package on request by the TNMSC Ltd., In case of any adverse report in the field, the B.M.R/B.P.R for the particular batch of the product(s) supplied shall be produced when demanded.
- 16.6. The products should conform to the standards of IP/BP/USP/EP/JP as the case may be. In case the product is not included in the said compendiums, the supplier, upon award of the contract, must provide the reference standards and testing protocols for quality control testing. For imported drugs, respective Country's Pharmacopial standards shall be acceptable (even if the product is official in IP).
- 16.7 In case of admixture of drugs / mixing of various products in the Secondary and/or Tertiary packing, such case will be treated as a violation of tender conditions and action will be initiated as per clause 19.3.

17. PAYMENT PROVISIONS

- 17.1. No advance payments towards costs of drugs, medicines etc., will be made to the Tenderer.
- 17.2. Payments towards the Supply of ARV Drugs will be made strictly as per the rules of the TNMSC Ltd., The payment will be made either by means of Cheque or through RTGS (Real Time Gross Settlement System)/Core Banking/NEFT. The Tenderer shall furnish the relevant details in original **(Annexure-XVI)** to make the payment through RTGS/Core Banking/NEFT and the change of Bank Account during the validity of the tender will not be entertained normally.
- 17.3. All Bills / Invoices should be raised in triplicate and in the case of excisable ARV Drugs, the bills should be drawn as per Central Excise Rules in the name of Tamil Nadu Medical Services Corporation Ltd., No.417, Pantheon Road, Egmore, Chennai - 600008 or in the name of any other authority as may be designated (or) as per the relevant Acts / rules in force at the time of billing.
- 17.4 (i) The Warehouses of TNMSC Ltd., has been divided in to 5 Divisions i.e., Chennai Division, Coimbatore Division, Madurai Division, Thiruchirapalli Division and Tirunelveli Division. The details of warehouse in each division are as follows.

- (a) Chennai Division - Anna Nagar, K.K.Nagar, Kancheepuram, Vellore, Villupuram and Tiruvallur.
- (b) Coimbatore Division - Coimbatore, Dharmapuri, Erode, The Nilgris, Salem, Krishnagiri, Namakkal and Tiruppur.
- (c) Madurai Division – Madurai, Dindigul, Ramanathapuram, Sivagangai, Theni and Virudhunagar.
- (d) Thiruchirapalli Division - Thiruchirapalli, Cuddalore, Thanjavur, Thiruvarur, Tiruvannamalai, Pudukottai, Karur, Perambalur and Nagapattinam.
- (e) Tirunelveli Division – Tirunelveli, Nagercoil and Thoothukudi.

Payments for supply will be considered only after supply of minimum 50% of the total ordered quantity in total and 40% in each divisions in the individual Purchase Order, provided reports of Standard Quality on samples testing are received from Government Analyst or Approved Laboratories of TNMSC Ltd.,

- (ii) However, in case of cancellation of a particular purchase order due to failure in delivery, payment for part supplies less than 50% of the purchase order quantity on the date of cancellation of the purchase order may be considered for release of payment subject to the following:

- (a) If the Tenderer have supplied at least 50% of the quantity ordered in the subsequent purchase order within **70 days** from the issue of such purchase order.
 - (b) If further purchase order is not placed with the supplier due to any reason, not attributable to the supplier, the amount eligible will be paid after **70 days** from the date of last supply.
 - (c) The payment for part supply as mentioned above will subject to the deduction of liquidated damages, penalty towards unexecuted quantity, risk and cost etc., as per the tender conditions.
- (iii) In all other cases, which are not covered under para (i) and (ii) above, the issue related to the settlement of payments will be decided by the TNMSC Ltd., on merits of the case subject to various terms and conditions of the tender.

17.5. If at any time during the period of contract, the price of tendered drugs is reduced or brought down by any law or Act of the Central or State Government or by the Tenderer himself, the Tenderer shall be bound to inform the TNMSC Ltd., immediately about such reduction in the contracted prices. Tender Inviting Authority is empowered to unilaterally effect such reduction as is necessary in rates in case the Tenderer fails to notify or fails to agree for such reduction of rates.

17.6. (a) In case of any increase or decrease in the taxes, such as customs duty, GST (or) as per the relevant Acts / rules in force after the date of submission of tenders and during the tender period, such variation in the taxes will be to the account of the TNMSC Ltd., For claiming the additional cost on account of the increase in taxes, the Tenderer should produce the proof of having paid additional amount on this account on the goods supplied to TNMSC Ltd., from the concerned authorities and also must claim the same in the invoice separately. However the basic price structure and the price of the Drugs approved under the tender shall not be altered. Similarly if there is any reduction in the taxes and statutory levies as notified by the Govt., after the date of submission of tender, the Tenderer will be paid based on the unit rate worked out on the basis of the reduced taxes / statutory levies without any change in the basic price or the price structure of the drugs approved under the tender. Any increase or decrease in taxes and statutory levies will be considered based on the notification issued by the Government.

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

18. **LIQUIDATED DAMAGES AND OTHER PENALTIES:**

- 18.1. If the supply reaches the designated places or District Drug Warehouse between 5 PM of the 60th day and 5 PM of 70th days from the date of issue of the purchase order, a liquidated damages will be levied at 0.25% per day and between 71st day and 5.00 PM of 80th day a liquidated damage will be levied at 0.50% per day irrespective of the fact that whether the TNMSC Ltd., has suffered any damage/loss or not, on account of delay in effecting supply. If the due day happens to be a holiday the supply will be accepted on the next working day without any penalty.
- 18.2 If there are any unexecuted orders after 5 PM of 80th day /upto the date of delivery extension granted whichever falls later (as the case may be), from the date of purchase order, the order shall stand cancelled automatically after levying penalty @ 25% on the value of unexecuted order and such penalty is recoverable from any amount payable to the supplier. In case of alternate purchase effected due to unexecution, the differential cost incurred or the unexecuted fine whichever is higher will be levied.
- 18.3. If the supply is received in damaged condition it shall not be accepted. In case of damage in the packing, the supply will be accepted only after levying penalty to the extent of damaged value of supply received at the destination place. ***In case of damaged/broken goods in transit unexecuted penalty will be relaxed for a maximum of 2% on each order quantity.***

18.4. All the Tenderers are required to supply the product(s) with printed logogram of appropriate size on the strips, blisters, vials, ampoules & bottles and with prescribed packing specification. If there are any deviation in these Tender conditions, action will be taken to blacklist the product and/or a separate damages will be levied @ 2% of value of the defaulted quantity irrespective of the Tender Inviting Authority having actually suffered any damage/loss or not, without prejudice the rights of alternative purchase specified in Clause No.15.11 and 14.4.

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

19.1. If the samples do not conform to statutory standards, the Tenderer will be liable for relevant action under the existing laws and the entire stock in such batch has to be taken back by the Tenderer within a period of 30 days of the receipt of the letter from the TNMSC Ltd., Such stock shall be taken back at the expense of the Tenderer. The TNMSC Ltd., has the right to destroy such “NOT OF STANDARD QUALITY DRUGS” if the Tenderer does not take back the goods within the stipulated time. The TNMSC Ltd., will arrange to destroy the “NOT OF STANDARD QUALITY DRUGS” after the expiry of 30 days mentioned above without further notice. **The tenderer should not claim the quantity/ amount for the rejected goods after this period.** TNMSC Ltd., will incinerate these rejected goods and incineration cost will be recovered from the supplier on the basis of weight (in Kg) of rejected Not of Standard Quality (NSQ) Drugs. The

rates for the above disposal will be the same on the rates fixed by TNMSC Ltd., for the waste management for the Govt. Hospitals in TamilNadu.

- 19.2. If any of the Drug/Medicines supplied by the Tenderer have been partially or wholly used or consumed after supply and are subsequently found to be in bad odour, unsound, inferior in quality or description or otherwise faulty or unfit for consumption, **then the contract price for the quantity not consumed and informed to take back, will be recovered from the tenderer, if payment have already been made. In other words the Tenderer will not be entitled for any payment whatsoever for the drugs found to be of “NOT OF STANDARD QUALITY” which is not consumed and informed to take back and the Tender Inviting Authority is entitled to deduct the cost of such batch of drugs of quantity returned will be made from any amount payable to the Tenderer.** On the basis of the nature of failure, action will be initiated to blacklist the product/supplier.
- 19.3. For the supply of Adulterated/Spurious/Misbranded drugs to TNMSC Ltd., the firm/company shall be blacklisted by TNMSC Ltd., and no further supplies shall be accepted from the firm/company. The Tenderer shall also not be eligible to participate in tenders of Tender Inviting Authority of TNMSC Ltd., for Supply of ARV Drugs for a period of 5 years from the date of blacklisting. In case of supply of NOT OF STANDARD QUALITY drug(s) to TNMSC Ltd., the product shall be blacklisted by TNMSC Ltd., and no further supplies shall be accepted for the

particular drug(s). The Tenderer shall also not be eligible to participate in tenders of TNMSC Ltd., for supply of such ARV Drugs for a period of 2 years from the date of blacklisting. In addition, the Director of Drugs Control of concerned State will be informed for initiating necessary action on the Tenderer in their state. Performance Security paid will also be forfeited without any intimation. In case of supply of admixed product(s) to TNMSC Ltd., the product(s) shall be blacklisted by TNMSC Ltd., and no further supplies shall be accepted for the particular drug(s). The Tenderer shall also not be eligible to participate in tenders of TNMSC Ltd., for supply of such ARV Drugs for a period of 2 years from the date of blacklisting. In addition, the Director of Drugs Control of concerned State will be informed for initiating necessary action on the Tenderer in their state. Performance Security paid will also be forfeited without any intimation.

- 19.4. The Tenderer shall furnish the source of procurement of raw material utilized in the formulations, if required by the TNMSC Ltd., The TNMSC Ltd., reserves the right to cancel the purchase orders, if the source of supply is not furnished.
- 19.5. The decision of the TNMSC Ltd., or any officer authorized by him, as to the quality of the supplied drugs, medicines etc., shall be final and binding.
- 19.6. The TNMSC Ltd., will be at liberty to terminate, without assigning any reasons thereof, the contract either wholly or in part on 30 days-notice. The Tenderer will

not be entitled for any compensation whatsoever in respect of such termination besides forfeiture of Performance Security.

- 19.7. For infringement of the stipulations of the contract or for other justifiable reasons, the contract may be terminated by the TNMSC Ltd., and the Tenderer shall be liable to pay for all losses sustained by the TNMSC Ltd., in consequence of the termination which may be recovered personally from the Tenderer or from his properties, as per rules besides forfeiture of Performance Security.
- 19.8. Non-performance of any of the contract conditions and provisions will disqualify a firm from participating in the tender for the next 2 years besides forfeiture of Performance Security.
- 19.9. In the event of making Alternative Purchase, as specified in Clause 13.4, Clause 15.11 and in Clause 16.3 penalty will be imposed on the supplier. The excess expenditure over and above contracted prices incurred by the TNMSC Ltd., in making such purchases from any other sources or in the open market or from any other Tenderer who has quoted higher rates and other losses sustained in the process, shall be recovered from the Performance Security or from any other money due and become due to the supplier and in the event of such amount being insufficient, the balance will be recovered personally from the supplier as per rules.

19.10. In all the above conditions, the decision of the TNMSC Ltd., shall be final and binding.

20. BLACK LISTING IN THE EVENT OF WITHDRAWAL FROM THE TENDER, AND NON-ADHERENCE TO THE QUALITY STANDARDS AND SUPPLY SCHEDULE

20.1. BLACKLISTING OF PRODUCT/TENDERER ON WITHDRAWAL OF TENDER

(a) If the Tenderer(s) fails to execute the agreement / to deposit performance security / to perform the obligations under the tender conditions / commits default in the performance of the contract, such Tenderers will be blacklisted for a period of 2 years by TNMSC Ltd., from the date of intimation besides forfeiture of Performance Security.

(b) The Tenderers who have withdrawn after submitting their bids in the tender and before TNMSC Ltd., finalizes and places the purchase orders either fully or partially, will be blacklisted for a period of 2 years from the date of intimation by TNMSC Ltd., apart from forfeiture of the Performance Security /EMD.

BLACKLISTING FOR QUALITY FAILURE

20.2.1. Quality Test by the Empanelled Laboratories of TNMSC Ltd.,

(a) Each and every batch of Drug/Medicines supplied by the supplier shall be subjected to quality test by the Empanelled laboratories.

- (b) The samples are collected from the Warehouses from each batch of supply of the same drugs and after eliminating the common batch, samples shall be taken in random, decoded and will be sent to the empanelled testing laboratories for testing the quality of drugs.
- (c) If such sample passes quality test in all respects, TNMSC Ltd., will instruct its Warehouses to issue such drugs of drugs to various hospitals/Institutions.
- (d) Such quality passed batches if received after declaration of result of the earlier supply, the same will be again subjected to testing and the latest report of that particular batch will prevails upon the earlier results and binding on the entire quantity of the batch supplied and recovery will be made for the entire quantity of that batch irrespective of purchase order date or date of supply etc.
- (e) If the sample fails in quality test and report is received certifying that sample is “NOT OF STANDARD QUALITY”, one more sample shall be drawn from the same batch and to be sent to Government Laboratory for quality testing.
 - (i) If such sample passes the quality test as per the report of Government Laboratory, the drugs representing the sample shall be qualified for issue to various Institutions.

- (ii) If such sample fails in the quality test, as per the report of the Government Laboratory, the drugs of the batch are not qualified for issue and the supplier shall take back the drugs supplied in that batch, besides taking other actions as per the Tender conditions by TNMSC Ltd.,
- (iii) If such Sample fails in quality test for ASSAY content of less than 50% as per the Government Analyst report, such product of the tenderer will be blacklisted for two years.
- (iv) However, TNMSC Ltd., reserves the right to reject the drugs based on reports from empanelled laboratories with the applicable penal provisions.
- (f) If 3 batches of a particular drug supplied by the supplier is reported to be failing in ASSAY content (above 50% but below prescribed limit) / Sterility / BET / Toxicity / and or other parameters, then the particular drug of the firm shall be blacklisted after observing procedure laid down in Para 20.2.4 besides forfeiture of Performance Security of that particular product(s).
- (g) In all the cases the reports received from the Government Drug Testing Laboratory/decision of TNMSC Ltd will be conclusive and final and binding on the suppliers.

20.2.2. Quality Test by Statutory Authorities:

- (a) On complaint from Drug Inspector(s) during their Test of statutory sampling, that the particular drug has been reported to be of “NOT OF STANDARD QUALITY”, the issue of available stock of the particular drug will be stopped. Further, the available stock of the product in hospitals will be retrieved. If the sample is reported to have less than 50% of content, the particular product will be **blacklisted for 2 years from the date of intimation of blacklisting.**

- (b) If 3 batches of a particular drug supplied by the supplier is reported to be failing in **ASSAY content (above 50% but below prescribed limit) and/or other parameters.** then the particular drug of the firm shall be blacklisted for a period of **2 years** from the date of intimation after observing procedure laid down in Para 20.2.4.

- (c) If a single batch of any product(s) supplied by the company/firm declared as Adulterated/spurious/ Misbranded by the Government Authorities during the shelf life of the product supplied irrespective of tender period, the company/firm shall be blacklisted for a period of **5 years from the date of intimation** after observing procedure laid down in Para 20.2.4.

20.2.3 BLACKLISTING OF THE SUPPLIER FOR QUALITY FAILURE:

- (a) In case of any sample even in one batch, declared as Adulterated/spurious/ Misbranded by the Government Authorities during, the company/firm shall be blacklisted for a period of **5 years** from the date of intimation besides forfeiture of performance security in full after observing the procedure laid down in Para 20.2.4.
- (b) If the supplier supplied more than one drug and 50% of such drugs are blacklisted, the firm is liable to be blacklisted for a period of 2 years from the date of intimation after observing the procedure laid down in Para 20.2.4.

20.2.4 PROCEDURE FOR BLACKLISTING

- (i) On receipt of report from Govt. Analyst/Drug Testing Laboratory indicating that a particular Drug/Drug is **“NOT OF STANDARD QUALITY/ ADULTERATED/ SPURIOUS/ MIS-BRANDED** (As the case may be), a show cause notice shall be issued to the supplier calling for explanation within 7 days from the date of notice. On receipt of explanation from the supplier, the Managing Director, TNMSC Ltd., may take appropriate action on merits of the case and impose penalty **@ 25% of the value of the failed batch (or) 7.5% of the total supply value made in the particular purchase order (which ever higher) and / or** the blacklisting of the particular drug(s) of the product/company or firm as deemed fit besides forfeiture of Performance Security.

- (ii) If a particular drug of the drug has been blacklisted according to the procedure stated above, the supplier is not eligible to participate in any of the tenders for that particular drug floated by the TNMSC Ltd., until the period of blacklisting is over.
- (iii) If a supplier company/firm is blacklisted according to the procedure stated above, such supplier is not eligible to participate in any of the tenders floated by the TNMSC Ltd., until the period of blacklisting is over.

20.3 BLACKLISTING FOR NON-SUPPLY:

Notwithstanding various actions and penalties for non-supply and/or delayed supply of the ARV Drugs as stipulated in the terms and conditions of the tender, the TNMSC Ltd., shall take action against the supplier as follows:

- (a) If the supplier fails to execute at least **70%** of the ordered quantity as mentioned in a single Purchase order and such part supply for **any three Purchase orders of the same drug,** then the product of the supplier will be blacklisted and becomes ineligible to participate in any of the tenders for that particular drug(s) by TNMSC Ltd., for a period of **2 years** from the date of intimation for blacklisting besides forfeiture of performance security of that product(s).

- (b) If the supplier supplies more than one drug and 50% of such drugs are blacklisted, the firm is liable to be blacklisted for a period of **2 years from the date of intimation** besides forfeiture of performance security in full.

20.4. Purchase orders, if any, already issued before taking any blacklisting action or 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.

20.5. The blacklisting of particular product or company/firm will be done without prejudice to other penalties which may be imposed as per the conditions of Tender documents and also to other actions which may be initiated under Drugs and Cosmetics Act 1940 or any other law of Land. TNMSC Ltd., will display names of such blacklisted product(s) and company/firm on its website and also circulate the same among other state Government / Central Government and its Drug procurement agencies including respective State Drugs Control Department where the company or firm is located.

21. SAVING CLAUSE

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

22. RESOLUTION OF DISPUTES

- (i) The TNMSC Ltd., and the supplier shall make every effort to resolve, amicably by direct informal negotiation any disagreement or dispute arising between them under or in connection with the contract,
- (ii) In case of a dispute or difference arising between the TNMSC Ltd., and a supplier relating to any matter arising out of or connected with this agreement, such dispute or difference shall be settled in accordance with the Arbitration and Conciliation Act, 1996. The venue of arbitration shall be Chennai alone.
- (iii) In case of discrepancy in payment provisions such as levy of Liquidated damages, Unexecuted fine, short passing of bills etc., the same shall be represented to TNMSC Ltd., within **15 days** from the date of receipt of payment, failing which TNMSC Ltd., will not entertain any claim thereafter.

23. APPEAL:

This tender is governed by the provisions of Tamilnadu Transparency in Tenders Act 1998 and the Rules there under.

- (i) Any Tenderer aggrieved by the order passed by the Tender Accepting Authority under section 10 of the said Act, may appeal to the Government within ten days from the date of receipt of order and the Government shall dispose the appeal within fifteen days from the date of receipt of such appeal.

- (ii) No Appeal shall be preferred while the tender is in process and until tender is finalized and Notification of award is issued by the TNMSC Ltd.,

24. CONTACTING THE TNMSC LIMITED BY THE BIDDER:

- (i) No bidder shall contact the *TNMSC Ltd.*, on any matter relating to its bid, from the time of bid opening to the time the contract is awarded.
- (ii) Any effort by a bidder to influence the *TNMSC Ltd.*, in the *Purchaser's* bid evaluation, bid comparison or contract award decisions may result in rejection of the bidder's bid.
- (iii) The bidder shall not make any attempt to establish unsolicited and unauthorized contact with the Tender Accepting Authority, Tender Inviting Authority or Tender Scrutiny Committee after opening of the bids and prior to the notification of award and any attempt by any bidder to bring to bear extraneous pressures on the Tender Accepting Authority, Inviting Authority or Tender Scrutiny Committee, shall be sufficient reason to disqualify the bidder.
- (iv) Notwithstanding anything contained in clause (iii) above the Tender Inviting Authority or the Tender Accepting Authority, may seek bonafide clarifications from bidders relating to the bids submitted by them during the evaluation of bids.

25. FRAUDULENT AND CORRUPT PRACTICES:

(1) FOR BIDDERS:

It is purchaser's policy to require that the bidders, suppliers and contractors and

their authorized representatives/agents observe the highest standard of ethics during the procurement and execution of such contracts. *(In this context, any action taken by a bidder, supplier, contractor, or by their authorized representatives/agent, to influence the procurement process or contract execution for undue advantage is improper)* In pursuance of this policy, the purchaser;

(a) defines, for the purposes of this provision, the terms set forth below as follows:

(i) “corrupt practice” is the offering, giving, receiving or soliciting, directly or indirectly, of anything of value to influence improperly the actions of another party (*“another party” refers to a public official acting in relation to the procurement process or contract execution*). *In this context, “public official” includes staff and employees of other organizations taking or reviewing procurement decisions.*

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

or contract execution; and the “act or omission” is intended to influence the procurement process or contract execution).

- (iii) “collusive practice” is an arrangement between two or more parties designed to achieve an improper purpose, including to influence improperly the actions of another party [*“parties” refers to participants in the procurement process (including public officials) attempting to establish bid prices at artificial, non competitive level*].
- (iv) “coercive practice” is impairing or harming, or threatening to impair or harm, directly or indirectly, any party or the property of the party to influence improperly the actions of a party (a *“party” refers to a participant in the procurement process or contract execution*).
- (v) “obstructive practice” is
 - (aa) deliberately destroying, falsifying, altering or concealing of evidence material to the investigation or making false statements to investigators in order to materially impede a investigation into allegations of a corrupt, fraudulent, coercive or collusive practice; and/or threatening, harassing or intimidating any party to prevent it from

disclosing its knowledge of matters relevant to the investigation or from pursuing the investigation; or

(bb) acts intended to materially impede the exercise of the purchaser's inspection and audit rights provided for under sub-clause (e) below.

- (b) will reject a proposal for award if it determines that the bidder considered for award has, directly or through an agent, engaged in corrupt, fraudulent, collusive, coercive or obstructive practices in competing for the contract in question;
- (c) will cancel the contract if the purchaser determines at any time that the bidder, supplier and contractors and their sub contractors engaged in corrupt, fraudulent, collusive, or coercive practices.
- (d) will sanction a firm or individual, including declaring ineligible, either indefinitely or for a stated period of time, to be awarded a contract if it at any time determines that the firm has, directly or through an agent, engaged in corrupt, fraudulent, collusive, coercive or obstructive practices in competing for, or in executing, a contract; and
- (e) will have the right to inspect the accounts and records of the bidders, supplier, and contractors and their subcontractors/authorized

representatives and to have them audited by auditors appointed by the purchaser.

(2) FOR SUPPLIERS:

If the TNMSC Ltd., determines that a Supplier has engaged in corrupt, fraudulent, collusive, coercive or obstructive practices, in competing for or in executing the Contract, then the TNMSC Ltd., may, after giving 7 days notice to the Supplier, terminate the Supplier's employment under the Contract and cancel the contract, and the procurement will be made at the risk and cost of the supplier besides blacklisting the bidder for 5 years with forfeiture of Performance Security apart from other penal actions.

(a) For the purposes of this Sub-Clause:

- (i) "corrupt practice" is the offering, giving, receiving or soliciting, directly or indirectly, of anything of value to influence improperly the actions of another party;
- (ii) "fraudulent practice" is any act or omission, including a misrepresentation, that knowingly or recklessly misleads, or attempts to mislead, a party to obtain a financial or other benefit or to avoid an obligation;
- (iii) "collusive practice" is an arrangement between two or more parties designed to achieve an improper purpose, including to influence improperly the actions of another party;

- (iv) “coercive practice” is impairing or harming, or threatening to impair or harm, directly or indirectly, any party or the property of the party to influence improperly the actions of a party;
- (v) “obstructive practice” is
 - (aa) deliberately destroying, falsifying, altering or concealing of evidence material to the investigation or making false statements to investigators in order to materially impede a purchaser investigation into allegations of a corrupt, fraudulent, coercive or collusive practice; and/or threatening, harassing or intimidating any party to prevent it from disclosing its knowledge of matters relevant to the investigation or from pursuing the investigation; or
 - (bb) acts intended to materially impede the exercise of the purchaser’s inspection and audit rights provided for.

26. JURISDICTION

In the event of any dispute arising out of the tender such dispute would subject to the jurisdiction of the Civil Court within the city of Chennai only and **official communicating language shall be Tamil (or) English, even if any dispute is raised in any other fora.**

DECLARATION

I do hereby declare that I will supply the **ARV Drugs** as per the designs in Primary, Secondary packing and labels given in enclosures to this Annexure and as per the instructions given in this regard.

Signature of the Tenderer
Name in capital letters with Designation

Attested by Notary Public.

ANNEXURE-II
Ref. Clause No. 4.1 (n) (ii) and 14.1

DECLARATION

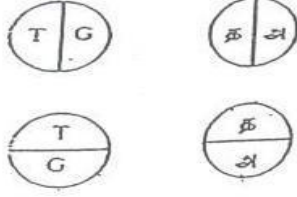
I / We do hereby declare that I will supply the **ARV Drugs** by affixing the logo for Secondary / Primary packing for the imported drugs along with Brand / trade Names as per the designs given in enclosures to this Annexure and as per the instructions given in this regard.

Signature of the Tenderer
Name in capital letters with Designation

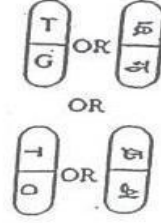
Attested by Notary Public.

ENCLOSEURE-I TO ANNEXURE-I and II – REFER CLAUSE NO.4.1.(n)
DESIGN FOR

TABLET



CAPSULE



DESIGN For STRIP

பாறாசிட்டு மால் 500 மி.சி.	PARACETAMOL 500 mg	PARACETAMOL 500 mg
	TAMILNADU GOVERNMENT SUPPLIES NOT FOR SALE	T G
பாறாசிட்டு மால் 500 மி.சி.	பாறாசிட்டு மால் 500 மி.சி.	PARACETAMOL 500 mg
	தமிழ்நாடு அரசு வழங்குவது விற்பனைக்கு அல்ல	த அ
பாறாசிட்டு மால் 500 மி.சி.	PARACETAMOL 500 mg	PARACETAMOL 500 mg
	TAMILNADU GOVERNMENT SUPPLIES NOT FOR SALE	PARACETAMOL 500 mg
பாறாசிட்டு மால் 500 மி.சி.	பாறாசிட்டு மால் 500 மி.சி.	PARACETAMOL 500 mg
	தமிழ்நாடு அரசு வழங்குவது விற்பனைக்கு அல்ல	PARACETAMOL 500 mg

REAR SIDE
MANUFACTURED BY

MFC. LICENCE NO
BATCH NO
DATE OF MANUFACTURE
DATE OF EXPIRY

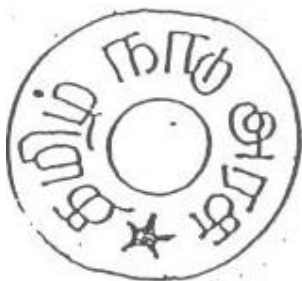
SCHEDULE

NOTE:
BRAND NAME OF THE DRUG
SHOULD NOT BE
PRINTED ANY WHERE

ENCLOSURE-III TO ANNEXURE-II and ANNEXURE-II (A)

SPECIMEN LABEL FOR
OUTER CARTON (20cm x 15cm)

**TAMIL NADU GOVT. SUPPLY
NOT FOR SALE**



~~~~~  
**DOLUTEGRAVIR TAB 50MG**  
~~~~~

Batch. : xxxxxxx
Mfg Date: JUN- 2022
Exp Date: MAY -2024

Quantity Packed:

Manufactured by:

DECLARATION

I/We M/s. _____ represented by its Proprietor / Managing Partner /
Managing Director having its Registered Office at
_____ and its Factory Premises at
_____ do declare that I/We have
carefully read all the conditions of tender in Ref.No.011/M(P)/ARV-DRUG/TNMSC/2022,
Dated.05.07.2022 for supply of **ARV Drugs** under rate contract system for one year from the date
of acceptance floated by the TamilNadu Medical Services Corporation Ltd., Chennai - 600008 and
accepts all conditions of the Tender.

I/We declare that we possess the valid licence and GMP Certificate as per revised
Schedule-'M'/WHO-GMP issued by the Competent Authority and complies and continue to comply
with the conditions laid in Schedule M of Drugs & Cosmetics Act, 1940 and the Rules made
thereunder. I/We furnish the particulars in this regard in enclosure to this declaration.

I am/we are aware of the Tender Inviting Authority's right to forfeit the Earnest Money
Deposit and/or Security Deposit and blacklisting me/us for a period of 5 years if, any information
furnished by us proved to be false at the time of inspection and not complying the conditions as
per Schedule M of the said Act for a period of 5 years.

	Signature	:
Seal	Name & Address	:

To be attested by the Notary.

Enclosure to Annexure – III Clause 4(1) (j)

DECLARATION FOR COMPLIANCE OF cGMP/WHO-GMP
(In Case of Loan Licensee, the following details to be provided for the Manufacturing Facility)

01. Name and Address of The Firm :
02. Name of Proprietor / Partner / Director :
03. Name and Designation of Person Present :
04. GMP Certificate **As per Revised Schedule “M”/WHO GMP**
05. Details of Licenses Held With Validity :
06. Number of Workers Employed :Ladies :
Gents :
07. Whether Workers Provided with Uniform : Yes / No
08. Whether Medical Examination done
for the Workers : Yes / No
09. **Hygienic Condition**
- (I) Surrounding : Satisfactory / Not Satisfactory
- (II) Production Areas : Satisfactory / Not Satisfactory
- (III) Other Areas : Satisfactory / Not Satisfactory
10. Provision For Disposal of Waste : Yes / No
11. Heating System : Yes / No
12. Whether Benches Provided in all
Working Area : Yes / No

13. Water Supply

- (A) Source :
- (B) Storage Condition : Satisfactory / Not Satisfactory
- (C) Testing
(With reference to Pathogenic Organization) : Yes / No
- (D) Cleaning Schedule In Water Supply
System With Proper Records : Yes / No
- (E) Type of Machinery installed as to Semiautomatic
or Fully Automatic plant for water purification system
along with cost and whether this is working, and if so
the flow rate of Pharmaceutical water to must the
requires preparation :

**14. Air handling system along with list of machine
and cost of the unit. Separately for sterile and
non sterile preparation :**

15. Whether the pollution control clearance is valid for
Air and Water and if so the period upto which valid
(copy of the certificate to be enclosed) :

16. Raw Material Storage Area
(Storage Facilities / Hygienic Condition) :

- (I) Quarantine : Provided / Not Provided
- (II) Passed Materials : Provided / Not Provided
- (III) Rejected Materials : Provided / Not Provided

17. Finished Product Storage Area
(Hygienic / Storage) :

- (I) Quarantine : Provided / Not Provided
- (II) Released Material : Provided / Not Provided

18. Details of Technical Staff

<u>Name</u>	<u>Qualification</u>	<u>Experience</u>
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For Manufacturing :

For Testing :

19. Testing Facilities (List of Equipments to be furnished Separately in the format to meet the bench mark vide Annexure)

Chemical Method : Yes / No

Instrumental : Yes / No
(Type of Instrument Provided as indicated in Annexure)

Biological : Yes / No

Micro Biological : Yes / No

Animal Testing : Yes / No

20. Remarks

(A) Whether Products Quoted to TNMSC are Endorsed in the Licence : Yes / No

(B) Whether the drugs Quoted to TNMSC have been Manufactured Earlier (Last 3 Years) : Yes / No

If Yes, Details Like

Sl.No	Date of Manufacturer	Name of the Drug	Batch No.	Batch Size	Date of Release

(C) Production Capacity (Section Wise)

PRODUCTION CAPACITY:**Tablet Section**

Type of Equipments (1)	No. of Equipments (2)	Production Capacity of all the Equipments in column 2 per shift (3)	No of shift (4)	Production Capacity allotted for TNMSC (5)
Planetary mixer				
Fluidized bed drier				
Tray drier				
Mechanical shifter				
Multi mill				
Tablet compression machine				
1) With _____ number of station				
2) With _____ number of station				
3) With _____ number of station				
4) With _____ number of station				
Coating pan.				
Blister Packing machine				
Strip packing machine				

Capsule Section

Type of Equipments (1)	No. of Equipments (2)	Production Capacity of all the Equipments in column 2 per shift (3)	No of shift (4)	Production Capacity allotted for TNMSC (5)
Double cone blender				
Automatic capsule filling machine				

Type of Equipments (1)	No. of Equipments (2)	Production Capacity of all the Equipments in column 2 per shift (3)	No of shift (4)	Production Capacity allotted for TNMSC (5)
Semi-Automatic Capsule filling machine				
Hand filling machine				
Blister packing machine				
strip packing machine				

Parenteral Section

Type of Equipments (1)	No. of Equipments (2)	Production Capacity of all the Equipments in column 2 per shift (3)	No of shift (4)	Production Capacity allotted for TNMSC (5)
Small volume Parenteral				
Mixing Vessel				
Laminar Flow unit				
Filtration unit				
Ampoule filling machine (with No of head)				
Vial filling Machine (with No of head)				
Vial sealing machine				
Powder filling machine				
Autoclave for terminal Sterilization				
Ampoule labeling machine				
Vials labeling machine				

Large Volume Parenterals

Type of Equipments (1)	No. of Equipments (2)	Production Capacity of all the Equipments in column 2 per shift (3)	No of shift (4)	Production Capacity allotted for TNMSC (5)
Mixing vessel				
Filtration Unit.				
Filling Machine Autoclave for terminal Sterilization				
Labeling Machine				

Ointment/ Cream

Type of Equipments (1)	No. of Equipments (2)	Production Capacity of all the Equipments in column 2 per shift (3)	No of shift (4)	Production Capacity allotted for TNMSC (5)
Stream jacket vessel for mixing				
Ointment/cream filling machine				

Liquid Section

Type of Equipments (1)	No. of Equipments (2)	Production Capacity of all the Equipments in column 2 per shift (3)	No of shift (4)	Production Capacity allotted for TNMSC (5)
Bottle washing machine				
SS tank with capacity				
Filter press				
Colloidal mill				
Bottle Filling Machine				
Labeling Machine				

External Preparation

Type of Equipments (1)	No. of Equipments (2)	Production Capacity of all the Equipments in column 2 per shift (3)	No of shift (4)	Production Capacity allotted for TNMSC (5)
Mixing Vessel				
Filling machine				
Labeling machine				

- (D) Any, Not Of Standard Quality : Yes / No
Reports Of Product Quoted/
Approved By TNMSC
(If Not, Nil Statement)
- (E) Any Prosecution After : Yes / No
Submission of Tender Documents.
(If Not, Nil Statement)
- (F) Chances Of Cross Contamination : Yes / No
at Raw Materials/In Process/
Finished Product Stages And Steps/Facilities
- (G) Validation of Equipments done : Yes / No
- (H) Cleaning Schedule
- (I) For Premises :
- (II) For Equipments :
- (I) Adverse Reaction, If Any and :
Reported

Sl.No.	Description	Remarks
1	Whether any drug(s) manufactured by the tenderer has / have been recalled during last five years? If yes given details	

Sl.No.	Description	Remarks
2	What are the results of investigations on the recalled drug(s)?	
3	What action have been taken to prevent recurrence of recall of drug(s) on that particular account?	

(J) Complaints Received If Any :
and Steps taken.

Sl.No.	Description	Remarks
1	Whether any drug(s) manufactured by the tenderer has / have been recalled during last five years? If yes given details	
2	What are the results of investigations on the recalled drug(s)?	
3	What action have been taken to prevent recurrence of recall of drug(s) on that particular account?	

Signature and Seal of
Proprietor / Partner / Director

To be attested by the Notary.

Annexure

Sl. No.	Name of the Instruments	No. of Instruments	Cost of Instruments	Whether it is in working condition
(1)	(2)	(3)	(4)	(5)
1	Analytical Balance			
2	Infra-Red Spectrometer			
3	Karl Fisher Tritator			
4	Melting Point			
5	Brookfield Viscometer			
6	Polarimeter			
7	Autoclave			
8	Refractometer			
9	Sampling Booth			
10	UV-Vis Spectrometer			
11	HPLC			
12	Muffle Furnace			
13	Fuming Cupboard			
14	Micrometer			
15	Dissolution Tester			
16	Disintegration Tester			
17	Friability Tester			

Sl. No.	Name of the Instruments	No. of Instruments	Cost of Instruments	Whether it is in working condition
(1)	(2)	(3)	(4)	(5)
18	Vernier Calipers			
19	IR Balance			
20	Hardness Tester			
21	Leak Test Apparatus			
22	Laminar Air Flow			
23	BOD Incubator			
24	Vacuum oven			
25	Bulk Density Apparatus			
26	Water Activity Meter			
27	Anaerobic System			
28	Gas Chromatograph			
29	LAL Kit			
30	Sterility Test Kit			
31	Particle Counter			
32	Air Sampler			
33	Flame Photometer			
34	Tap Density Tester			

ANNEXURE-IV
Ref. Clause No. 2 (f)

DECLARATION

I _____ Managing Director /
Director / Partner / Proprietor of M/s. _____ having its
manufacturing or import unit / registered office
at _____ do hereby declare that we
have not been blacklisted either by Tender Inviting Authority or by any State Government or
Central Government Organization for the following products quoted in the tender. We are eligible
to participate in the tender Ref. No. **011/M(P)/ARV-DRUG/TNMSC/2022, Dated.05.07.2022** for
the following products.

Sl. No.	Drug Code	Name of the Drug

M/s. _____

Company seal

To be attested by the Notary.

ANNEXURE-V
Ref. Clause No.7.1 (a)

DETAILS OF E.M.D. SUBMITTED

We herewith submit the E.M.D. of Rs._____ in the form of Demand
Draft bearing No._____ Dated: _____ drawn on
_____ Bank _____ Branch in
favour of TNMSC for the following drugs of drugs.

Sl. No.	Drug Code*	Name of the Drug	Amount of E.M.D.
		Total :	

*in ascending order as in **Annexure-IX**.

Signature & Seal

NOTARISED UNDERTAKING

(In 20- Rupees stamp paper)

I _____, S/o _____, Proprietor / Partner / Managing
Director of _____ (Proprietary Concern / Firm /
Company Ltd.) execute this Undertaking for myself and on behalf of
_____ (Proprietary Concern / Firm / Company Ltd.).

2. Whereas, TNMSC Ltd., (Tender Inviting Authority) has invited Tender for supply of
ARV Drugs under rate contract system for one year from the date of acceptance and in
pursuant to the conditions in the tender documents.

M/s. _____ (Proprietary Concern/ Firm / Company
Ltd.), having its Office at

_____ is exempted from payment of Earnest Money Deposit as
indicated in the Annexure-IX of tender document.

3. And whereas, in pursuant to the conditions in Clause Nos. 7.2&7.3(viii) of the tender,
the sum equivalent to the Earnest Money Deposit can be forfeited by the Tender Inviting
Authority in case of violation of any of the conditions and for non-performance of the obligation
under tender document.

4. In consideration of exempting M/s._____ (Proprietary Concern/ Firm / Company Ltd.) from payment of Earnest Money Deposit as indicated in the Annexure-IX of tender document, I undertake to pay the said sum without any demur on receipt of demand issued by the tender inviting authority.

M/s _____

for Self and Firm / Company Ltd.

Signature and Seal

Witness:-

(1)

(2)

PROFORMA FOR PERFORMANCE STATEMENT

**(FOR A PERIOD OF LAST 3 YEARS)
(2018-2019, 2019-2020 AND 2020-2021)**

Name of firm _____

Sl.	Name of the product	Year	No. of batches manufactured / imported & supplied.	Batch No.	Name and full address of the purchaser
	1	2	3	4	5
1.					
2.					
3.					

Note : Proof for the manufacturing (BMR) / importing of the drug quoted to be produced.

Signature and seal of the Tenderer _____

ANNUAL TURN OVER STATEMENT

The Annual Turnover of M/s. _____ for last 3 years
are given below and certified that the statement is true and correct.

Sl.No.	Financial Year	Turnover in Lakhs (Rs)
1.	2018 - 2019	-
2.	2019 - 2020	-
3.	2020 - 2021	-
Total		- Rs. _____ Lakhs.

Average turnover per annual - Rs. _____ Lakhs.

Date:

Seal:

Signature of Auditor/
Chartered Accountant
(Name in Capital)

TENDER FOR THE SUPPLY OF ARV DRUGS TO TANSACS UNDER RATE CONTRACT SYSTEM FOR ONE YEAR FROM THE DATE OF ACCEPTANCE

Sl. No.	Drug Code	Drug Name / Specification	Unit	Approx Tender Qty in Units	EMD Value (₹)	Drug Category
1	RC140	Tenofovir + Lamivudine + Efavirenz (TLE) Tablet Each Tablet Contains Tenofovir - 300mg + Lamivudine - 300mg + Efavirenz - 600mg	30 Tablets	10,400	50,000.00	A
2	RC141	Efavirenz Tablet - 200mg	30 Tablets	2,520	50,000.00	A
3	RC142	Efavirenz Tablet - 600mg	30 Tablets	1,200	50,000.00	A
4	RC143	Dolutegravir Tablet - 50mg	30 Tablets	60,000	50,000.00	A
5	RC160	Zidovudine + Lamivudine + Nevirapine Tablet Each Tablet Contains Zidovudine - 300mg + Lamivudine - 150mg + Nevirapine - 200mg	60 Tablets	14,600	50,000.00	A
6	RC161	Tenofovir + Lamivudine + Dolutegravir Tablet Each Tablet Contains Tenofovir - 300mg + Lamivudine - 300mg + Dolutegravir - 50mg	30 Tablets	4,12,800	50,000.00	A
7	RC162	Zidovudine + Lamivudine Tablet Each Tablet Contains Zidovudine - 300mg + Lamivudine - 150mg	60 Tablets	50,600	50,000.00	A
8	RC163	Tenofovir + Lamivudine Tablet Each Tablet Contains Tenofovir - 300mg + Lamivudine - 300mg	30 Tablets	68,160	50,000.00	A

TENDER FOR THE SUPPLY OF ARV DRUGS TO TANSACS UNDER RATE CONTRACT SYSTEM FOR ONE YEAR FROM THE DATE OF ACCEPTANCE

Sl. No.	Drug Code	Drug Name / Specification	Unit	Approx Tender Qty in Units	EMD Value (₹)	Drug Category
9	RC164	Nevirapine Tablet - 200mg	60 Tablets	1,200	50,000.00	A
10	RC165	Abacavir + Lamivudine Tablet Each Tablet Contains Abacavir - 600mg + Lamivudine - 300mg	30 Tablets	20,000	50,000.00	A
11	RC166	Zidovudine + Lamivudine + Nevirapine Tablet Each Tablet Contains Zidovudine - 60mg + Lamivudine - 30mg + Nevirapine - 50mg	60 Tablets	570	50,000.00	A
12	RC167	Abacavir + Lamivudine Tablet Each Tablet Contains Abacavir - 60mg + Lamivudine - 30mg	60 Tablets	33,320	50,000.00	A
13	RC168	Nevirapine Tablet - 50mg	60 Tablets	1,000	50,000.00	A
14	RC169	Zidovudine + Lamivudine Tablet Each Tablet Contains Zidovudine - 60mg + Lamivudine - 30mg	60 Tablets	11,400	50,000.00	A
15	RC170	Lopinavir + Ritonavir Tablet Each Tablet Contains Lopinavir - 200mg + Ritonavir - 50mg	120 Tablets	14,500	50,000.00	A

TENDER FOR THE SUPPLY OF ARV DRUGS TO TANSACS UNDER RATE CONTRACT SYSTEM FOR ONE YEAR FROM THE DATE OF ACCEPTANCE

Sl. No.	Drug Code	Drug Name / Specification	Unit	Approx Tender Qty in Units	EMD Value (₹)	Drug Category
16	RC171	Lopinavir + Ritonavir Tablet Each Tablet Contains Lopinavir - 100mg + Ritonavir - 25mg	120 Tablets	4,200	50,000.00	A
17	RC172	Lopinavir / Ritonavir Pellets	120 Tablets	800	50,000.00	A
18	RC173	Atazanavir (Adult) + Ritonavir (RTV) Tablets Each Tablet Contains Atazanavir – 300mg + Ritonavir – 100mg	30 Tablets	86,000	50,000.00	A
19	RC174	Ritonavir Tablet - 100mg	30 Tablets	8,000	50,000.00	A
20	RC175	Darunavir Tablet - 600mg	30 Tablets	8,000	50,000.00	A
21	RC176	Colour Coded KIT- 1 (Grey) Each Kit Contains Azithromycin Tablet - 1gm Single Dose + Cefixime Tablet - 400mg Single Dose	Kit	1,87,500	50,000.00	A
22	RC177	Colour Coded KIT- 2 (Green) Each Kit Contains Secnidazole Tablet - 2gm Single Dose + Fluconazole Tablet - 150mg Single Dose	Kit	1,87,500	50,000.00	A

TENDER FOR THE SUPPLY OF ARV DRUGS TO TANSACS UNDER RATE CONTRACT SYSTEM FOR ONE YEAR FROM THE DATE OF ACCEPTANCE

Sl. No.	Drug Code	Drug Name / Specification	Unit	Approx Tender Qty in Units	EMD Value (₹)	Drug Category
23	RC178	Colour Coded KIT- 3 (White) Each Kit Contains Azithromycin Tab - 1gm Single Dose + Benzathine Penicillin Inj - 2.4 MU (1) + Disposable Syringe - 10ml with 21 G Needle (1) + Sterile Water - 10ml (1)	Kit	18,750	50,000.00	A
24	RC179	Colour Coded KIT- 4 (Blue) Each Kit Contains Doxycycline Tab - 100mg (30) + Azithromycin Tab - 1gm (1)	Kit	33,125	50,000.00	A
25	RC180	Colour Coded KIT- 5 (Red) Each Kit Contains Acyclovir Tab - 400mg TID for 7 Days	Kit	36,250	50,000.00	A
26	RC181	Colour Coded KIT- 6 (Yellow) Each Kit Contains Cefixime Tab - 400mg Single Dose + Metronidazole Tab - 400mg BID for 14 Days + Doxycycline Cap - 100mg BID for 14 Days	Kit	91,250	50,000.00	A

TENDER FOR THE SUPPLY OF ARV DRUGS TO TANSACS UNDER RATE CONTRACT SYSTEM FOR ONE YEAR FROM THE DATE OF ACCEPTANCE

Sl. No.	Drug Code	Drug Name / Specification	Unit	Approx Tender Qty in Units	EMD Value (₹)	Drug Category
27	RC182	Colour Coded KIT- 7 (Black) Each Kit Contains Doxycycline Tab - 100mg BID for 21 Days + Azithromycin Tab - 1gm Dingle Dose	Kit	4,375	50,000.00	A
28	RC156	RPR (Rapid Plasma Reagin) Testing Kits (Specification of the Kit as mentioned Below)	Kit	20,00,000	50,000.00	A

SPECIFICATION OF RPR (RAPID PLASMA REAGIN) TESTING KITS (CODE NO.: RC156)

1. The kit should have approval of the statutory authority in its country of origin.
2. In case of imported kits it should be registered and licensed under the provisions of Drugs & Cosmetics Act and rules and / or Medical Devices Rules 2017 in India.
3. In case of indigenous manufacturers should be licensed under the provisions of Drugs & Cosmetics Acts and rules and or Medical Devices Rules 2017 issued by the competent authority defined under Drugs and Cosmetics Act, 1940.
4. The assay should allow for qualitative sand semi quantitative determination of Reagin antibodies in serum or plasma for sero – diagnosis of syphilis based on flocculation principle using non treponemal antigens.
5. The assay should be suitable to perform with either serum or plasma

TENDER FOR THE SUPPLY OF ARV DRUGS TO TANSACS UNDER RATE CONTRACT SYSTEM FOR ONE YEAR FROM THE DATE OF ACCEPTANCE

6. The assay should have sensitivity of more than or equal to 85% in primary syphilis and a specificity of more than or equal to 93%.
7. The assay should be calibrated to WHO reference serum and the same should be supported by statements in kit insert and certificate from manufacturer.
8. The test should be able to yield results within 20 minutes.
9. The pack size of RPR test kit should be less than or equal to 50 tests per kit.
10. The assay components should include positive and negative serum controls sufficient for conducting 20 % of the tests (10% negative and 10% positive controls)
11. The kit should have all essential accessories required for the test such as cards, droppers, applicator, etc., in adequate quantities for the number of tests to be performed.
12. The kit should have minimum shelf-life of 5/6th or 12 months (whichever is more) at the port of discharge of consignees.
13. The cumulative time temperature indicator technology used should be pre-qualified by WHO.
14. Literature detailing the components, methodologies, validity criteria, performance characteristics storage conditions, manufacturing and expiry dates should be provided with each kit.

Technical Specifications of cumulative time temperature indicator

1. Cumulative time/temperature indicator should indicate the exposure to high temperature above 8 degree C
2. The cumulative time- temperature indicator technology used should be prequalified by WHO

TENDER FOR THE SUPPLY OF ARV DRUGS TO TANSACS UNDER RATE CONTRACT SYSTEM FOR ONE YEAR FROM THE DATE OF ACCEPTANCE

3. The indicator should change color uniformly, irreversibly and the rate of reaction should be predictable by appropriate kinetic parameters.
4. The color change should have a well-defined start point and end point that can be correlated to the heat stability of the kit.
5. Should be mounted on card with clear instruction of interpretation.
6. The card should be self adhesive and placed on each kit box to monitor heat exposure during transit and storage of the kits till its expiry.

The manufacturer/authorized agent should ensure maintenance of cold chain during storage and transport of the kits at 2-8 degree.

I. SCHEDULE FOR PACKAGING OF ARV DRUGS
GENERAL SPECIFICATIONS

1. No corrugate package should weigh more than 15 kgs (ie., product + inner carton + corrugated box).
2. All Corrugated boxes should be of 'A' grade paper ie., Virgin.
3. All drugs should be packed only in first hand boxes only.

FLUTE:

4. The corrugated boxes should be of narrow flute.

JOINT:

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

STITCHING:

6. Every box should be stitched using pairs of metal pins with an interval of two inches between each pair. The boxes should be stitched and not joined using calico at the corners.

FLAP:

7. The flaps should uniformly meet but should not over lap each other. The flap when turned by 45 - 60° should not crack.

TAPE:

8. Every box should be sealed with Gum Tape/BoPP (Biaxially Oriented Polypropylene) tape running along the Top and lower opening.

CARRY STRAP:

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

LABEL:

10. Every corrugated box should carry a large outer label clearly indicating that the product is for **"Tamilnadu Govt. Supply - Not For Sale"**. The lower one third of the large label should indicate in bold, the value of the product as depicted in Annexure II of this document.
11. The product label on the carton should be large atleast 15cms x 10cms dimension. It should carry the correct technical name, strength of the product, date of manufacturing, date of expiry, quantity packed and net weight of the box.

OTHERS:

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

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

- (1) The box should not weigh more than 7-8 kgs. The grammage of outer box should be 150 gsm and inside partition / lining should be 120gsm.
- (2) The box should be of 5 ply with Bursting strength of 9 Kg/ Cm²

AGREEMENT

THIS AGREEMENT made the..... day of, 20 Between Tamilnadu Medical Services Corporation Limited, No.417, Pantheon Road, Egmore, Chennai – 600 008. (Name of purchaser) of (Country of Purchaser) (here in after "the Purchaser") of the one part and (Name of Supplier) of (City and Country of Supplier) (here in after called "the Supplier") of the other part :

WHEREAS the Purchaser is desirous that certain Goods and ancillary services viz;. Supply of ARV Drugs to TANSACS under rate contract system for one year from the date of acceptance in the tender Reference No. **011/M(P)/ARV-DRUG/TNMSC/2022, Dated.05.07.2022** (Brief Description of Goods and Services) and has accepted a bid by the Supplier for the supply of those goods and services for the sum of(Contract Price in Words and Figures) (hereinafter called "the Contract Price") for a period of one year from the date of execution of this agreement.

NOW THIS AGREEMENT WITNESSETH AS FOLLOWS:

1. In this Agreement words and expressions shall have the same meanings as are respectively assigned to them in the Conditions of Contract referred to, and they shall be deemed to form and be read and construed as part of this agreement.

2. The following documents shall be deemed to form and be read and construed as part of this Agreement, viz.:

- (a) The Letter of Acceptance issued by the purchaser.

- (b) The Notice Inviting Tender
- (c) The supplier's bid including enclosures, Annexure, etc.
- (d) The Terms and Conditions of the Contract
- (e) The Schedule of Requirement
- (f) The Technical Specification
- (g) Any other document listed in the supplier's bid and replies to queries, clarifications issued by the purchaser, such confirmations given by the bidder which are acceptable to the purchaser and the entire Addendum issued as forming part of the contract.

3. In consideration of the payments to be made by the Purchaser to the Supplier as hereinafter mentioned, the Supplier hereby covenants with the Purchaser to provide, the goods and services and to remedy defects therein in conformity in all respects with the provisions of the Contract.

4. The purchaser hereby covenants to pay the Supplier in consideration of the provision of the goods and services and the remedying of defects therein, the Contract Price or such other sum as may become payable under the provisions of the Contract at the times and in the manner prescribed by the Contract.

Brief particulars of the goods and services which shall be supplied / provided by the Supplier are as under.

Sl. No	Drug Code	Brief Description of Goods & Services	Unit	Tender Qty in Unit*	Unit Price	GST %	Total value inclusive of GST
Total contract value							

* Tender quantity indicated here is tentative and may vary subjected to various terms and conditions of the tender.

DELIVERY SCHEDULE:

The supplier shall supply at least **50%** of the ordered quantity within **45 days** from the date of purchase order and the balance quantity within **60 days** from the date of purchase order at the destinations mentioned in the purchase order.

IN WITNESS where of the parties here to have caused this Agreement to be executed in accordance with their respective laws the day and year first above written.

Signed, Sealed and Delivered by the

said..... (For the Purchaser)

in the presence of

Signature

Name

Address

Signed, Sealed and Delivered by the

Said (For the Supplier)

in the presence of

Signature

Name

Address

Performance Security Bank Guarantee

(unconditional)

To : **Tamilnadu Medical Services Corporation Limited** (Name of **Purchaser**)
No.417, Pantheon Road, Egmore, Chennai – 600 008.

WHEREAS (Name of the Supplier) herein called
“the Supplier” has undertaken, in pursuance of Tender No.011/M(P)/ARV-DRUG/TNMSC/2022,
Dated.05.07.2022 to supply of **ARV Drugs to TANSACS under rate contract system for one
year from the date of acceptance.** (Description of Goods and Services) hereinafter called “the
Contract”.

AND WHEREAS it has been stipulated by you in the said Contract that the Supplier
shall furnish you with a Bank Guarantee by a recognized 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, upto 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 Contract and without cavil or

argument, any sum or sums within the limit of (Amount of the
Guarantee in Words and Figures) 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 20.....

Signature and Seal of Guarantors

.....
.....
.....

Date 20

Address
.....
.....

DETAILS OF MANUFACTURING /IMPORTING UNIT

(In Case of Loan Licensee, the details of manufacturing facility to be furnished)

Name of the Tenderer and
office Address :

Factory Address * :

PAN Number :

GST Number :

Phone Nos. :

Fax :

Mobile Number
(SMS shall be alerts to this Mobile Number) :

E-Mail
(e-purchase order(s) shall be to this Mail ID) :

Date of Inception :

Licence No. & Date :

Issued by :

Valid up to :

Details of installed Production Capacity :

Details of Installed Production Capacity for 30 days
(In Terms of Unit Packs)

Tablets :

Capsules

General :

Beta-Lactum :

Injections

Ampoules :

Vials :

I.V.Fluids :

Sterile Powder :

Liquids

Suspension :

Syrups :

Drops :

Ointment :

Powders :

Antiseptics /
Disinfectants :

Name & designation of the authorized signatory :

Specimen signature of the authorized Signatory :

*** The details of manufacturing unit should be for the premises where drugs quoted are actually manufactured**

THE DETAILS OF FACTORY PREMISES

Person In-charge of Factory

Name :

Phone No. :

Mobile No. :

Nearest Land mark of Factory :

Layout

Km from Airport :

Name of the Airport and City :

Km from Railway Station :

Name of the Railway station
and city :

Km from Bus Stand :

Name of the Bus Stand
and City :

Name & designation of the authorized signatory

ANNEXURE – XIV
Ref. clause 4.1 (r)

List of Drugs quoted

1. Name of the firm and address
as given in Drug licence :
2. Drug Licence No. in form 25 & 28 /
25-A & 28-A or import Licence No. :
3. Date of issue & validity :
4. Revised schedule M compliance
Certificate obtained on :
5. Non-conviction Certificate
Obtained on :
6. Market standing Certificate
obtained on :

In the event of the bidder becoming L1 for more than one drug, if the total annual quantity for such drugs is more than the capacity earmarked to TNMSC, TNMSC reserve the rights to decide any appropriate drug(s) within his production capacity.

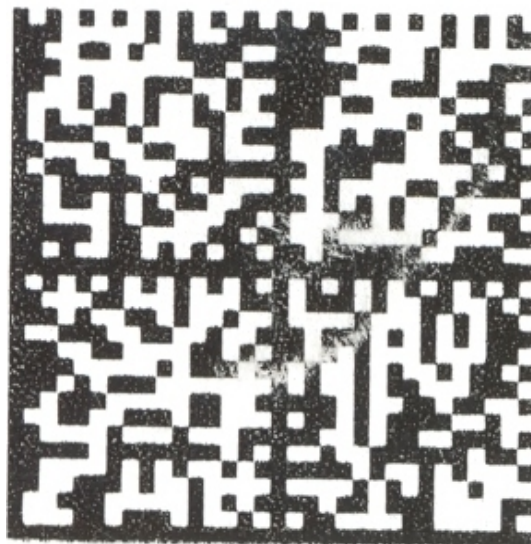
7. Details of Endorsement for all products quoted :

Sl. No	Drug Code	Drug Name	IP / BP / USP	Date of Endorsement obtained from the State Drugs Controller	Whether Endorsement is in Generic or Trade Name	Expiry Period for the drugs quoted should be specified in Months (for Eg. Griseofulv in Tab. – 48 Months)	Yearly Mfg. Capacity Earmarked to TNMSC for each drug quoted (in Units)	Mfg / Importing unit location (State from which supplies will be made)	Value of EMD
1	2	3	4	5	6	7	8		9
1.									
EMD Total									

Authorised signatory :
Date :

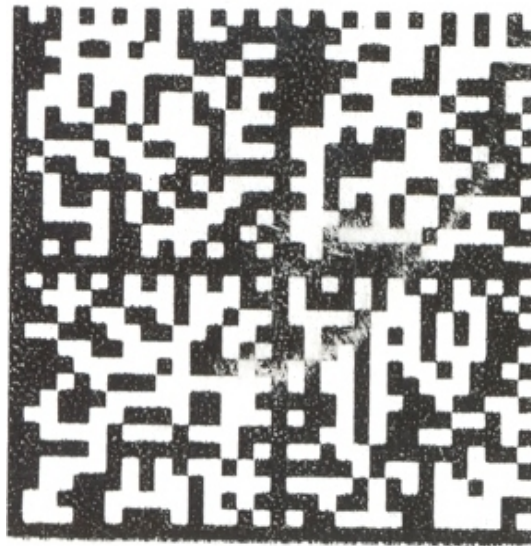
2D Bar Coding on Tertiary Packing

BOX NO :
PO NUMBER :
SUPPLIER CODE :
SUPPLIER NAME :
DRUG CODE :
DRUG NAME :
BATCH NO :
MFG DATE :
EXPIRY DATE :
BATCH QUANTITY :
INVOICE NO :
D C NO :



2D Bar Coding on Secondary Packing

DRUG CODE :
DRUG NAME :
BATCH NO :
SUPPLIER CODE :
SUPPLIER NAME :
MFG DATE :
EXPIRY DATE :



MANDATE FORM

01	Company Name	
02	Postal Address of the company with Telephone No., Fax No. and Mail I.D.	
03	Name of the Managing Director / Director / Manager Mobile No. / Phone No. E-mail I.D.	
04	Name and Designation of the authorized company official Mobile No. E-mail ID	

Date: Company Seal

Signature

Place: (Name of the person signing & designation)

01	Name of the Bank . Branch Name& address. Branch Code No. Branch Manager Mobile No. Branch Telephone no. Branch E-mail ID	
02	9 digit MICR code number of the bank and branch appearing on the MICR cheque issued by the bank.	
03	IFSC code of the Branch	
04	Type of Account (Current / Savings).	
05	Account Number (as appear in cheque book)	

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

I /We hereby declare that the particulars given above are correct and complete. If the transaction is delayed or not effected at all for reasons of incomplete or incorrect information, I would not hold M/s. Tamilnadu Medical Services Corporation Limited (TNMSC) responsible. I

have read the conditions of the tender/agreement entered and agree to discharge the responsibility expected of me / from the company as a tenderer /successful tenderer.

Date:	Company Seal	Signature
Place:	(Name of the person signing & designation)	
<hr/>		

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

Bank Seal with address.	Signature of the authorized official of the bank.
<hr/>	

FORMAT FOR EARNEST MONEY DEPOSIT BANK GUARANTEE

Whereas (*hereinafter called "the Tenderer"*) has submitted its tender in the **Ref.No.011/M(P)/ARV-DRUG/TNMSC/2022, Dated.05.07.2022** dated (*date of submission of tender*) for the supply of (*name and/or description of the goods*) (*hereinafter called "the Tender"*).

KNOW ALL PEOPLE by these presents that WE (*name of bank*) of (*name of country*), having our registered office at (*address of bank*) (*hereinafter called "the Bank"*), are bound unto (*name of purchaser*) (*hereinafter called "the Purchaser"*) in the sum of _____ for which payment well and truly to be made to the said Purchaser, the Bank binds itself, its successors, and assigns by these presents.

Sealed with the Common Seal of the said Bank this _____ day of _____ 20____.

THE CONDITIONS of this obligation are :

1. If the Tenderer
 - a) withdraws its Tender during the period of tender validity specified by the Tenderer on the tender Form; or
 - b) does not accept the correction of errors in accordance with the Tender; or

- c) rejected on inspection for the compliance of Good Manufacturing Practice as per revised schedule-M of Drugs & Cosmetics Act.
2. If the Tenderer, having been notified of the acceptance of its tender by the Purchaser during the period of tender validity :
- a) fails or refuses to execute the Agreement if required; or
 - b) fails or refuses to furnish the security deposit, in accordance with the Instruction to Tenderer;

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

This guarantee will remain in force up to and including Sixty (60) days after the period of the tender validity, i.e. 180 days from the date of opening of the tender, and any demand in respect thereof should reach the Bank not later than the above date.

.....
(Signature of the Bank)

1 Name of Tenderer

CHECK LIST

ANNEXURE - XVIII
Ref. Clause. 4.1.(s)

COVER - A.

Sl.	Description	Page No.	Yes	No
1.	Checklist – Annexure-XVII.	1	✓	
2.	EMD in the form of Demand Draft/Bank Guarantee shall be kept in an envelope as per (Annexure-V) (Domestic Enterprise under MSME- Exempted).			
3.	Annual Turnover Statement of last 3 years (Annexure-VIII).			
4.	Under taking of Domestic Enterprise under MSME (Annexure-VI).			
5.	Declaration for Non Blacklisting of the Firm / Products (Annexure-IV).			
6.	List of drugs quoted with production capacity and without rates (Annexure-XIV).			
7.	Duly attested photocopy of Manufacturing License for the product duly approved by the Licensing authority for each and every product(s) quoted.			
8.	Duly attested photocopy of Import License, along with whole sale Drug license.			
9.	Details of Technical personnel employed in the manufacture and testing.			
10.	Market Standing Certificate issued by the Licensing Authority.			
11.	Non-conviction Certificate / Conviction Status Certificate issued by the Licensing Authority.			
12.	Good Manufacturing Practices Certificate.			

Sl.	Description	Page No.	Yes	No
13.	Documentary evidence for the constitution of the company /concern.			
14.	List of Board of Directors/ Partners / Proprietor.			
15.	The instruments such as power of attorney, resolution of board etc. for authorized signatory			
16.	Authorization letter nominating the employee of the tenderer to transact the business with the Tender inviting Authority.			
17.	Copies of Balance Sheet and Profit & Loss Account details for Last 3 years.			
18.	GST returns from 01.01.2021 to 31.03.2021 (as applicable) along with GST registration copy of the tenderer.			
19.	Undertaking for embossment of logo (Annexure-I). (Affixing the logo for Secondary / Primary packing for the Manufacturing drugs)			
20.	Undertaking for embossment of logo (Annexure-II). (Affixing the logo for Secondary / Primary packing for the imported drugs along with Brand / trade Names)			
21.	Declaration Form for cGMP/WHO GMP along with enclosure (Annexure-III).			
22.	Details of Manufacturing/Importing Unit in Annexure-XIII .			
23.	WHO, UNICEF, ISO certificates if any.			
24.	Mandate Form (Annexure-XVI).			
25.	Proforma for Performance Statement (Annexure-VII).			

Sl.	Description	Page No.	Yes	No
26.	True copy of record of manufacture to establish 3 years market standing.			
27.	The Tender document signed by the tenderer/authorized signatory in all pages with official seal.			
28.	Signed and Sealed copy of the Corrigendum issued for the tender conditions (Issued If Any)			

TAMILNADU MEDICAL SERVICES CORPORATION LTD., CHENNAI - 600 008

TENDER FOR THE SUPPLY OF ARV DRUGS TO TANSACS UNDER RATE CONTRACT SYSTEM FOR ONE YEAR FROM THE DATE OF ACCEPTANCE

Break up of Landed price per unit

**Annexure-XIX
Ref. Clause 5**

P Sl. No	Drug Code	Name of the Drug and Strength	Unit	HSN Code No	Basic Price Inclusive of Incidental Charge (Per Unit) (Rs.P)	Packing & Forwarding Charges (Per Unit) (Rs.P)	Freight and Insurance Charges (Per Unit) (Rs.P)	Customs Duty (Per Unit) (if Applicable) (Rs.P)	Total Landed Price (in Figure) (Per Unit) (5 + 6 + 7 + 8) (Rs.P)	Rate per Unit in Words (Exclusive of Tax)	SGST (% Only)	CGST (% Only)	IGST (or) UTGST (% Only)	Drug Category
(1)	(2)	(3)		(4)	(5)	(6)	(7)	(8)	(9)	(10)	(11)	(12)	(13)	(14)
1	RC140	Tenofovir + Lamivudine + Efavirenz (TLE) Tablet Each Tablet Contains Tenofovir - 300mg + Lamivudine - 300mg + Efavirenz - 600mg	30 Tablets											A
2	RC141	Efavirenz Tablet - 200mg	30 Tablets											A
3	RC142	Efavirenz Tablet - 600mg	30 Tablets											A
4	RC143	Dolutegravir Tablet - 50mg	30 Tablets											A
5	RC160	Zidovudine + Lamivudine + Nevirapine Tablet Each Tablet Contains Zidovudine - 300mg + Lamivudine - 150mg + Nevirapine - 200mg	60 Tablets											A

Note: The firms shall indicate the break up prices at Column 5 to 8 and 11 to 13 separately and wording like "Included" shall not be substituted for the same.

Signature of the Authorized Signatory : _____
Name of the Firm / Company : _____
Manufacturing Premises Located at (State) : _____

TAMILNADU MEDICAL SERVICES CORPORATION LTD., CHENNAI - 600 008

TENDER FOR THE SUPPLY OF ARV DRUGS TO TANSACS UNDER RATE CONTRACT SYSTEM FOR ONE YEAR FROM THE DATE OF ACCEPTANCE

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(1)	(2)	(3)		(4)	(5)	(6)	(7)	(8)	(9)	(10)	(11)	(12)	(13)	(14)
6	RC161	Tenofovir + Lamivudine + Dolutegravir Tablet Each Tablet Contains Tenofovir - 300mg + Lamivudine - 300mg + Dolutegravir - 50mg	30 Tablets											A
7	RC162	Zidovudine + Lamivudine Tablet Each Tablet Contains Zidovudine - 300mg + Lamivudine - 150mg	60 Tablets											A
8	RC163	Tenofovir + Lamivudine Tablet Each Tablet Contains Tenofovir - 300mg + Lamivudine - 300mg	30 Tablets											A
9	RC164	Nevirapine Tablet - 200mg	60 Tablets											A
10	RC165	Abacavir + Lamivudine Tablet Each Tablet Contains Abacavir - 600mg + Lamivudine - 300mg	30 Tablets											A

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(1)	(2)	(3)		(4)	(5)	(6)	(7)	(8)	(9)	(10)	(11)	(12)	(13)	(14)
11	RC166	Zidovudine + Lamivudine + Nevirapine Tablet Each Tablet Contains Zidovudine - 60mg +Lamivudine - 30mg + Nevirapine - 50mg	60 Tablets											A
12	RC167	Abacavir + Lamivudine Tablet Each Tablet Contains Abacavir - 60mg + Lamivudine - 30mg	60 Tablets											A
13	RC168	Nevirapine Tablet - 50mg	60 Tablets											A
14	RC169	Zidovudine + Lamivudine Tablet Each Tablet Contains Zidovudine - 60mg + Lamivudine - 30mg	60 Tablets											A
15	RC170	Lopinavir + Ritonavir Tablet Each Tablet Contains Lopinavir - 200mg + Ritonavir - 50mg	120 Tablets											A

Note: The firms shall indicate the break up prices at Column 5 to 8 and 11 to 13 separately and wording like "Included" shall not be substituted for the same.

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TENDER FOR THE SUPPLY OF ARV DRUGS TO TANSACS UNDER RATE CONTRACT SYSTEM FOR ONE YEAR FROM THE DATE OF ACCEPTANCE

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Ref. Clause 5

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(1)	(2)	(3)		(4)	(5)	(6)	(7)	(8)	(9)	(10)	(11)	(12)	(13)	(14)
16	RC171	Lopinavir + Ritonavir Tablet Each Tablet Contains Lopinavir - 100mg + Ritonavir - 25mg	120 Tablets											A
17	RC172	Lopinavir / Ritonavir Pellets	120 Tablets											A
18	RC173	Atzanavir (Adult) + Ritonavir (RTV) Tablets Each Tablet Contains Atzanavir – 300mg + Ritonavir – 100mg	30 Tablets											A
19	RC174	Ritonavir Tablet - 100mg	30 Tablets											A
20	RC175	Darunavir Tablet - 600mg	30 Tablets											A

Note: The firms shall indicate the break up prices at Column 5 to 8 and 11 to 13 separately and wording like “Included” shall not be substituted for the same.

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(1)	(2)	(3)		(4)	(5)	(6)	(7)	(8)	(9)	(10)	(11)	(12)	(13)	(14)
21	RC176	Colour Coded KIT- 1 (Grey) Each Kit Contains Azithromycin Tablet - 1gm Single Dose + Cefixime Tablet - 400mg Single Dose	Kit											A
22	RC177	Colour Coded KIT- 2 (Green) Each Kit Contains Secnidazole Tablet - 2gm Single Dose + Fluconazole Tablet - 150mg Single Dose	Kit											A
23	RC178	Colour Coded KIT- 3 (White) Each Kit Contains Azithromycin Tab - 1gm Single Dose + Benzathine Penicillin Inj - 2.4 MU (1) + Disposable Syringe - 10ml with 21 G Needle (1) + Sterile Water - 10ml (1)	Kit											A

Note: The firms shall indicate the break up prices at Column 5 to 8 and 11 to 13 separately and wording like "Included" shall not be substituted for the same.

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Name of the Firm / Company : _____
Manufacturing Premises Located at (State) : _____

TAMILNADU MEDICAL SERVICES CORPORATION LTD., CHENNAI - 600 008

TENDER FOR THE SUPPLY OF ARV DRUGS TO TANSACS UNDER RATE CONTRACT SYSTEM FOR ONE YEAR FROM THE DATE OF ACCEPTANCE

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Ref. Clause 5**

P Sl. No	Drug Code	Name of the Drug and Strength	Unit	HSN Code No	Basic Price Inclusive of Incidental Charge (Per Unit) (Rs.P)	Packing & Forwarding Charges (Per Unit) (Rs.P)	Freight and Insurance Charges (Per Unit) (Rs.P)	Customs Duty (Per Unit) (if Applicable) (Rs.P)	Total Landed Price (in Figure) (Per Unit) (5 + 6 + 7 + 8) (Rs.P)	Rate per Unit in Words (Exclusive of Tax)	SGST (% Only)	CGST (% Only)	IGST (or) UTGST (% Only)	Drug Category
(1)	(2)	(3)		(4)	(5)	(6)	(7)	(8)	(9)	(10)	(11)	(12)	(13)	(14)
24	RC179	Colour Coded KIT- 4 (Blue) Each Kit Contains Doxycycline Tab - 100mg (30) + Azithromycin Tab - 1gm (1)	Kit											A
25	RC180	Colour Coded KIT- 5 (Red) Each Kit Contains Acyclovir Tab - 400mg TID for 7 Days	Kit											A
26	RC181	Colour Coded KIT- 6 (Yellow) Each Kit Contains Cefixime Tab - 400mg Single Dose + Metronidazole Tab - 400mg BID for 14 Days + Doxycycline Cap - 100mg BID for 14 Days	Kit											A

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(1)	(2)	(3)		(4)	(5)	(6)	(7)	(8)	(9)	(10)	(11)	(12)	(13)	(14)
27	RC182	Colour Coded KIT- 7 (Black) Each Kit Contains Doxycycline Tab - 100mg BID for 21 Days + Azithromycin Tab - 1gm Dingle Dose	Kit											A
28	RC156	RPR (Rapid Plasma Reagin) Testing Kits (Specification of the Kit as mentioned Annexure-IX)	Kit											A

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