MINISTRY OF HEALTH & FAMILY WELFARE

Department of AIDS Control National AIDS Control Organization



National HIV/AIDS Control Programme

NATIONAL COMPETITIVE BIDDING

BID DOCUMENT

For

PROCUREMENT OF SINGLE BLOOD BAGS (350ML), DOUBLE BLOOD BAGS 350ML & 450 ML, TRIPLE BLOOD BAGS 350ML & 450ML, QUADRUPLE BLOOD BAGS 350ML & 450ML

IFB NO.:- RITES/MSM/NACP/11/2013



(Procurement Agent)

Materials System Management Division
RITES Ltd., RITES Office Complex, Annex Building, 4th Floor,
Plot No.144, Sector 44
Gurgaon - 122003, Haryana, India

Fax: 91(124)2571659/2571660 Tel: 91(124) 2728-408/405/403

MINISTRY OF HEALTH & FAMILY WELFARE **Department of AIDS Control National AIDS Control Organization**

Through

RITES Ltd., RITES Office Complex, Annex Building, 4th Floor, Plot No.144, Sector 44 Gurgaon - 122003, Haryana, India Fax: 91(124) 2571659/2571660

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NAME OF THE PROJECT: - National HIV/AIDS Control Programme

BID REFERENCE: -RITES/MSM/NACP/11/2013

DATE OF COMMENCEMENT

OF SALE OF BID DOCUMENT: 3rd September 2013

DATE AND TIME OF PRE-BID

12th September 2013 at 1400 Hrs. (IST) **CONFERENCE:**

LAST DATE AND TIME FOR

1st October 2013 up to 1400 Hrs. (IST) **RECEIPT OF BID:**

TIME AND DATE OF OPENING

1st October 2013 at 1415 Hrs. (IST) **OF BIDS:**

PLACE OF OPENING OF BIDS: RITES Ltd..

> MSM Division, RITES Office Complex, Annex Building, 4th Floor, Plot No.144, Sector 44, Gurgaon-122003 (Haryana), India

Fax: 91(124)2571659/2571660 Tel: 91(124) 2728-408/405/403

ADDRESS FOR COMMUNICATION: RITES Ltd.,

> MSM Division, RITES Office Complex, Annex Building, 4th Floor, Plot No.144, Sector 44, Gurgaon-122003 (Harvana), India

Fax: 91(124)2571659/2571660 Tel: 91(124) 2728-408/405/403

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Invitation For Bids

Invitation for Bids (IFB) 5

Invitation for Bids (IFB)

Country : India

Name of Project : National HIV/AIDS Control Programme

Name of Goods : SINGLE BLOOD BAGS (350ML), DOUBLE BLOOD BAGS

350ML & 450 ML, TRIPLE BLOOD BAGS 350ML & 450ML,

QUADRUPLE BLOOD BAGS 350ML & 450ML

IFB No : RITES/MSM/NACP/11/2013

1. National Aids Control Organization, Ministry of Health & Family Welfare, Govt. of India intends to utilise part of its domestic budget for eligible payments under the contracts for Procurement of SINGLE BLOOD BAGS (350ML), DOUBLE BLOOD BAGS 350ML & 450 ML, TRIPLE BLOOD BAGS 350ML & 450ML, QUADRUPLE BLOOD BAGS 350ML & 450ML against Schedule I to XVII for which this invitation for bid is issued under National HIV/AIDS Control Programme.

- 2. RITES Ltd. (A Govt. of India Enterprise), acting as procurement agent on behalf of Ministry of Health & Family Welfare, Govt. of India now invites sealed bids from eligible bidders for the Procurement of SINGLE BLOOD BAGS (350ML), DOUBLE BLOOD BAGS 350ML & 450 ML, TRIPLE BLOOD BAGS 350ML & 450ML, QUADRUPLE BLOOD BAGS 350ML & 450ML for the quantity as per Schedule of Requirement to the consignees located at various states all over India.
- 3. Bidding will be conducted through the National Competitive Bidding procedures as per the requirements, under GFR 2005 of Ministry of Finance, GOI, as applicable.
- 4. Interested eligible Bidders may obtain further information from RITES Ltd. and inspect the bidding documents at the address given below from 1000 to 1600 hrs. (IST) on all working days.
- 5. A complete set of bidding documents in English may be purchased by interested bidders on the submission of a written application to the address below and upon payment of a non-refundable fee of **Rs. 5000.** The method of payment will be by Demand Draft/Pay Order in favour of RITES Ltd., Payable at Gurgaon, India. The document may be purchased from 3rd September 2013 to 1st October 2013 from the address mentioned below in S. No. 6. The document will be sent by courier on payment of an extra amount of **Rs 900** for domestic bidder if requested by mail.

Bidders can also download the bid document from RITES website "<u>www.rites.com</u>" or www.nacoonline.org. For downloaded bid document, no fee is required. The bidders, who have downloaded the bid documents, shall be solely responsible for checking these websites for any addendum/amendment issued subsequently to the bid document and take into consideration the same while preparing and submitting the bids.

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6. The bidders or their official representatives are invited to attend a pre bid meeting which will take place on 12th September 2013 at 1400 hrs (IST) at the address mentioned below in S. No. 6. Please note that non-attendance at the pre-bid conference will not be the cause of disqualification of the bidders.

7. Bids must be delivered to the address below before **1400 hrs (IST)** on **1**st **October 2013**. All bids must be accompanied by a bid security as specified in the "Section VI – Schedule of Requirements" of the bidding document. Late bids will be rejected. Bids will be opened in the presence of the bidders' representatives who choose to attend at the address below at **1415 hrs (IST)** on **1**st **October 2013**.

Group General Manager/MSM RITES Ltd., MSM Division, RITES Office Complex, Annex Building, 4th Floor, Plot No.144, Sector 44, Gurgaon-122003 (Haryana), India Fax: 91(124)2571659/2571660

Tel: 91(124) 2728-408/405/403

Email: rites_naco@rediffmail.com, rites_naco@rites.com

8. Bid documents are non transferable.

SECTION I. INSTRUCTIONS TO BIDDERS

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A. INTRODUCTION

1. Scope of Bid

- 1.1 RITES Ltd., RITES Office Complex, Annex Building, 4th Floor, Plot No.144, Sector 44, Gurgaon-122003 (Haryana), India for and on behalf of Ministry of Health & Family Welfare (Govt. of India) invites bids for SINGLE BLOOD BAGS (350ML), DOUBLE BLOOD BAGS 350ML & 450 ML, TRIPLE BLOOD BAGS 350ML & 450ML. Detailed description of goods and specification are given in schedule of requirement and technical specification respectively. Identification number of contract is RITES/MSM/NACP/11/2013.
- 1.2 Throughout these bidding documents, the terms "writing" means any handwritten, typewritten, or printed communication, including telex, cable, and facsimile transmission, and "day" means calendar day. Singular also means plural.
- 2. Source of Funds
- 2.1 The Government of India.

3. Fraud and Corruption

- 3.1 It is the Government of India policy that Bidders/Suppliers/Contractors under the contracts, observe the highest standard of ethics during the procurement and execution of such Contracts. In pursuance of this policy, the Purchaser:
 - (a) defines, for the purposes of this provision, the terms set forth below as follows:
 - (i) "corrupt practice" means the offering, giving, receiving, or soliciting of anything of value to influence the action of a public official in the procurement process or in Contract execution; and
 - (ii) "fraudulent practice" means 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" means 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" means 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;
 - (b) will declare a firm 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 engaged in corrupt or fraudulent or collusive or coercive practices in competing for, or in executing, the contract.

- 3.2 Furthermore, bidders shall be aware of the provision stated in Sub-Clauses 6.4 and 23.1 (c) of the General Conditions of Contract.
- 3.3 In pursuance of the policy defined in ITB Sub-Clause 3.1, the purchaser will cancel the Contract for Goods or works if it at any time determines that corrupt or fraudulent or collusive or coercive practices were engaged during the procurement or the execution of the Contract.

4. Eligibility

- 4.1 Except as provided in ITB Sub-Clauses 4.2 this bidding process is open to all Indian bidders. Non manufacturer bidders will have to submit Manufacturer's Authorization Form 7 in Section V.
- 4.2 A firm declared ineligible by the Purchaser in accordance with ITB Sub-Clause 3.1(b) shall be ineligible to bid for the contract during the period of time determined by the Purchaser.
- 5. Documents
 Establishing
 conformity of
 Goods and
 Services to
 Bidding
 Documents
- 5.1 The documentary evidence of conformity of the goods and services to the Bidding Documents may be in the form of literature, drawings, and data and shall consist of:
 - (a) a detailed description of the essential technical and performance characteristics of the Goods;
 - (b) an item-by-item commentary on the Purchaser's Technical Specifications demonstrating substantial responsiveness of the Goods and Services to those specifications, or a statement of deviations and exceptions to the provisions of the Technical Specifications;
 - (c) The Goods offered should meet the specified pharmaceuticals standards as stated in the Technical Specifications. If the Goods offered are not included in one of the specified pharmacopoeias (e.g., the case of new drug), the Bidder will provide testing protocols and alternative standards.
- 5.2 The Goods to be supplied under the Contract shall be registered with the relevant authority in the Purchaser's country. A Bidder who has already registered its Goods by the time of bidding should submit a copy of the Registration Certificate with its bid. Otherwise, the successful Bidder, by the time of Contract signing, shall submit to the Purchaser:
 - (1) Copy of Registration Certificate establishing registration of Goods to be supplied under the Contract, with the National Regulatory Authority of India viz. Central Drugs Standard Control

Organization (CDSCO).

(2) Copy of documentation indicating that the goods proposed to be supplied under this contract are registered and licensed for use in India by the DCG (I) (Drugs Controller General of India) for imported pharmaceuticals and by the competent authority defined under the Drugs and Cosmetics Act 1940, as amended, after appropriate evaluation by centers approved by the DCG (I) (Drugs Controller General of India) for pharmaceuticals produced by indigenous manufacturers.

Note: Bidders are requested to inquire in advance about the registration requirements and procedures in order to avoid any delays due to involvement of various government agencies. Purchaser shall not be responsible for any delay on this account.

- 5.2.1 The Purchaser shall at all times cooperate with the successful Bidder to facilitate the registration process within the Purchaser's country. The agency and contact person able to provide additional information about the requirements for registration can be obtained from the Website: www.cdsco.nic.in.
- 5.2.2 If the Goods of the successful Bidder have not been registered in the Purchaser's country at the time of Contract signing, then the Contract shall become effective upon such date as the Certificate of Registration is obtained.
- 5.3 For purposes of the commentary to be furnished pursuant to ITB Clause 5.1 (b) above, the Bidder shall note that standards as well as references to brand names designated by the Purchaser in its Technical Specifications are intended to be descriptive only and not restrictive. The Bidder may substitute alternative standards, brand names, and/or catalogue numbers in its bid, provided that it demonstrates to the Purchaser's satisfaction that the substitutions ensure substantial equivalence to those designated in the Technical Specifications and meet the Pharmacopoeial standards.

6. Qualifications of the Bidder

Qualification requirements for Bidders are listed below:

The qualification criteria and the supporting document/information to be submitted along with the bid are detailed below:

(A) Manufacturer Bidders

(i) that, in the case of a Bidder offering to supply Goods under the Contract which the Bidder manufactures or otherwise produces (using ingredients supplied by primary manufacturers) that the Bidder:

- (a) is incorporated in the country of manufacture of the Goods;
- (b) has been licensed by the regulatory authority in the country of manufacture to supply the Goods covered by the IFB;
- (c) has manufactured and marketed the specific good covered by the bidding document for at least **one** (1) **year in the last five** (5) **years,** and for **similar goods** (**viz. Blood Bags**) for at least **three** (3) **years in the last five** (5) **years.** In support of this, data on past performance should be submitted as per Form 6 in Section V;
- (d) has shown evidence of compliance [for the factory where the specific goods are manufactured and are being offered for supply] with ISO 13485:2003 (or FDA 21 CFR 820) by way of accreditation by an independent recognized certification body.
- (e) provides the evidence that it has the financial, technical and production capability necessary to perform the contract as under:
 - 1. that it has successfully completed at least one (1) contract for similar goods (any type of Blood Bags) within the period of last five years (preceding two months before the date of bid opening) for supply of goods. Minimum value of completed contract for each schedule should be as per Appendix 'A'. Bidder shall submit list of major supply contracts conducted within the last five years as per form 6 (Proforma for Performance Statement) in Section V.
 - 2. that it has achieved an actual annual production of similar goods (any type of Blood Bags) of equal quantity as specified in relevant schedule in "Section III Schedule of Requirements" during any one of the last three (3) financial years; certified by chartered accountant. If the bidder quotes for more than one schedule the above criteria will be cumulative.
 - 3. that it has generated an annual turnover of the value of at least equal to as specified in Appendix 'B', during any one of the last three financial years, to qualify for a particular schedule. If the bidder quotes for more than one Schedule, the above criteria shall be cumulative. The turnover is to be supported by audited financial statements of accounts (including balance sheet, profit and loss account, auditor's reports and IT returns) for the past three financial years duly certified by the auditor of

the Company.

When offering their bid for more than one schedule, the bidder must provide evidence that it meets or exceeds the sum of all the individual requirements for the schedules being applied for in regard to

- (I) Actual annual production (sub-clause (f) (2) above) and
- (II) Actual annual turnover (sub-clause (f) (3) above).

Hence, if the bidder quotes for more than one schedule, the above criteria shall be cumulative. In case a bidder fails to fully meet any of these criteria, it will be qualified only for those schedules for which the bidder meets the above requirements and the combination of schedules to be awarded to such bidder will be decided based on the lowest cost of the combination to the Purchaser. The decision of the buyer in this regard shall be final and binding on the bidder.

- Note: 1. However, the cumulative criteria will not be applicable for one successfully completed contract within the last five years (subclause A (i)(f) (1) above) that mean if a firm has completed one contract of value more than Rs. x Million then it will qualify for all schedules whose value less than Rs. x Million.
- (ii) The Bidder shall also submit the following documents with respect to above qualification criterias:
 - 1. A copy of its manufacturing license and a statement of installed manufacturing capacity.
 - 2. details of on-site quality control laboratory facilities and services and range of tests conducted;
 - 3. copies of its audited financial statements for the past three fiscal years.
 - 4. A copy of the achieved annual production rate certified by Chartered Accountant.
 - 5. list of major supply contracts executed/ secured (Completed and ongoing) within the last five years as per Form 6 in Section V.

(B) Non Manufacturer Bidder

a) In the case of a Bidder offering to supply Goods under the Contract that the Bidder does not manufacture or otherwise produce, (all supporting documents that the Bidder should be duly authorized by the manufacturer of the Goods

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who meets the criteria under (A) above /information as asked above for manufacturer shall be submitted with the bid), as per authorization Form 7 in Section V;

- b) The bidder has successfully completed at least one similar contract within the period of last five (5) years (preceding two months before the date of opening of bids) for supply of goods. Minimum value of the completed contract should be at least 50% of the value indicated against each schedule as indicated in 'Appendix A' and that includes comparable products e.g. any type of Blood Bags or any pharmaceutical products. Bidder shall submit list of major supply contracts conducted within the last five years as per form 6 (Proforma for Performance Statement) in Section V
- that it has generated an annual turnover of at least 50% of the value as given in 'Appendix B', in any one of the last three (3) financial years, to qualify for a particular schedule. If the bidder quotes for more than one Schedule, the above criteria shall be cumulative. The turnover is to be supported by audited financial statements of accounts (including balance sheet, profit and loss account, auditor's reports, and IT returns) for the past three fiscal years duly certified by the auditor of the Company.
- d) NOTE: In case any bidder is lowest evaluated & responsive in more than one schedule but fails to meet the cumulative requirement of turn over for those schedules, consideration of bid for specific schedule wherein he meets the requirement of the schedule, will be at the sole decision of the buyer.
- e) The bidder will also submit the list of major supply contracts completed within the last five years as per Form 6 in Section V.
- f) Copies of its audited financial statements for the past three fiscal years.
- NOTE- 1. An agent submitting a bid in its own name will be treated as a non-manufacturer bidder.

(C) For both (A) and (B)

The Bidder shall also submit the following additional information:

1. The bidder and the manufacturer whose product is offered by

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the bidder shall disclose instance of previous past performance of his and the manufacturer whose product is procured by the bidder, that may have resulted into adverse actions taken against the bidder during the **last two years**. Such adverse actions taken against the bidder or manufacturer may be treated as unsatisfactory performance history while deciding the award of contract. If no adverse action has been taken against the Bidder, the Bidder must provide a statement in its bid saying that there has been no such previous past performance resulting in adverse actions being taken against him.

- 2. The bidder shall provide an undertaking that:
 - (a) The proprietor/promoter/director of the firm, its employee, partner or representative is not convicted by a court of law following prosecution for offence involving moral turpitude in relation to business dealings including malpractices such as bribery, corruption, fraud, substitution of bids, interpolation, misrepresentation, evasion, or habitual default in payment of tax levied by law; etc.
 - (b) The firm does not employ a government servant, who has been dismissed or removed on account of corruption.
- Copies of original documents defining the constitution or legal status, place of registration, and principal place of business; written power of attorney of the signatory of the Bid to commit the Bidder;

Note:

- (a) The bidder must complete the check list given in Form 12 in Section V and submit it along with the Bid. It is essential that Bidders review carefully this Checklist to ensure that their Bid is complete and includes all required information.
- (b) The bidder should serially number all the documents of his bid, provide a summery table & sign/initial all the pages.
- (c) Details of two persons that RITES may contact for requests for clarification during bid evaluation:

Name	
Telephone No	
(direct)	
Email address	

(d) The Bank details from where the Bank Guarantee has been issued along with Phone, fax numbers and email Ids. For Banks from

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outside India the details of the correspondent Bank in India.

(e) Bidder should furnish Authority to the Purchaser to seek references from the Bidder's bankers.

7. One Bid per Bidder

7.1 A firm shall submit only one bid either individually or as a partner of a joint venture A firm that submits either individually or, as a member of a joint venture, more than one bid will cause all the proposals with the firm's participation to be disqualified.

8. Cost of Bidding

8.1 The Bidder shall bear all costs associated with the preparation and submission of its bid, and the Purchaser will in no case be responsible or liable for those costs, regardless of the conduct or outcome of the bidding process.

B. THE BIDDING DOCUMENTS

9. Content of Bidding Documents

9.1 The Bidding Documents are those stated below and should be read in conjunction with any addendum issued in accordance with ITB Clause 11.

Section I. Instructions to Bidders (ITB)
Section II. General Conditions of Contract (GCC)
Section III. Schedule of Requirements
Section IV. Technical Specifications
Section V. Sample Forms (including Contract Agreement)

9.2 The "Invitation for Bids" does not form part of the Bidding Documents and is included as a reference only. In case of discrepancies between the Invitation for Bid and the Bidding Documents listed in 9.1 above, said Bidding Documents will take precedence.

10. Clarification of Bidding Documents

10.1 A prospective Bidder requiring any clarification of the Bidding Documents shall contact the **Purchaser** in writing or by cable (for these ITB, the term "cable" is deemed to include electronic mail, telex, or facsimile) at the **Purchaser's** address **indicated in the clause 21.2 (b) of ITB.** The **Purchaser** will respond **in writing to any request for clarification received no later than fourteen (14) calendar days** prior to the deadline of submission of bids. Copies of the Purchaser's response shall be sent to all prospective Bidders who have purchased the Bidding Documents, including a description of the inquiry but without identifying its source.

Add as clause 10.2 to the ITB the following

Pre Bid meeting: - The bidder or his official representatives is invited to attend a pre bid meeting which will take place as per details given

Date: 12th September 2013

Time: 1400 hrs (IST)

Venue:

MSM Division,

RITES Ltd., RITES Office Complex,

Annex Building, 4th Floor, Plot No. 144, Sector 44,

Gurgaon - 122003, Harvana, India

Non-attendance at the pre bid meeting will not be a cause for disqualification of a bidder.

11. Amendment of Bidding Documents

- 11.1 At any time prior to the deadline for submission of bids, the Purchaser may amend the Bidding Documents by issuing Addenda.
- 11.2 Any addendum thus issued shall be part of the Bidding Documents pursuant to ITB Sub-Clause 9.1 and shall be communicated in writing to all purchasers of the Bidding Documents and will be binding on them. Bidders are required to immediately acknowledge receipt of any such amendment, and it will be assumed that the information contained in the amendment will have been taken into account by the Bidder in its bid.
- 11.3 To give prospective Bidders reasonable time in which to take the amendment into account in preparing their bids, the Purchaser shall extend, at its discretion, the deadline for submission of bids, in which case, the Purchaser will notify all Bidders by cable confirmed in writing of the extended deadline.

C. PREPARATION OF BIDS

12. Language of Bid

12.1 The bid, as well as all correspondence and documents relating to the bid exchanged by the Bidder and the Purchaser, shall be written in English language. Supporting documents and printed literature furnished by the Bidder may be in another language provided they are accompanied by an accurate translation of the relevant passages in the English language.

13. Documents Constituting the Bid

13.1 The bid submitted by the Bidder shall comprise the following:

- (a) duly filled-in Form of Bid and Price Schedule, in accordance with the forms indicated in Section V;
- (b) 'Integrity Pact' in accordance with ITB Clause 42.
- (b) original form of bid security in accordance with the provisions

of ITB Sub-Clause 18.3 (Bid Security);

- (c) written power of attorney authorizing the signatory of the bid to commit the Bidder;
- (d) documentary evidence establishing to the Purchaser's satisfaction, and in accordance with ITB Clause 6 that the Bidder/Manufacturer is qualified to perform the Contract if its bid is accepted.
- (e) Manufacturer's authorization Form 7, Section –V for bidder.

14. Bid Form

The Bidder shall complete the Bid Form and the Price Schedule furnished in the Bidding Documents, indicating the Goods to be supplied, a brief description of the Goods, their country of origin, and unit prices. (All details of the price components like taxes, duties etc. may also be indicated)

15. Bid Prices

- 15.1 The Bidder shall indicate on the Price Schedule, the unit price of each item, it proposes to supply under the Contract.
- 15.2 The bidder shall quote the prices on "Door Delivery Basis" to all consignees. The list of probable consignees is attached in schedule of requirement. However the list of consignees is the tentative list. The purchaser reserves the right to change any consignee at the time of placement of order.
- 15.3 Deleted.
- 15.4 The rate quoted should be both in words and figures. No figure or word should, be over written. Correction if any should be rewritten under the full signature of the person signing the tender.
- 15.5 The rate of Excise Duty and quantum of Excise should be shown distinctly. Similarly, Sales Tax/VAT, if any, where legally leviable and intended to be claimed extra should be shown distinctly as percentage along with the price quoted, separately. Where this is not done, no claim for excise duty and or Sales Tax/VAT will be admitted at any later stage on any ground.
- 15.6 (a) **Indigenous goods**:- Prices indicated on the Price Schedule shall be entered separately in the following manner:
 - (i) the price of the Goods quoted EXW (ex works, ex factory, ex warehouse, ex showroom, or off-the-shelf, as applicable), including all duties and sales tax and other duties and taxes already paid or payable: on the components and raw material

used in producing or manufacturing the Goods quoted ex works or ex factory;

- (ii) the rate and quantum of Excise duty and Sales Tax/VAT if any that will be payable on the Goods if the Contract is awarded.
- (iii) the price for inland transportation and other local costs incidental to delivery of the Goods to their final destination. The final destination is specified in Schedule of Requirements (Section III)
- (b) Imported goods: Offers for Imported origin goods shall clearly indicate firm, "All inclusive lump sum price" calculated in equivalent Indian Rupees and giving break up of as CIF (Indian Port), custom charges and other charges including inland transportation etc. The all inclusive lump sum price shall take care of impact of foreign exchange rate fluctuations etc., and accordingly arrive at the all inclusive lump sum price in equivalent Indian Rupees and this shall be the ceiling amount payable.

The terms EXW, CIF etc., shall be governed by the rules prescribed in the current edition of *Incoterms 2010* published by the International Chamber of Commerce, Paris.

- 15.7 The prices quoted by the bidder should be on firm and fixed basis during the performance of the contract. A bid submitted with adjustable price quotation will be treated as non responsive and will be rejected pursuant to ITB clause 28.
- 15.8 The bidder's separation of price components in accordance with clause above will be solely for the purpose of facilitating the comparison of bids by the purchaser and will not in any way limit the purchaser's right to contract on any of the terms offered.
- 15.9 The purchaser shall not be liable to any claim on account of fresh imposition and/or increase of Excise Duty, Customs duty, Sales Tax on raw materials and/or components used directly in the manufacture of the contracted stores taking place during the pendency of the contract.
- 15.10 Statutory variation in taxes and duties on finished product will be on purchaser's account.

16. Currencies of Bid

- 16.1 Prices shall be quoted in Indian Rupees only.
- 17. Period of
- 17.1 Bids shall remain valid for the period of **150 days** after the date of

Validity of Bids

bid submission specified in ITB Clause 22 i.e. up to **28th February 2014**. A bid valid for a shorter period shall be rejected by the Purchaser as non-responsive.

17.2 In exceptional circumstances, prior to expiry of the original bid validity period, the Purchaser may request that the Bidders extend the period of validity for a specified additional period. The request and the responses thereto shall be made in writing. A Bidder may refuse the request without forfeiting its bid security.

18. Bid Security

18.1 the Bidder shall furnish, as part of its bid, a bid security against each schedule in fixed amount as specified in Section –III, Schedule of Requirement. The amount of bid security against each schedule(s) should be in fixed amount as specified in the Schedule of Requirements.

If the bidder is submitting bid for more than one schedule, the amount of the bid security shall be the sum of bid securities required for the respective schedules. The bidder has the option to submit individual bid security instrument for different schedules.

If the amount of bid security furnished is less than the required for total quoted schedules by the bidders, and then Bid security will be considered valid only for the quoted schedules (in serial order of the Schedule of Requirement). The later schedule(s) for which Bid security fall short, will be treated as non-responsive.

- 18.2 The bid security shall remain valid for a period of **45 days** beyond the validity period for the bid i.e. up to **14**th **April 2014** and beyond any extension subsequently requested under Sub-clause 17.1.
- 18.3 The bid security shall be denominated in Indian Rupees, and shall be, at the Bidder's option, in one of the following forms:
 - (a) a crossed demand draft or a pay order drawn in favour of the Purchaser;
 - (b) a (bank) guarantee issued by a nationalized/scheduled bank in India. The format of the (bank) guarantee shall be in accordance with the form of bid security included in Section V.
- 18.4 Any bid not accompanied by an acceptable bid security shall be rejected by the Purchaser as non-responsive.
- 18.5 The bid securities of unsuccessful Bidders will be returned as promptly as possible.
- 18.6 The bid security of the successful Bidder will be returned when the Bidder has signed the Agreement and furnished the required

performance security.

18.7 The bid security may be forfeited

- (a) if the Bidder withdraws its bid, except as provided in ITB Sub-Clauses 17.2 and 24.3; or
- (b) if the Bidder does not accept the correction of its bid price, pursuant to ITB Clause 29; or
- (c) in the case of a successful bidder, if the Bidder fails within the specified time limit to:
 - (i) sign the contract, or
 - (ii) furnish the required performance security, or
 - (iii) In case of any false, incorrect or misleading information provided in the bid.

19. Alternative Proposals by Bidders

Alternative bids shall not be accepted. The bidder should not submit more than one bid for any Schedule.

20. Format and Signing of Bid

- 20.1 The Bidder shall prepare one original and one copy of the bid, clearly marking each one as "ORIGINAL BID" and "COPY OF BID," as appropriate. In the event of any discrepancy between them, the original shall govern.
- 20.2 The original and all copies of the bid, each consisting of the documents listed in ITB Sub-Clause 13.1, shall be typed or written in indelible ink and shall be signed by the Bidder or a person or persons duly authorized to bind the Bidder to the Contract. The later authorization shall be indicated by written power of attorney, which pursuant to ITB Sub-Clause 13.1 (c) shall accompany the bid.
- 20.3 Any interlineations, erasures, or overwriting to correct errors made by the Bidder should be initialled by the person or persons signing the bid.

D. Submission of Bids

21. Sealing and Marking of Bids

21.1 The Bidder shall enclose the original and each copy of the bid in separate sealed envelopes, duly marking the envelopes as "ORIGINAL" and "COPY." The envelopes containing the original and copies shall then be enclosed in another envelope.

- 21.2 The inner and outer envelopes shall:
 - (a) bear the name and address of the Bidder;
 - (b) be addressed to the Purchaser at the address given below

Group General Manager/MSM RITES Ltd., MSM Division, RITES Office Complex, Annex Building, 4th Floor, Plot No. 144, Sector 44, Gurgaon – 122003, Haryana, India

(c) The inner and outer envelopes shall bear the following additional identification marks:

Invitation for Bids Title: Invitation for Bids Number: Schedule Number: Time & Date of Submission of Bids: Name of the Goods

- (d) bear a statement "DO NOT OPEN BEFORE 1st October 2013 at 14:15 hrs" to be completed with the time and date specified in the ITB clause 22.1.
- 21.3 If the outer envelope is not sealed and marked as required by ITB Sub-Clause 21.2, the Purchaser will assume no responsibility for the misplacement or premature opening of the bid.
- 22. Deadline for Submission of Bids
- 22.1 Bids must be received by the Purchaser at the address specified in the ITB Sub-Clause 21.2 (b) no later than the time and date specified below:-

Bids must be delivered before 14:00 Hrs. on 1st October 2013. Late bids will be rejected.

"In event of the specified date for the submission of Bids being declared a holiday for the Purchaser, the Bids will be received up to the appointed time on the next working day".

- 22.2 The Purchaser may, at its discretion, extend the deadline for the submission of bids by amending the Bidding Documents in accordance with ITB Sub-Clause 11.3, in which case all rights and obligations of the Purchaser and Bidders previously subject to the deadline will thereafter be subject to the deadline as extended.
- 23. Late Bids
- 23.1 Any bid received by the Purchaser after the deadline for submission

of bids prescribed by the Purchaser in the ITB Clause 22 will be rejected and returned unopened to the Bidder.

24. Modification and Withdrawal of Bids

24.1 The Bidder may modify or withdraw its bid after submission, provided that written notice of the modification, or withdrawal of the bids duly signed by an authorized representative, is received by the Purchaser prior to the deadline prescribed for submission of bids.

Note: No bid may be modified subsequent to the deadline for submission of bid.

- 24.2 The Bidder's modification shall be prepared, sealed, marked, and dispatched as follows:
 - (a) The Bidder shall provide an original and the number of copies specified in the ITB clause 20.1 of any modifications to its bid, clearly identified as such, in two inner envelopes duly marked "BID MODIFICATION-ORIGINAL" and "BID MODIFICATION-COPY." The inner envelopes shall be sealed in an outer envelope, which shall be duly marked "BID MODIFICATION."
 - (b) Other provisions concerning the marking and dispatch of bid modifications shall be in accordance with ITB Sub-Clauses 21.2 and 21.3.
- 24.3 A Bidder wishing to withdraw its bid shall notify the Purchaser in writing prior to the deadline prescribed for bid submission. A withdrawal notice shall be received prior to the deadline for submission of bids. The notice of withdrawal shall:
 - (a) be addressed to the Purchaser at the address named in the ITB clause 21.2 (b)
 - (b) bear the specific identification of the bidding process (Contract name), the IFB title and IFB number, and the words "BID WITHDRAWAL NOTICE," and
 - (c) be accompanied by a written power of attorney authorizing the signatory of the withdrawal notice to withdraw the bid.
- 24.4 Bids requested to be withdrawn in accordance with ITB Sub-Clause 24.3, shall be returned unopened to the Bidders.
- 24.5 No bid may be withdrawn in the interval between the bid submission deadline and the expiration of the bid validity period specified in ITB Clause 17. Withdrawal of a bid during this interval may result in the

forfeiture of the Bidder's bid security, pursuant to ITB Sub-Clause 18.7.

E. OPENING AND EVALUATION OF BIDS

25. Bid Opening

25.1 The Purchaser will open all bids, including withdrawal notices and modifications, in public, in the presence of Bidders' representatives who choose to attend, at 14.15 hrs, on the date, and at the place specified below:

Time, date, and place for bid opening are: **1415 hrs** (Indian Standard Time) on **1**st **October 2013** at the following address:

RITES Ltd., MSM Division, RITES Office Complex, Annex Building, 4th Floor, Plot No. 144, Sector 44, Gurgaon – 122003, Haryana, India

Add at the end of this clause:

"In the event of the specified date of the bid opening being declared a holiday for the Purchaser, the bids shall be opened at the appointed time and Location on the next working day."

Bidders' representatives shall sign a register as proof of their attendance. All bids must be accompanied by a bid security as specified in Section –III, Schedule of Requirements.

In case the bidder uses an agent in any capacity (including for attending pre-bid meetings or bid opening meetings), the Purchaser will be informed in writing by the bidders regarding the appointment of such agent and a copy of the agreement signed between the bidder and the agent (which will include the scope of services provided by such agent and amount payable by the bidder) will be shared with the Purchaser in advance. The agreement should be legally binding with the clear understanding that the Bidder will be held responsible for unlawful actions (viz. fraudulent representation, bribing or collusion) of the agent. If this condition is not complied, such agents will not be allowed to attend the meetings and also no queries from such agents will be entertained by the Purchaser. In addition, the bidder will ensure that such agent should not work simultaneously for two or more competing bidders.

25.2 Envelopes marked "WITHDRAWAL" shall be read out and the envelope with the corresponding bid shall not be opened but returned to the Bidder. No bid withdrawal shall be permitted unless the corresponding withdrawal notice is read out at bid opening. Envelopes marked "MODIFICATION" shall be read out and opened

with the corresponding bid.

- 25.3 Bids shall be opened one at a time, reading out: the name of the Bidder and whether there is a modification; the bid price of each item or lot, as the case may be, including discounts the presence or absence of a bid security, if required; the presence or absence of requisite powers of attorney; and any other such details as the Purchaser may consider appropriate. No bid shall be rejected at bid opening except for late bids pursuant to Sub-Clause 23.1.
- 25.4 Bids (and modifications sent pursuant to ITB Sub-Clause 24.2) that are not opened and read out at bid opening shall not be considered further for evaluation, irrespective of the circumstances.
- 25.5 The Purchaser will prepare minutes of the bid opening at the end of the opening session, including, as a minimum: the name of the Bidder and whether there was a withdrawal or modification; the bid price; including any discounts or alternatives offered if permitted in the Bid Data Sheet; the presence or absence of a bid security; the presence or absence of requisite powers of attorney.
- 25.6 The Bidder's representatives who are present shall be requested to sign the minutes. The omission of a Bidder's signature on the minutes shall not invalidate the content and effect of the minutes. The minutes should be distributed to all Bidders who request them.

26. Clarification of Bids

26.1 During evaluation of the bids, the Purchaser may, at its discretion, ask the Bidder for a clarification of its bid. The request for clarification and the response shall be in writing, and no change in the prices or substance of the bid shall be sought, offered, or permitted, except to correct arithmetic errors identified by the Purchaser in the evaluation of the bids, in accordance with ITB Sub-Clause 29.1.

27.Confidentiality

- 27.1 Information relating to the examination, clarification, evaluation, and comparison of bids, and recommendations for the award of a Contract shall not be disclosed to bidders or any other persons not officially concerned with such process until the notification of Contract award is made to all Bidders.
- 27.2 Any effort by the bidder to influence the Purchaser in the Purchaser's bid evaluation, bid comparison, or contract award decisions may result in the rejection of the Bidder's bid.
- 27.3 From the time of bid opening to the time of Contract award, if any Bidder wishes to contact the Purchaser on any matter related to its bid, it should do so in writing.

28. Examination of Bids and Determination of Responsivenes s

- 28.1 The Purchaser will examine the bids to determine whether they are complete, whether any computational errors have been made, whether required sureties have been furnished, whether the documents have been properly signed, and whether the bids are generally in order.
- 28.2 The Purchaser may waive any minor informality, nonconformity, or irregularity in a bid that does not constitute a material deviation, provided such waiver does not prejudice or affect the relative ranking of any Bidder.
- Prior to the detailed evaluation, pursuant to ITB Clause 30, the Purchaser will determine whether each bid is of acceptable quality, is complete, and is substantially responsive to the Bidding Documents. For purposes of this determination, a substantially responsive bid is one that conforms to all the terms, conditions, and specifications of the Bidding Documents without material deviations, exceptions, objections, conditionality's, or reservations. A material deviation, exception, objection, conditionality, or reservation is one: (i) that limits in any substantial way the scope, quality, or performance of the Goods and related Services; (ii) that limits, in any substantial way that is inconsistent with the Bidding Documents, the Purchaser's rights or the successful Bidder's obligations under the Contract; and (iii) that the acceptance of which would unfairly affect the competitive position of other Bidders who have submitted substantially responsive bids.

The following clauses are the critical provisions deviations from or objections or reservations to which, will be treated as material deviations:

- Non submission of Bid Form
- Bid Validity (ITB Clause 17)
- Bid Security (ITB Clause 18);
- Validity of Bid Security (ITB Clause 18.2)
- Performance Security (GCC Clause 8);
- Delivery Terms (GCC Clause 11 & Schedule of Requirements)
- Warranty (GCC Clause 15);
- Payment terms (GCC Clause 16)
- Force Majeure (GCC Clause 24);
- Limitation of liability (GCC Clause 28)
- Applicable Law (GCC Clause 30);
- Taxes and Duties (GCC Clause 32);
- Technical Specification (As per Section IV)
- Delivery Period (Schedule of Requirements)

Above list is not exhaustive

28.4 If a bid is not substantially responsive, it will be rejected by the Purchaser and may not subsequently be made responsive by the Bidder by correction of the nonconformity. The Purchaser's determination of a bid's responsiveness is to be based on the contents of the bid itself without recourse to extrinsic evidence.

29. Correction of Errors

29.1 Arithmetical errors will be rectified as follows. If there is a discrepancy between the unit price and the total price that is obtained by multiplying the unit price and quantity, the unit or subtotal price shall prevail. If there is a discrepancy between subtotals and the total price, the total price shall be corrected. If there is a discrepancy between words and figures, the amount in words will prevail. If a Bidder does not accept the correction of errors, its bid will be rejected and its bid security may be forfeited.

30. Evaluation and Comparison of Bids

30.1 The Purchaser will evaluate and compare the bids that have been determined to be substantially responsive, pursuant to ITB Clause 28.

- 30.2 The Purchaser's evaluation of a bid will take into account the total unit cost of the item at the consignee's destination inclusive of all duties, taxes and other charges.
- 30.3 The contract shall be awarded only to the bidder who are substantially responsive, offer competitive rates, and meet the qualification requirement stipulated in the bidding documents.
- 30.4 Bidder may bid for one or more schedules. Bids will be evaluated for each schedule separately and the contract will comprise the schedules(s) awarded to the successful bidder. Bidders must quote for the entire quantity of each schedule. Bidders who do not quote for full quantity of the schedule will be treated as non-responsive.
- 30.5 Deviations in the delivery schedule and Payment schedule are not permitted.
- 30.6 In exercising of the powers conferred in Section 11 of the Micro, Small and Medium Enterprises Development (MSMED) Act 2006, the Government has notified a new Public Procurement Policy for Micro & Small Enterprises effective from 1st April 2012.

In accordance to the above notification the participating Micro and Small Enterprises (MSEs) in a Bid, quoting price within the band of L 1+15% would be allowed to supply a portion of the requirement by bringing down their price to the L 1 price, in a situation where L 1 price is from someone other than an MSE. Such MSEs would be allowed to supply up to 20% of the total Bid value. In case there are more than one such eligible MSE, the 20% supply will be shared equally. Out of 20% of the quantity earmarked for supply from MSEs, 4% quantity is earmarked for procurement from MSEs owned by SC/ST entrepreneurs. However, in the event of failure of such MSEs to participate in the Bid process or meet the Bid requirements and the L 1 price, the 4% quantity earmarked for MSEs owned by

SC/ST entrepreneurs will be met from other participating SMEs.

The MSEs participating in the bid shall enclose with their Bid a copy of their valid registration certificate with District Industries Centres or Khadi and Village Industries Commission or Coir Board or NSIC or any other body specified by Ministry of Micro and Small enterprices in support of their being an MSE, failing which their offer will be liable to be ignored.

F. AWARD OF CONTRACT

31. Post qualification

- 31.1 The Purchaser will determine to its satisfaction whether the Bidder that is selected as having submitted the lowest evaluated responsive bid is qualified to perform the Contract satisfactorily, in accordance with the criteria listed in ITB Sub-Clause 6.1.
- 31.2 The determination will evaluate the Bidder's financial, technical, and production capabilities. It will be based on an examination of the documentary evidence of the Bidder's qualifications submitted by the Bidder, pursuant to ITB Sub-Clause 6.1, as well as other information the Purchaser deems necessary and appropriate.
- 31.3 An affirmative post qualification determination will be a prerequisite for award of the contract to the lowest evaluated Bidder. A negative determination will result in rejection of the Bidder's bid, in which event the Purchaser will proceed to the next-lowest evaluated Bidder to make a similar determination of that Bidder's capabilities to perform satisfactorily.

32. Award Criteria

- 32.1 Pursuant to ITB Clauses 30, 31, and 35, the Purchaser will award the Contract to the Bidder whose bid has been determined to be substantially responsive and has been determined to be the lowest evaluated bid, provided further that the Bidder is determined to be qualified to perform the Contract satisfactorily, pursuant to ITB Clause 31.
- 33. Purchaser's
 Right to Accept
 Any Bid and to
 Reject Any or
 All Bids
- 33.1 The Purchaser reserves the right to accept or reject any bid, or to annul the bidding process and reject all bids at any time prior to contract award, without thereby incurring any liability to the affected Bidder or Bidders. No reason for such action of Purchaser shall be given. 333
- 34. Purchaser's right to vary quantities during currency of contract
- 34.1 The purchaser reserves the right to increase or decrease the ordered quantity of goods by 25% during the currency of contract.

35. Notification of Award

- 35.1 Prior to the expiration of the period of bid validity, the Purchaser will notify the successful Bidder in writing by registered letter or by fax, to be subsequently confirmed in writing by registered letter, that its bid has been accepted for award of contract.
- 35.2 Upon the successful Bidder's furnishing of the signed Contract Form and performance security pursuant to ITB Clause 38, the Purchaser will promptly notify each unsuccessful Bidder and will discharge its bid security, pursuant to ITB Clause 18.
- 36. Publication of Bid result
- 36.1 The name and address of Successful bidder(s) will be declared and published appropriately.
- 37. Signing of Contract
- 37.1 Promptly after the Purchaser notifies the successful Bidder that its bid has been accepted, the Purchaser will send the Bidder the Contract Form provided in the Bidding Documents, incorporating all agreements between the parties.
- 37.2 Within twenty-one (21) days of receipt of the Contract Form, the successful Bidder shall sign the Contract Form and return it to the Purchaser.

38. Performance Security

- 38.1 With in twenty one days (21) days of the receipt of notification of award from the purchaser, the successful bidder shall furnish the performance security in accordance with the conditions of contract, using the performance security form provided in the bidding documents, or any another form acceptable to the purchaser.
- 38.2 Failure of the successful Bidder to comply with the requirement of ITB Clause 37 or ITB Sub-Clause 38.1 shall constitute sufficient grounds for the annulment of the award and forfeiture of the bid security, in which event the Purchaser may make the award to the next-lowest evaluated bid submitted by a qualified Bidder or call for new bids.

39. Clarification on Duties & Taxes

39.1 **EXCISE DUTY**

- 39.1.1 The price quoted should be-EXW and the rate of excise duty and quantum of Excise Duty separately should be shown distinctly. In the absence of any such stipulation it will be presumed that the price includes Excise Duty and no claim for the same will be entertained. If case of stipulation like excise duty extra as applicable, the quoted prices will be loaded with the maximum quantum of excise duty which is normally applicable on the item in question for the purpose of comparing the prices with other bidders.
- 39.1.2 If a bidder is exempted from payment of excise duty up to any

monetary limit of supplies, he should clearly state that no excise duty will be charged by him up to the limit of exemption which he may have. If any concession is available in regard to the rate/quantum of Central Excise Duty, it should be brought out clearly. Stipulations like excise duty presently not applicable but the same will be charged if it becomes leviable later on, will not be accepted (unless in such cases it is clearly stated by the bidder that excise duty will not be charged by him even if the same becomes applicable later on). In respect of the bidders who fail to comply with this requirement, their quoted prices will be loaded with the maximum quantum of excise duty which is normally applicable on the item in question for the purpose of comparing their prices with other bidders.

- 39.1.3 Any change in Excise Duty upward/downward as a result of any statutory variation in excise, on the finished goods, taking place during currency of contract shall be allowed to the extent of actual quantum of excise duty paid by the supplier. Similarly in case of downward revision in excise duty, the actual quantum of reduction in excise duty shall be reimbursed to the Purchaser by the Supplier. All such adjustments shall include all relief's, exemptions, rebates, concessions etc if any obtained by the supplier.
- 39.1.4 Bidders should note that in case any refund of excise duty is granted to them by excise Authorities in respect of goods-supplied under the contract they will pass on the credit to the purchaser immediately along with a certificate from their Director /Manager/ Proprietor/Accountant that the credit so passed on relates to the excise originally paid for the goods supplied under the contract. In case of failure to do so within 10 days of the issue of the excise duty refund orders to them by the Excise Authorities, the purchaser would be empowered to deduct a sum equivalent to the amount refunded by the Excise authorities without any further reference to them from any of their outstanding bills against the contract or any other pending Government contract and that no disputes on this account would be raised by them.
- 39.1.5 The purchaser shall not be liable for any claim on account of fresh imposition and/ or increase of Excise Duty on raw materials and/or components used directly in the manufacture of the contracted stores taking place during the pendency of the contract.
- 39.1.6 The tenderer should indicate in their offer whether they are registered with Excise authorities for availing CENVAT credit or not. If they are availing CENVAT CREDIT, they should take into account the entire credit on inputs available under CENVAT CREDIT Scheme while quoting the price and furnish a declaration

to this effect.

39.2 **SALES TAX /VAT**

39.2.1 The price quoted should be exclusive, of Sales Tax/VAT. The element of CST/VAT leviable should be specifically stated and shown distinctly as a percentage along with the price-quoted, separately. Where this is not done, no claim for sales tax will be admitted at any later stage on any ground. Further in the absence of any such stipulation regarding sales tax in the bid, it will be presumed that the prices quoted by the bidder are inclusive of sales tax and no liability for payment of sales tax will be devolved up on the purchaser. If case of stipulation like Sales Tax/VAT extra as applicable, the quoted prices will be loaded with the maximum quantum of Sales Tax/VAT which is normally applicable on the item in question for the purpose of comparing the prices with other bidders.

Any change in Sales Tax upward/downward as a result of any statutory variation in element of CST/VAT leviable, on the finished goods, taking place during currency of contract shall be allowed to the extent of actual quantum of CST/VAT paid by the supplier. Similarly in case of downward revision in CST/VAT, the actual quantum of reduction in CST/VAT shall be reimbursed to the Purchaser by the Supplier. All such adjustments shall include all relief's, exemptions, rebates, concessions etc if any obtained by the supplier.

- 39.2.2 For the bidder quoting sales tax extra, sales tax will be paid to the bidder at the rate at which it is liable to be assessed or has actually been assessed provided the transaction of sales is legally liable to sales tax and the same is payable as per terms of the contract.
- 39.2.3 The purchaser shall not be liable for any claim on account of fresh imposition and/or increase of sales tax/VAT on raw materials and or components used directly in the manufacture of the contracted goods taking place during the pendency of the contract.
- 39.2.4 The bidder shall unconditionally pass on applicable input tax credit or set off of tax paid on raw material under the relevant VAT/Sales Tax Act availed on inputs used in manufacture of the finished product. The bidder shall furnish a declaration to this effect.

39.3 OCTROI DUTY AND LOCAL T AXES

39.3.1 Goods to be supplied to Govt. Departments against Government Contracts are exempted from levy of Town duty, Octroi Duty, Terminal Tax and other levies of local bodies. The local Town/Municipal Body regulations at times, however, provide for such Exemption only on Production of such exemption certificate

from an authorised officer. Supplier should ensure that, goods ordered against contracts placed by this department are exempted from levy of Town Duty, Octroi Duty, Terminal Tax or other Local Taxes and Duties. Wherever required, supplier should obtain the exemption certificate from the concerned office to avoid local taxes or duties.

39.3.2 In case where the Municipality or other local body insists upon payment of these duties or taxes, the same should be paid by the supplier to avoid delay in supplies and possible demurrage charges. The receipt obtained for such payment should be forwarded to the officer concerned without delay together with a copy of the relevant act or by laws/notifications of the Municipality or the Local body concerned to enable him to take up the question of refund with the concerned bodies, if admissible under the said acts or rules.

39.4 **CUSTOMS DUTY**

In respect of imported stores offered, the bidder shall specify the rate as well as the total amount of customs duty payable, on the quoted goods in the price schedule. The bidder shall also indicate the corresponding Indian Customs Tariff Number applicable for the goods in question.

Any variation to the custom duty during the currency of the contract will be reimbursed to the bidder/refunded by the bidder. However no upward variation will be reimbursed to the bidder after the expiry of the original delivery period.

40. Purchase preference

The Purchaser reserves the right to give purchase preference to the Micro and Small Scale Enterprises as per the policies of Govt. of India in vogue, for which bidder should produce valid copy of his registration as Micro or Small Scale Enterprise.

41. Registration of Imported goods

Bidder intending to supply the imported goods must ensure that the goods and the manufacturing facilities of the manufacturer are registered with the relevant authorities in India, as for relevant laws of the country on the date of bid opening. Bidders are advised to visit website www.cdsco.nic.in for necessary information on the subject. Bidders are required to furnish a copy of the aforesaid registration along with their bid.

42. Integrity Pact

42.(i) The Bidder/Supplier is required to enter into an Integrity Pact with the Purchaser, in the Format at Sample Forms Section V. The Integrity Pact enclosed as Form No.12 will be signed by RITES for and on behalf of Purchaser as its Agent/Power of Attorney Holder at the time of execution of Agreement with the successful Bidder. While submitting the Bid, the Integrity Pact shall be signed by the duly authorized signatory of the Bidder/Lead Member of JV. In case of

failure to submit the Integrity Pact duly signed and witnessed, along with the Bid, the Bid is likely to be rejected.

42.(ii) In case of any contradiction between the Terms and Conditions of the Bid Document and the Integrity Pact, the former will prevail.

Name and Address of the Independent External Monitor (In case value of contract is Rs.10 crores or more): Shri B. S. Minhas, A-29, Bhairon Marg, Hanuman Nagar, Jaipur-302021

Name, Designation and Address of RITES' Liaison Officer (in case value of contract is less than Rs.10 crores): Shri Y. K. Sharma, GM/CP

SECTION II. GENERAL CONDITIONS OF CONTRACT

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General Conditions of Contract

1. Definitions

- 1.1 In this Contract, the following terms shall be interpreted as indicated:
- (a) "The Contract" means the agreement entered into between the Purchaser and the Supplier, as recorded in the Contract Form signed by the parties, including all attachments and appendices thereto and all documents incorporated by reference therein.
- (b) "The Contract Price" means the unit price payable to the Supplier under the Contract for the full and proper performance of its contractual obligations.
- (c) "Day" means calendar day.
- (d) "Effective Date" means the date on which this Contract becomes effective i.e. date of notification of Award.
- (e) "GCC" means the General Conditions of Contract contained in this section.
- (f) "The Goods" means all of the pharmaceuticals including nutritional supplement and oral and injectable forms of contraception, vaccines, and condoms that the Supplier is required to supply to the Purchaser under the Contract.
- (g) "The Purchaser" means Ministry of Health & Family Welfare, Govt. of India through RITES Ltd, New Delhi.
- (h) "The Purchaser's Country" is India.
- (i) "Registration Certificate" means the certificate of registration or other documents in lieu thereof establishing that the Goods supplied under the Contract are registered for use in India in accordance with the Applicable Law.
- (j) "The Services" means those services ancillary to the supply of the Goods, such as transportation and insurance, and any other incidental services, such as provision of technical assistance, training, and other such obligations of the Supplier covered under the Contract.
- (k) "The Site," where applicable, means the place or places named in the Schedule of requirement.
- (1) "The Supplier" means the individual or firm supplying the Goods

and Services under this Contract.

(m) End user means the organization(s) where the goods will be used. The end user is the consignee stated in the Schedule of Requirements

2. Imports

For Import origin goods quoted, the supplier or the Indian agent shall have to arrange at his own cost, all import/custom clearance handling facilities. The purchaser shall not be liable to any claim on account of fresh imposition and/or increase of Excise Duty, Custom Duty, Sales Tax on raw materials and /or components used directly in the manufacture of the contracted goods taking place during the pendency of the contract.

- 3. Application
- 3.1 These General Conditions shall apply to the extent that they are not superseded by provisions of other parts of the Contract.
- 4. Country of Origin
- 4.1 Any Goods and Services supplied under the Contract shall have their origin in India or eligible countries (in case of imported goods offered).
- 5. Standards
- 5.1 The Goods supplied under this Contract shall conform to the standards mentioned in the Technical Specifications.
- 6. Use of Contract
 Documents and
 Information;
 Inspection and
 Audit by the
 Purchaser
- 6.1 The Supplier shall not, without the Purchaser's prior written consent, disclose the Contract, or any provision thereof, or any specification, plan, drawing, pattern, sample, or information furnished by or on behalf of the Purchaser in connection therewith, to any person other than a person employed by the Supplier in the performance of the Contract. Disclosure to any such employed person shall be made in confidence and shall extend only so far as may be necessary for purposes of such performance.
- 6.2 The Supplier shall not, without the Purchaser's prior written consent, make use of any document or information enumerated in GCC Sub-Clause 6.1 except for purposes of performing the Contract.
- 6.3 Any document, other than the Contract itself, enumerated in GCC Sub-Clause 6.1 shall remain the property of the Purchaser and shall be returned (all copies) to the Purchaser on completion of the Supplier's performance under the Contract if so required by the Purchaser.
- 6.4 The Supplier shall permit the Purchaser to inspect the Supplier's accounts and records relating to the performance of the Contract and to have them audited by auditors appointed by the Purchaser, if

so required by the Purchaser.

7. Patent Rights

7.1 The Supplier shall indemnify the Purchaser against all third-party claims of infringement of patent, trademark, or industrial design rights arising from use of the Goods or any part thereof in India.

8. Performance Security

- 8.1 Within twenty-eight (28) days of receipt of the notification of Contract award, the successful Bidder shall furnish to the Purchaser the performance security in the amount equal to 10 % of the total contract price.
 - a) In the event of any amendment issued to the Contract, the Supplier shall, within twenty-one (21) days of issue of the amendment, furnish the corresponding amendment to the Performance Security (as necessary) rendering the same valid in all respects in terms of the Contract, as amended.
 - b) The performance security shall be valid till **60 days** after the date of completion of all contractual obligations including warranty.
- 8.2 The proceeds of the performance security shall be payable to the Purchaser as compensation for any loss resulting from the Supplier's failure to complete its obligations under the Contract.
- 8.3 The performance security shall be denominated in Indian Rupees, and shall be in one of the following forms:
 - (a) The performance security shall be in the form of a Bank guarantee and the named beneficiary shall be "RITES Ltd" (acting as procurement agent on behalf of Ministry of Health & Family Welfare Government of India), issued by a nationalized/scheduled bank located in India and acceptable to the Purchaser, in the format provided in the Bidding Documents; or
 - (b) a crossed demand draft or a pay-order drawn in favour of RITES Ltd.
- 8.4 The performance security will be discharged by the Purchaser and returned to the Supplier not later than thirty (30) days following the date of completion of the Supplier's performance obligations under the Contract, including any warranty obligations.

In the event of any amendment issued to the contract, the supplier shall, within twenty one (21) days of issue of the amendment, furnish the corresponding amendment to the Performance Security (as necessary) rendering the same valid in all respects in terms of the contract, as amended.

9. Inspections and Tests

9.1 The Purchaser or its representative shall have the right to inspect and/or to test the Goods to confirm their conformity to the Contract specifications.

The Technical Specifications (Section IV) shall specify what inspections and tests the Purchaser requires. Further,

- (a) Pre-dispatch inspection of the supplies shall be conducted by purchaser or its authorised representative retained by the purchaser for these purposes. The Purchaser shall notify the Supplier in writing, in a timely manner, of the identity of any representatives retained for these purposes. The Supplier shall at the earliest furnish details of number of batches and visits for inspection and testing to enable the pre-dispatch inspection and testing when undertaken.
- (b) Said inspection and testing is for the Purchaser's account. In the event that inspection and testing is required prior to dispatch, the Goods shall not be shipped unless a satisfactory inspection and quality control report has been issued in respect of those Goods.

The related costs of the pre-shipment inspection for the first inspection of goods shall be borne by the Purchaser. However, if goods are offered for inspection in smaller lots than specified in contract then supplier will have to bear the additional inspection charges. The goods consumed during tests will be on suppliers account. The cost of subsequent inspections and related costs, due to rejection of Goods at the first inspection shall be borne by the Supplier. Inspection will be done by a Purchaser's agent to ascertain whether the Goods are in conformity with the technical specifications of the contract or not.

The supplier shall put up the goods for such inspection to the purchaser's inspector 15-25 days (depending on the time required for pre-dispatch inspection & testing) ahead of the contractual delivery period, so that deliveries to the consignees are completed as per the contractual delivery period.

- (c) The Supplier may have an independent quality test conducted on a batch ready for shipment. The cost of such tests will be borne by the Supplier.
- (d) Upon receipt of the Goods at place of final destination, the

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end user/consignee shall have the right to inspect the Goods or part of the Goods to ensure that they conform to the condition of the Contract and advise the Purchaser that the Goods were received in apparent good order. The end user/consignee will issue an Acceptance Certificate to the Supplier in respect of such Goods (or part of Goods). The Acceptance Certificate should normally be issued within twenty one (21) days of receipt of the Goods or part of Goods at place of final destination.

- (e) Batch wise inspection of goods shall be carried out by Purchaser's representative.
- 9.2 Where the Supplier contests the validity of the rejection by the Purchaser or his representative, of any inspection as required by 9.1 above, conducted before shipment or at ultimate destination, whether based on product or packing grounds, a sample drawn jointly by the Supplier and Purchaser or his or her representative and authenticated by both, will be forwarded for umpire analysis within four weeks of the time the Supplier contests to an independent agency mutually agreed by the Purchaser and Supplier. The umpire's finding, which will be promptly obtained, will be final and binding on both parties. The cost of umpire analysis will be borne by the losing party.

10. Packing

- 10.1 The Supplier shall provide such packing of the Goods as is required to prevent their damage or deterioration during transit to their final destination, as indicated in the Contract. The packing shall be sufficient to withstand, without limitation, rough handling during transit and exposure to extreme temperatures, salt, and precipitation during transit and open storage. Packing case size and weights shall take into consideration, where appropriate, the remoteness of the Goods' final destination and the absence of heavy handling facilities at all points in transit.
- 10.2 The packing, marking, and documentation within and outside the packages shall comply strictly with such special requirements as shall be expressly provided for in the Contract, including additional requirements strictly as per Technical Specifications, and in any subsequent instructions ordered by the Purchaser.

Packing and Marking shall be strictly as per Technical Specifications and will be inspected in terms of provisions of specifications before clearing for dispatch. The Bar coding requirement shall also be properly understood and marked on the package as per the provision of the specification.

11. Delivery and

Documents

(A) Documents to be submitted to purchaser:-

Upon the delivery of the Goods, the Supplier shall notify the Purchaser in writing and deliver to the Purchaser four sets of documents comprising of the following:

- One original and three copies of commercial invoice, indicating the RITES Ltd as the Purchaser on behalf of Ministry of Health & Family Welfare, Govt. of India, the Contract number, credit number; Goods' description, quantity, unit price, and total amount. Invoices must be signed in original and stamped or sealed with the company stamp/seal;
- ii. Four copies of Proof of Dispatch (POD), viz., Railway consignment note/road consignment note or multimodal transport document showing Purchaser as RITES Ltd. on behalf of Ministry of Health & Family Welfare, Govt. of India and delivery up to final destination as stated in the Contract
- iii. One original & 3(three) copies of Acknowledgement of receipt of Goods/Final Acceptance Certificate by the Consignees, as per the format.
- iv. Four copies of packing list identifying contents of each package
- v. One original and three copies of the manufacturer's or Supplier's Warranty certificate covering all items supplied
- vi. Four copies of Certificate of Inspection furnished to Supplier by the nominated inspection agency (where inspection is required)
- vii. Four copies of Internal Test Analysis Report of drugs and pharmaceuticals of the Manufacturer
- viii. Four copies of notification of the local tax authority in support of rate of tax indicated in invoice.
- ix. Any other/additional procurement-specific document(s) s required for delivery/payment purposes.

(B) Documents to be submitted to Consignee:-

The Supplier should intimate the Consignee at least 7 days in advance before the dispatch of Goods, the expected date of

arrival of Goods along with quantity of Goods. Along with each consignment the Supplier should provide the Consignee one set of the documents mentioned below:

- (i) Copy of NOA
- (ii) Copy of Invoice containing particulars as per (A)(i) above;
- (iii) Packing list identifying contents of each package
- (iv) Manufacturer's or Supplier's Warranty certificate covering all items supplied.

12. Insurance

Deleted

13. Transportation

- 13.1 Where the Supplier is required under the Contact to transport the Goods to a specified place of destination within India, defined as the Site, transport to such place of destination in India, including insurance and storage, as shall be specified in the Contract, shall be arranged by the Supplier, and related costs shall be included in the Contract Price.
- 14. Incidental Services
- 14.1 The Supplier shall provide such incidental services:-
- (a) The Supplier shall provide all necessary licenses and permissions for use of the Goods in India that may be required for the Goods. The cost shall be deemed to be included in the Contract Price.
- (b) The Supplier shall provide such other services as are stated in the Technical Specifications.

15. Warranty

15.1 All goods must be of fresh manufacture and must bear the dates of manufacture and expiry.

The Supplier further warrants that all Goods supplied under the Contract will have remaining a minimum of five-sixths (5/6) of the specified shelf life upon delivery at site or named place of destination in India for goods with a shelf life of more than two years and three-fourths (3/4) for goods with a shelf life of two years or less, have "overages" within the ranges set forth in the Technical Specifications, where applicable; are not subject to recall by the applicable regulatory authority due to unacceptable quality or an adverse drug reaction; and in every other respect will fully comply in all respects with the Technical Specifications and with the conditions laid down in the Contract.

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- 15.2 The Purchaser shall have the right to make claims under the above warranty up to the **full period of shelf life of goods**. Upon receipt of a written notice from the Purchaser, the Supplier shall, with all reasonable speed, replace the defective Goods without cost to the Purchaser. The Supplier will be entitled to remove, at his own risk and cost, the defective Goods once the replacement Goods have been delivered.
- 15.3 In the event of a dispute by the Supplier, a counter analysis will be carried out on the manufacturer's retained samples by an independent neutral laboratory agreed by both the Purchaser and the Supplier. If the counter analysis confirms the defect, the cost of such analysis will be borne by the Supplier as well as the replacement and disposal of the defective goods. In the event of the independent analysis confirming the quality of the product, the Purchaser will meet all costs for such analysis.
- 15.4 If, after being notified that the defect has been confirmed pursuant to GCC Sub-Clause 15.2 above, the Supplier fails to replace the defective Goods within the period of **30 days**, the Purchaser may proceed to take such remedial action as may be necessary, including removal and disposal, at the Supplier's risk and expense and without prejudice to any other rights that the Purchaser may have against the Supplier under the Contract. The Purchaser will also be entitled to claim for storage in respect of the defective Goods for the period following notification and shall have the right to deduct the sum from payments due to the Supplier under this Contract or any other contract.

15.5 Recalls

In the event any of the Goods are recalled, the Supplier shall notify the Purchaser within fourteen (14) days, providing full details of the reason for the recall and promptly replace, at its own cost, the items covered by the recall with Goods that fully meet the requirements of the Technical Specification and arrange for collection or destruction of any defective Goods. If the Supplier fails to fulfil its recall obligation promptly, the Purchaser will, at the Supplier's expense, carry out the recall.

16. Payment

- 16.1 The method and conditions of payments to be made to the supplier shall be paid upon under this contract shall be as follows:-
- (i) On Receipt: Ninety (90) percent of the Contract Price of the Goods delivered to the Consignee shall be paid within 60 days of submission of documents specified in GCC Clause 11 along with the Acknowledgement of receipt of Goods (Form 8 of the bid

document) through ECS of the bank.

- (ii) On Acceptance: Ten (10) percent of the Contract Price of Goods received shall be paid within sixty (60) days of acceptance of the Goods upon submission of an invoice (indicating the RITES Ltd., as the Purchaser on behalf of Ministry of Health & Family Welfare, Govt. of India; the Contract number, description of payment and total amount, signed in original, stamped or sealed with the company stamp/seal) supported by the Final Acceptance Certificate (Form 9 of the bid document) issued by the Consignee through ECS of the bank.
- 16.2 The Supplier's request (s) for payment shall be made to the Purchaser in writing, accompanied by an invoice describing, as appropriate, the Goods delivered and Services performed, and by documents submitted pursuant to GCC Clause 11 & 16.1, and upon fulfilment of other obligations stipulated in the Contract.

17. Prices

17.1 Prices charged by the Supplier for Goods delivered and Services performed under the Contract shall not vary from the prices quoted by the Supplier in its bid for the duration of the Contract. Prices shall be fixed and firm for the duration of the Contract. However, sales tax or Vat wherever payable shall be paid as applicable at the time of supply. Statutory variations are permitted during the original delivery schedule and not in the extended delivery schedule.

18. Change Orders

- 18.1 The Purchaser may at any time, by a written order given to the Supplier pursuant to GCC Clause 31, make changes within the general scope of the Contract in any one or more of the following:
- (a) specifications, where Goods to be furnished under the Contract are to be specifically manufactured for the Purchaser;
- (b) the method of shipment or packing;
- (c) the place of delivery; and/or
- (d) the Services to be provided by the Supplier.
- 18.2 If any such change causes an increase or decrease in the cost of, or the time required for, the Supplier's performance of any provisions under the Contract, an equitable adjustment shall be made in the Contract Price or delivery schedule, or both, and the Contract shall accordingly be amended. Any claims by the Supplier for adjustment under this clause must be asserted within thirty (30) days from the date of the Supplier's receipt of the Purchaser's

change order.

19. Contract Amendments

- 19.1 Subject to GCC Clause 18, no variation in or modification of the terms of the Contract shall be made except by written amendment signed/agreed by the Purchaser and Supplier.
- 20. Assignment
- 20.1 The Supplier shall not assign, in whole or in part, its obligations to perform under this Contract, except with the Purchaser's prior written consent. Assignment and sub-contracting, which is not disclosed in bid, are not permitted.

21. Delays in the Supplier's Performance

21.1 DELAYS IN THE SUPPLIES PERFORMANCE OF THE CONTRACT:

Delivery of the goods shall be made by the supplier in accordance with the time schedule specified in the contract. Any deviation in performance of its delivery obligations shall render the supplier liable to any or all of the following action.

- (a) Forfeiture of its Performance Security and / or
- (b) Imposition of liquidated damages and/or
- (c) Termination of the contract for default.
- 21.2 If at any time during the performance of the contract, the supplier should encounter conditions impending timely delivery of the goods, the supplier shall promptly notify the purchaser in writing of the facts of the delay, its likely duration and its cause(s). As soon as practicable after receipt of the suppliers notice, the purchaser shall evaluate the, situation and may at its discretion extend the supplier time for performance in which case the extension shall be ratified by the parties by amendment to the contract. The extension of the delivery period will be subject to the following conditions.
- a) The Purchaser shall deduct from the supplier under the provision of Clause 22 liquidated damages on the goods, which the supplier has failed to deliver within the delivery period fixed for delivery.
- b) That no increase in price on account of any statutory increases in or fresh imposition of customs duty, excise duty or sales tax or on account of any other tax or duty leviable in respect of the goods specified in the contract which takes place after the date of the delivery period stipulated in the contract, shall be admissible on such of the said goods as are delivered after the date of delivery stipulated in the contract.
- c) But nevertheless, the purchaser shall be entitled to the benefit of any decrease in price on account of reduction in or remission of

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Customs duty, Excise Duty, Sales Tax or on *account of* any other tax or duty or on any other grounds which takes place after the expiry of the date of delivery stipulated in the contract.

21.3 Except as provided under GCC Clause 24, a delay by the Supplier in the performance of its delivery obligations shall render the Supplier liable to the imposition of liquidated damages pursuant to GCC Clause 22, unless an extension of time is agreed upon pursuant to GCC Clause 21.2 without the application of liquidated damages.

22. Liquidated Damages

22.1 Subject to GCC Clause 24, if the Supplier fails to deliver any or all of the Goods or to perform the Services within the period(s) specified in the contract, the Purchaser shall, without prejudice to its other remedies under the Contract, deduct from the contract prices as liquidated damages, a sum equivalent to the 0.5 percent per week or part thereof of the delivered price of the delayed Goods or unperformed Services for each week or part thereof of delay until actual delivery or performance, up to a maximum deduction of the 10 percent of the value of delayed Goods. Once the maximum is reached, the Purchaser may consider termination of the contract pursuant to GCC Clause 23.

23. Termination for Default

- 23.1 The Purchaser, without prejudice to any other remedy for breach of Contract, by written notice of default sent to the Supplier, may terminate the Contract in whole or in part:
 - (a) if the Supplier fails to deliver any or all of the Goods within the period(s) specified in the contract, or within any extension thereof granted by the Purchaser pursuant to GCC Clause 21; or/and
 - (b) if the Goods do not meet the Technical Specifications stated in the Contract; or/and
 - (c) if the Supplier, in the judgment of the Purchaser, has engaged in corrupt or fraudulent or collusive or coercive practices in competing for or in executing the Contract.

For the purpose of this clause:

"corrupt practice" means the offering, giving, receiving, or soliciting of any thing of value to influence the action of a public official in the procurement process or in Contract execution.

"fraudulent practice" means a misrepresentation of facts in order to influence a procurement process or the execution of a Contract to the detriment of the Purchaser, and includes collusive practice among Bidders (prior to or after bid submission) designed to establish bid prices at artificial non-competitive levels and to

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deprive the Purchaser of the benefits of free and open competition.

- (d) if the Supplier fails to perform any other obligation(s) under the Contract.
- 23.2 In the event the Purchaser terminates the Contract in whole or in part, pursuant to GCC Clause 23.1, the Purchaser may procure, upon such terms and in such manner as it deems appropriate, Goods or Services similar to those undelivered, and the Supplier shall be liable to the Purchaser for any excess costs for such similar Goods or Services. However, the Supplier shall continue performance of the Contract to the extent not terminated.

24. Force Majeure

- 24.1 Notwithstanding the provisions of GCC Clauses 21, 22, and 23, the Supplier shall not be liable for forfeiture of its performance security, imposition of liquidated damages, or termination for default if and to the extent that its delay in performance or other failure to perform its obligations under the Contract is the result of an event of Force Majeure.
- 24.2 For purposes of this clause, "Force Majeure" means an event beyond the control of the Supplier and not involving the Supplier's fault or negligence and not foreseeable. Such events may include, but are not restricted to, acts of the Purchaser in its sovereign capacity, wars or revolutions, fires, floods, epidemics, quarantine restrictions, and freight embargoes.
- 24.3 If a Force Majeure situation arises, the Supplier shall promptly notify the Purchaser in writing of such condition and the cause thereof. Unless otherwise directed by the Purchaser in writing, the Supplier shall continue to perform its obligations under the Contract as far as is reasonably practical and shall seek all reasonable alternative means for performance not prevented by the Force Majeure event.

25. Termination for Insolvency

25.1 The Purchaser may at any time terminate the contract by giving written notice to the Supplier if the Supplier becomes bankrupt or otherwise insolvent. In this event, termination will be without compensation to the Supplier, provided that such termination will not prejudice or affect any right of action or remedy that has accrued or will accrue thereafter to the Purchaser.

26. Termination for Convenience

26.1 The Purchaser, by written notice sent to the Supplier, may terminate the contract, in whole or in part, at any time for its convenience. The notice of termination shall specify that termination is for the Purchaser's convenience, the extent to which performance of the Supplier under the Contract is terminated, and the date upon which

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such termination becomes effective.

- 26.2 The Goods that are complete and ready for shipment within thirty (30) days after the Supplier's receipt of notice of termination shall be accepted by the Purchaser at the Contract terms and prices. For the remaining Goods, the Purchaser may elect:
 - (a) to have any portion completed and delivered at the contract terms and prices; and/or
 - (b) to cancel the remainder and pay to the Supplier an agreed amount for partially completed Goods and Services and for materials and parts previously procured by the Supplier.

27. Settlement of Disputes

- 27.1 If any dispute or difference of any kind whatsoever shall arise between the Purchaser and the Supplier in connection with or arising out of the Contract, the parties shall make every effort to resolve amicably such dispute or difference by mutual consultation.
- 27.2 If, after thirty (30) days, the parties have failed to resolve their dispute or difference by such mutual consultation, then either the Purchaser or the Supplier may give notice to the other party of its intention to commence arbitration, as hereinafter provided, as to the matter in dispute, and no arbitration in respect of this matter may be commenced unless such notice is given.
- 27.2.1 Any dispute or difference in respect of which a notice of intention to commence arbitration has been given in accordance with this Clause shall be finally settled by arbitration. Arbitration may be commenced prior to or after delivery of the Goods under the Contract.
- 27.2.2 Arbitration proceedings shall be conducted in accordance with the rules of procedure which are as follows:-.
- (a) In case of Dispute or difference arising between the Purchaser and a domestic supplier relating to any matter arising out of or connected with this agreement, such disputes or difference shall be settled in accordance with the Arbitration and Conciliation Act, 1996. The arbitral tribunal shall consist of 3 arbitrators one each to be appointed by the Purchaser and the Supplier. The third Arbitrator shall be chosen by the two Arbitrators so appointed by the Parties and shall act as Presiding arbitrator. In case of failure of the two arbitrators appointed by the parties to reach upon a consensus within a period of 30 days from the appointment of the arbitrator appointed subsequently, the Presiding Arbitrator shall be

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appointed by the Medical Council of India.

- (b) The Arbitration and Conciliation Act of 1996 the rules herewith and any statutory modification or re-enactment thereof shall apply to arbitration proceedings
- (c) Where the value of the contract is Rs.10 million and below, the disputes or differences arising shall be referred to the Sole Arbitrator. The Sole Arbitrator should be appointed by agreement between the parties; failing such agreement, by the Medical Council of India.
- (d) If one of the parties fails to appoint its arbitrator in pursuance of sub-clause (a) above, within 30 days after receipt of the notice of the appointment of its arbitrator by the other party, then the Medical Council of India shall appoint the arbitrator. A certified copy of the order of the Medical Council of India making such an appointment shall be furnished to each of the parties.
- (e) The venue of Arbitration shall be the place from where the contract is issued and the language of the arbitration proceedings and that of all councils and communications between the parties shall be English.
- (f) The decision of the majority of arbitrators shall be final and binding upon parties. In case there is no majority decision, the decision of the Presiding arbitrator shall be final. The cost and expenses of Arbitration proceedings will be paid as determined by the arbitral tribunal. However, the expenses incurred by each party in connection with the preparation, presentation, etc. of its proceedings as also the fees and expenses paid to the Counsel appointed by such party or on its behalf shall be borne by each party itself.
- 27.3 Notwithstanding any reference to arbitration herein,
 - (a) the parties shall continue to perform their respective obligations under the Contract unless they otherwise agree; and
 - (b) the Purchaser shall pay the Supplier any monies due to the Supplier.

28. Limitation of Liability

- 28.1 Except in cases of criminal negligence or wilful misconduct, and in the case of infringement pursuant to Clause 7,
 - (a) the Supplier shall not be liable to the Purchaser, whether in contract, tort, or otherwise, for any indirect or consequential

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loss or damage, loss of use, loss of production, or loss of profits or interest costs, provided that this exclusion shall not apply to any obligation of the Supplier to pay liquidated damages to the Purchaser and

(b) the aggregate liability of the Supplier to the Purchaser, whether under the Contract, in tort or otherwise, shall not exceed the total price of contract, provided that this limitation shall not apply to the cost of repairing or replacing defective equipment.

- 29. Governing Language
- 29.1 The governing language of the contract shall be English. All correspondence and other documents pertaining to the Contract that are exchanged by the parties shall be written in the same language.
- 30. Applicable Law
- 30.1 The Contract shall be interpreted in accordance with the laws of Union of India.
- 31. Notices
- 31.1 Any notice given by one party to the other pursuant to this Contract shall be sent to the other party in writing or by cable, telex, or facsimile and confirmed in writing to the other party's address are as follows: -

The Purchaser's addresses for notice purposes is:

Group General Manager/MSM RITES Ltd., MSM Division, RITES Office Complex, Annex Building, 4th Floor, Plot No.144, Sector 44, Gurgaon-122003 (Haryana), India Fax: 91(124)2571659/2571660 Tel: 91(124) 2728-408/405/403

The Supplier's address for notice purposes is as mentioned in the NOA/contract.

- 31.2 A notice shall be effective when delivered or on the notice's effective date, whichever is later.
- 32. Taxes and Duties
- 32.1 The Supplier shall be entirely responsible for all taxes, duties, octroi, road permits, license fees, etc., incurred until delivery of the Goods to the Purchaser.
- 33. Jurisdiction

All disputes arising out of the contract shall (subject to clause 27) be subject to the jurisdiction of the appropriate court at New Delhi, India, only.

SECTION III. SCHEDULE OF REQUIREMENTS

SECTION III SCHEDULE OF REQUIREMENTS

Schedule	Description of Goods	Unit	Required	Bid Security
No.	-		Quantity	in Indian
				Rupees
I	Single Blood Bags (350ml) for West Zone	Nos.	302,000	300,000
II	Single Blood Bags (350ml) for North Zone	Nos.	263,700	260,000
III	Single Blood Bags (350ml) for South Zone	Nos.	247,300	240,000
IV	Single Blood Bags (350ml) for North East Zone	Nos.	40,500	40,000
V	Single Blood Bags (350ml) for East Zone	Nos.	180,400	180,000
	Total:		1,033,900	1,020,000
VI	Double Blood Bags (350ml) for West & North Zone	Nos.	128,520	240,000
VII	Double Blood Bags (350ml) for South, North East & East Zone	Nos.	105,840	210,000
	Total:		234,360	450,000
VIII	Double Blood Bags (450ml) for West & North Zone	Nos.	55,080	100,000
IX	Double Blood Bags (450ml) for South, North East & East Zone	Nos.	45,360	90,000
	Total:		100,440	190,000
X	Triple Blood Bags (350ml) for West & North Zone	Nos.	316,520	850,000
XI	Triple Blood Bags (350ml) for South, North East & East Zone	Nos.	260,280	720,000
	Total:		576,800	1,570,000
XII	Triple Blood Bags (450ml) for West & North Zone	Nos.	135,600	360,000
XIII	Triple Blood Bags (450ml) for South, North East & East Zone	Nos.	111,600	310,000
	Total:		247,200	670,000
XIV	Quadruple Blood Bags (350ml) for West & North Zone	Nos.	148,380	560,000
XV	Quadruple Blood Bags (350ml) for South, North East & East Zone	Nos.	121,980	460,000
	Total:		270,360	1,020,000
XVI	Quadruple Blood Bags (450ml) for West & North Zone	Nos.	63,630	240,000
XVII	Quadruple Blood Bags (450ml) for South, North East & East Zone	Nos.	52,260	200,000
	Total:		115,890	440,000
	Grand Total:			5,360,000

Delivery Schedule & Consignee details: (i) 50% quantity within 60 days after the issue of NOA, (ii) 25% within 61 to 90 days after the issue of NOA (iii) balance 25% within 91 to 120 days after the issue of NOA (Notification of Award).

Terms of Delivery: Final Destination at the consignee end (as per Schedule of Requirements)

Note: 1. The purchaser reserves the right to increase or decrease the ordered quantity of goods by **25%** during the currency of contract.

Consignee address and Consignee-wise Quantity distribution

S.	Consignee	SINGLE BLOOD BAGS (350ML)				
No		Quantity in No. of Bags				
		1 st	2 nd	3 rd	Total	
		Within 60 days	Within 61 to 90 days	Within 91 to 120		
		after the issue of	after the issue of	days after the		
		NOA	NOA	issue of NOA		
			Schedule-I			
1	Gujarat	36,500	18,200	18,200	72,900	
2	Ahmedabad	8,300	4,200	4,200	16,700	
3	Maharashtra	57,000	28,400	28,400	113,800	
4	Mumbai	17,200	8,500	8,500	34,200	
5	Rajasthan	30,400	15,200	15,100	60,700	
6	Dadra & Nagar Haveli	600	400	0	1,000	
7	Daman & Diu	400	200	0	600	
8	Goa	1,100	500	500	2,100	
		151,500	75,600	74,900	302,000	
			Schedule-II	2.100	12 = 00	
1	Chandigarh	6,400	3,200	3,100	12,700	
2	Chhattisgarh	5,700	2,900	2,800	11,400	
3	Delhi	27,800	14,000	14,000	55,800	
4	Haryana	9,500	4,700	4,800	19,000	
5	Himachal Pradesh	2,400	1,200	1,200	4,800	
6	Jammu & Kashmir	6,300	3,100	3,100	12,500	
7	Madhya Pradesh	19,000	9,500	9,400	37,900	
8	Punjab	15,200	7,600	7,600	30,400	
9	Uttar Pradesh	33,800	16,800	16,800	67,400	
10	Uttarakhand	5,900	3,000	2,900	11,800	
	Total	132,000	66,000	65,700	263,700	
	A 11 D 1 1	26.500	Schedule-III	10.200	72.000	
1	Andhra Pradesh	36,500	18,200	18,200	72,900	
2	Karnataka	27,900	13,900	13,900	55,700	
3	Kerala	21,300	10,600	10,600	42,500	
5	Pondicherry Tamil Nadu	2,100 29,200	1,000 14,600	1,000	4,100	
6		,	,	14,600 3,400	58,400	
0	Chennai	6,900	3,400 61,700	61,700	13,700 247,300	
		123,900	Schedule-IV	01,700	247,300	
1	Arunachal Pradesh	1,200	600	500	2,300	
2	Assam	11,000	5,500	5,500	22,000	
3	Manipur	2,100	1,100	1,000	4,200	
4	Meghalaya	700	300	400	1,400	
5	Mizoram	1,800	900	800	3,500	
6	Nagaland	800	400	300	1,500	
7	Sikkim	600	300	200	1,100	
8	Tripura	2,300	1,100	1,100	4,500	
- 0	inpura	20,500	10,200	9,800	4,500	
		40,500	Schedule-V	2,000	40,500	
1	Andaman & Nicobar	400	200	0	600	
		9,100	4,600	4,500	18,200	
		2.100				
2	Bihar	/	13 300	13 200	53 100	
2 3	Orissa	26,600	13,300	13,200	53,100	
2 3 4	Orissa West Bengal	26,600 45,600	22,800	22,700	91,100	
2 3	Orissa	26,600				

S.	Consignee	D	ouble Blood	Bags (350	ml)		Double Blood	Bags (450	ml)
No		Quan	tity in No. of	Bags	Grand		ntity in No. of	f Bags	Grand
		1 st	2 nd	3 rd	Total	1 st	2 nd	3 rd	Total
		Within	Within	Within		Within	Within	Within	
		60 days	61 to 90	91 to		60 days	61 to 90	91 to	
		after	days	120		after	days	120	
		the	after the	days		the	after the	days	
		issue of	issue of	after		issue of	issue of	after	
		NOA	NOA	the		NOA	NOA	the	
				issue of				issue of	
				NOA				NOA	
		0.000		ule-VI	4 - 7 - 0	2 100	_	ule-VIII	
1	Gujarat	8,280	4,140	4,140	16,560	3,600	1,740	1,740	7,080
2	Ahmedabad	1,860	960	960	3,780	840	420	360	1,620
3	Maharashtra	12,960	6,480	6,480	25,920	5,520	2,760	2,820	11,100
4	Mumbai	3,900	1,920	1,980	7,800	1,680	840	780	3,300
5	Rajasthan	6,900	3,420	3,480	13,800	3,000	1,500	1,440	5,940
6	Daman & Diu	240	0	0	240	120	0	0	120
7	Goa	480	0	0	480	180	0	0	180
8	Chandigarh	1,440	720	720	2,880	600	660	0	1,260
9	Chhatisgarh	1,260	660	660	2,580	1,140	0	0	1,140
10	Delhi	6,360	3,180	3,180	12,720	2,760	1,380	1,320	5,460
11	Haryana	2,160	1,080	1,080	4,320	900	480	480	1,860
12	Himachal Pd.	480	600	0	1,080	480	0	0	480
13	J & K	1,380	720	720	2,820	600	600	0	1,200
14	Madhya Pd.	4,320	2,160	2,160	8,640	1,800	960	960	3,720
15	Punjab	3,420	1,740	1,740	6,900	1,500	720	720	2,940
16	Uttar Pd.	7,620	3,840	3,840	15,300	3,240	1,620	1,680	6,540
17	Uttaranchal	1,320	660	720	2,700	1,140	0	0	1,140
	Total	64,380	32,280	31,860	128,520	29,100	13,680	12,300	55,080
				ule-VII	ı		Schedule-IX		T
1	Andhra Pradesh	8,280	4,140	4,140	16,560	3,540	1,740	1,800	7,080
2	Karnataka	6,300	3,180	3,180	12,660	2,640	1,380	1,380	5,400
3	Kerala	4,860	2,400	2,400	9,660	2,100	1,020	1,020	4,140
4	Pondicherry	480	480	0	960	420	0	0	420
5	Tamil Nadu	6,660	3,300	3,360	13,320	2,880	1,440	1,380	5,700
6	Chennai	1,560	780	780	3,120	660	360	300	1,320
7	Assam	2,520	1,260	1,260	5,040	1,080	540	540	2,160
8	Manipur	480	480	0	960	420	0	0	420
9	Meghalaya	300	0	0	300	120	0	0	120
10	Mizoram	420	360	0	780	360	0	0	360
11	Nagaland	360	0	0	360	180	0	0	180
12	Tripura	540	480	0	1,020	480	0	0	480
13	Andaman & Nicobar	180	0	0	180	60	0	0	60
14	Bihar	2,100	1,020	1,020	4,140	900	480	420	1,800
15	Orissa	6,120	3,000	3,000	12,120	2,580	1,320	1,260	5,160
16	West Bengal	10,380	5,160	5,160	20,700	4,440	2,220	2,220	8,880
17	Jharkhand	1,920	1,020	1,020	3,960	840	420	420	1,680
	Total	53,460	27,060	25,320	105,840	23,700	10,920	10,740	45,360
				Total:	234,360			Total:	100,440

S.	Consignee	7	Triple Blood	Bags (350 r	ml)			Triple Blood		nl)
No			tity in No. of	Bags	Grand		Quan	tity in No. of	Bags	Grand
		1 st	2 nd	3 rd	Total		1 st	2 nd	3 rd	Total
		Within	Within	Within			Within	Within	Within	
		60 days	61 to 90	91 to			60 days	61 to 90	91 to	
		after	days	120			after	days	120	
		the	after the	days			the	after the	days	
		issue of	issue of	after			issue of	issue of	after	
		NOA	NOA	the			NOA	NOA	the	
		1,012	1,012	issue of			1,012	1,012	issue of	
				NOA					NOA	
				lule-X					ule-XII	
1	Gujarat	20,400	10,200	10,200	40,800		8,760	4,360	4,360	17,480
2	Ahmedabad	4,720	2,320	2,320	9,360		2,000	1,000	1,000	4,000
3	Maharashtra	31,840	15,960	15,960	63,760		13,640	6,840	6,840	27,320
4	Mumbai	9,600	4,760	4,760	19,120		4,120	2,040	2,040	8,200
5	Rajasthan	17,040	8,480	8,480	34,000		7,280	3,640	3,640	14,560
6	Daman & Diu	560	0	0	560		240	0	0	240
7	Goa	560	600	0	1,160		480	0	0	480
8	Chandigarh	3,560	1,800	1,800	7,160		1,520	800	760	3,080
9	Chhatisgarh	3,200	1,600	1,600	6,400		1,360	680	680	2,720
10	Delhi	15,600	7,840	7,840	31,280		6,680	3,360	3,360	13,400
11	Haryana	5,320	2,680	2,640	10,640		2,280	1,160	1,120	4,560
12	Himachal Pd.	1,320	680	680	2,680		1,160	0	0	1,160
13	J & K	3,520	1,760	1,720	7,000		1,520	720	760	3,000
14	Madhya Pd.	10,640	5,320	5,280	21,240		4,560	2,280	2,280	9,120
15	Punjab	8,520	4,240	4,240	17,000		3,680	1,800	1,800	7,280
16	Uttar Pd.	18,840	9,440	9,440	37,720		8,080	4,040	4,040	16,160
17	Uttaranchal	3,280	1,680	1,680	6,640		1,400	720	720	2,840
	Total	158,520	79,360	78,640	316,520		68,760	33,440	33,400	135,600
				ule-XI					ıle-XIII	
1	Andhra Pradesh	20,400	10,200	10,200	40,800		8,720	4,400	4,360	17,480
2	Karnataka	15,600	7,800	7,760	31,160		6,680	3,360	3,320	13,360
3	Kerala	11,880	5,960	5,960	23,800		5,080	2,560	2,560	10,200
4	Pondicherry	1,160	560	560	2,280		560	440	0	1,000
5	Tamil Nadu	16,320	8,200	8,200	32,720	$\sqcup \downarrow$	7,000	3,520	3,520	14,040
6	Chennai	3,840	1,920	1,880	7,640	$\sqcup \!\! \perp$	1,640	840	800	3,280
7	Assam	6,160	3,080	3,080	12,320	$\sqcup \!\! \perp$	2,640	1,320	1,320	5,280
8	Manipur	1,160	600	600	2,360	$\sqcup \!\! \perp$	480	520	0	1,000
9	Meghalaya	760	0	0	760	$\sqcup \!\! \perp$	320	0	0	320
10	Mizoram	1,000	480	480	1,960	\sqcup	400	440	0	840
11	Nagaland	440	400	0	840	$\vdash \vdash$	360	0	0	360
12	Tripura	1,280	640	640	2,560	\sqcup	560	520	0	1,080
13	Andaman & Nicobar	360	0	0	360	\vdash	160	0	0	160
14	Bihar	5,080	2,560	2,560	10,200	$\vdash \vdash$	2,200	1,120	1,080	4,400
15	Orissa	14,880	7,440	7,440	29,760	-	6,400	3,200	3,160	12,760
16	West Bengal	25,520	12,760	12,720	51,000	\vdash	10,960	5,440	5,440	21,840
17	Jharkhand	4,880	2,440	2,440	9,760	\vdash	2,080	1,080	1,040	4,200
	Total	130,720	65,040	64,520	260,280	$\vdash \vdash$	56,240	28,760	26,600	111,600
				Total:	576,800				Total:	247,200

S.	Consignee	Qu	adruple Bloo	d Bags (35	0 ml)	Qı	adruple Bloo	od Bags (45	0 ml)
No			ntity in No. of		Grand	Quantity in No. of Bags			
		1 st	2 nd	3 rd	Total	1 st	2 nd	3 rd	
		Within	Within 61	Within		Within	Within 61	Within	
		60 days	to 90 days	91 to		60 days	to 90 days	91 to	
		after	after the	120		after	after the	120	
		the	issue of	days		the	issue of	days	
		issue of	NOA	after		issue of	NOA	after	
		NOA		the		NOA		the	
				issue of				issue of	Grand
				NOA				NOA	Total
		0.770	Schedu		10.110	4.000		ıle XVI	0.100
1	Gujarat	9,570	4,800	4,770	19,140	4,080	2,070	2,040	8,190
2	Ahmedabad	2,190	1,110	1,080	4,380	930	480	480	1,890
3	Maharashtra	14,940	7,470	7,470	29,880	6,420	3,210	3,180	12,810
4	Mumbai	4,470	2,250	2,250	8,970	1,920	960	960	3,840
5	Rajasthan	7,980	3,990	3,960	15,930	3,420	1,710	1,710	6,840
6	Daman & Diu	270	0	0	270	120	0	0	120
7	Goa	540	0	0	540	240	0	0	240
8	Chandigarh	1,680	840	840	3,360	720	360	360	1,440
9	Chhatisgarh	1,500	750	750	3,000	630	330	330	1,290
10	Delhi	7,350	3,660	3,660	14,670	3,180	1,560	1,560	6,300
11	Haryana	2,490	1,260	1,230	4,980	1,050	540	540	2,130
12	Himachal Pd.	660	300	300	1,260	540	0	0	540
13	J & K	1,650	810	810	3,270	720	360	330	1,410
14	Madhya Pd.	4,980	2,490	2,490	9,960	2,130	1,080	1,050	4,260
15	Punjab	3,990	1,980	1,980	7,950	1,740	840	840	3,420
16	Uttar Pd.	8,820	4,440	4,440	17,700	3,780	1,920	1,890	7,590
17	Uttaranchal	1,560	780	780	3,120	660	330	330	1,320
	Total	74,640	36,930	36,810	148,380	32,280	15,750	15,600	63,630
			Schedu	ıle XV		Schedule XVII			
1	Andhra Pradesh	9,570	4,770	4,770	19,110	4,110	2,040	2,040	8,190
2	Karnataka	7,320	3,660	3,630	14,610	3,150	1,560	1,560	6,270
3	Kerala	5,580	2,790	2,790	11,160	2,370	1,200	1,200	4,770
4	Pondicherry	540	270	270	1,080	450	0	0	450
5	Tamil Nadu	7,650	3,840	3,840	15,330	3,270	1,650	1,650	6,570
6	Chennai	1,800	900	900	3,600	780	390	360	1,530
7	Assam	2,910	1,440	1,440	5,790	1,260	630	600	2,490
8	Manipur	540	300	270	1,110	480	0	0	480
9	Meghalaya	360	0	0	360	150	0	0	150
10	Mizoram	480	420	0	900	390	0	0	390
11	Nagaland	390	0	0	390	180	0	0	180
12	Tripura	600	300	300	1,200	510	0	0	510
1.0	Andaman &		-		1.50				
13	Nicobar	150	0	0	150	60	0	0	60
14	Bihar	2,370	1,200	1,200	4,770	1,020	510	510	2,040
15	Orissa	6,990	3,480	3,480	13,950	3,000	1,500	1,470	5,970
16	West Bengal	11,970	5,970	5,970	23,910	5,160	2,550	2,550	10,260
17	Jharkhand	2,280	1,140	1,140	4,560	990	480	480	1,950
	Total	61,500	30,480	30,000	121,980	27,330	12,510	12,420	52,260
				Total:	270,360			Total:	115,890

The bidders shall inform the delivery schedule offered by them in the similar tables as above.

CONSIGNEE ADDRESSES

S. No	Name of the State	Address of the SACS	Email Id
1	Ahmedabad	Ahmedabad Municipal Corpn. AIDS Control Society, Old	ahmedabadmacs@gm
	MACS	Municipal Dispensary, behind Lal Bungalow, C.G. Road,	ail.com
		Ahmedabad.	
2	Andaman &	The Project Director,	andamansacs@gmail.
	Nicobar	Andaman & Nicobar AIDS Control Society,	com,
		G.B. Pant Hospital Complex, Port Blair-744104.	
3	Andhra Pradesh	Tele: 03192 - 236555, 237941 Andhra Pradesh State AIDS Control Society, Directorate of	sacsandhra@gmail.co
3	AndmaTraucsi	Medical and Health Services, Sultan Bazar Hyderabad -	m
		500059	
4	Arunachal Pradesh	The Project Director,	arunachalsacs@gmail
		Arunachal Pradesh State AIDS Control Society,	.com,
		Naharlagun, New Itanagar Arunachal Pradesh – 791110	
		Tele: 0360-2351016	
5	Assam	The Project Director,	assamsacs@gmail.co
		Assam State AIDS Control Society, Khanapara,	m,
		Guwahati – 781 022	
	7.0	Tele.: 0361 – 2360524, 2366388	1.1 0 1
6	Bihar	The Project Director,	biharsacs@gmail.co
		Bihar State AIDS Control Society, State Instt. Of Health & F W,	m,
		Sheikhpura, Patna-800014	
		Tele: 0612 – 2213383, 2290278, 2292494	
7	Chandigarh	The Project Director,	chandigarhsacs@gma
		Chandigarh State AIDS Control Society, Chandigarh	il.com,
		International Hostel, (Near PGIMER), Sector 15-A,	,
		Madhya Marg, Chandigarh- 160015	
8	Chennai MACS	Chennai Municipal Corpn. AIDS Control Society, 82 Thiru	chennaimacs@gmail.
		Vi-Ka-Salai, Mylapore, Chennai - 600003	com
9	Chhattisgarh	The Project Director,	chattisgarhsacs@gma
		Chattisgarh State Aids Control Society,	il.com,
		Chattisgarh Health Society (sub society aids), State health training centre, Kalibadi chowk, raipur, chattisgarh –	
		492001.	
		Tele: 0771-2235860, 2235240	
10	Dadra & Nagar	The Project Director,	
	Haveli	Dadra & Nagar Haveli State AIDS Control Society,	1.1
		1st Floor, Shri Vinoba Bhave Civil Hospital,	dnhsacs@gmail.com
		Silvassa – 396 230, Tele : 0260 - 2642061	
11	Daman & Diu	The Project Director,	ddsacs@gmail.com,
		State AIDS Control Society,	
		Union Territory of Daman & Diu, Primary Health Centre,	
12	D.II.!	Moti Daman, Daman – 396220	
12	Delhi	The Project Director, Delhi State AIDS Control Organization,	
		Dharmsala Block, Dr. Baba Saheb Ambedkar Hospital,	delhisacs@gmail.co
		Sector – 6, Rohini, Delhi - 110085,	m
		Tele: 011-27055722-24, 27055660, 27055725	
	l .	, , , , , , , , , , , , , , , , , , , ,	ı

S. No	Name of the State	Address of the SACS	Email Id
13	Goa	The Project Director,	
10		Goa State AIDS Control Society,	
		1st Floor, Dayanand Smruti Building, Swami Vivekanand	goaaids@gmail.com
		Road,	gountes o gillumito in
		Panaji, Goa – 403 001 Tele : 0832 – 2422519, 2427286	
14	Gujarat	The Project Director,	
	Gujurut	Gujarat State AIDS Control Society,	
		O-1 Block, New Mental Hospital Complex,	gsacs@icenet.net
		Menghaninagar, Ahmedabad - 380 016, Gujarat	godes e recinetimet
		Tele: 079 – 22681043, 22685210	
15	Haryana	The Project Director & DG,	haryanasacs@gmail.c
13		Haryana State AIDS Control Society	om,
		Plot No. C-15, Awas Bhawan, Sector-6, Panchkula, Haryana	OIII,
		1 lot 1vo. C-13, Awas Bhawan, Sector-0, I anchkula, Haryana	
		Tele: 0172-2563317, 2585413	
16	Himachal Pradesh	The Project Director,	
10		Himachal Pradesh State AIDS Control Society,	
		Hari Villa, Near Forest Rest House,	hpsacs@gmail.com
		Khalini, Shimla -2 0177-2625857,2621608	
17	Jammu & Kashmir	The Project Director,	jksacs@gmail.com,
1,	guinna & Ixasiinii	Jammu & Kashmir State AIDS Prevention & Control	jksues e gman.com,
		Society,	
		48, Samandar Bagh, Lal Chawk, Srinagar.	
		Tele: 0194-2477516,2486409, 2476642	
18	Jharkhand	The Project Director,	
10	Jimi ilimii	Jharkhand State AIDS Control Society,	
		Sardar Hospital Campus,	jharkhandsacs@gmai
		Puruliya Road, Ranchi –1, Jharkhand	1.com
		Tele: 0651-2309556, 2211018	
19	Karnataka	Karnataka State AIDS Control Society, No.4/13-1,	ksapspdp@gmail.co
		Crescent Road, High Grounds, Bangalore - 560001.	m
20	Kerala	The Project Director,	
		Kerala State AIDS Control Society,	
		IPP Building, Red Cross Road, Thiruvananathapuram -	keralasacs@gmail.co
		695037.	m
		Tele: 0471-2304882, 2327938,2305183	
21	Madhya Pradesh	The Project Director,	
		Madhya Pradesh State AIDS Control Society,	mpsacs@gmail.com/
		1 Arera Hills, 2 nd Floor, OILFED Building, Bhopal – 462	mpsacsb@sanchar.ne
		011	t.in
		Tele: 0755-2577016, 2559629,2577628 / 29	
22	Manipur	The Project Director,	
		Manipur State AIDS Control Society,	moninureosa@amail
		Medical Directorate, R & D Wing,	manipursacs@gmail.
		Lamphelpat, Imphal, Manipur -795 004.	com
		Tele: 0385-2410144	
23	Meghalaya	The Project Director,	
	"	Meghalaya State Aids Control Society,	meghalayasacs@gma
			il.com
		Ideal Lodge, Oakland, Shillong – 793001 Tele: 0364-2223140	il.com

S. No	Name of the State	Address of the SACS	Email Id
100		Mizoram State AIDS Control Society,	biz
		MV-124, Mission Veng South,	O12
		Aizawl – 796005. Mizoram	
		Tele: 0389-2321556/2321566	
25	MSACS, Mumbai	Project Director,	
		Maharashtra State AIDS Control Society	
		(MSACS), Ackworth Leprosy Hospital Compound, Behind	maharashtrasacs@gm
		S.I.W.S. College,	ail.com
		R. A. Kidwai Marg, Near Wadala Over Bridge,	
		Wadala (West), Mumbai – 400 031.	
		Tele: 022-24113097, 24115791,24115619	
26	Mumbai	Project Director,	
		Mumbai Districts AIDS Control Society,	1
		Municipal Corporation of Greater Mumbai	mumbaimacs@gmail.
		R.A. Kidwai Marg, Acworth Complex,	com
		Wadala (West), Mumbai – 400 031	
27	Nagaland	Tele: 022 – 24100250, 24100246-47 The Project Director,	
21	Nagaiand	Nagaland State AIDS Control Society,	
		Health & Family Welfare Department,	naglandsacs@gmail.c
		New Secretariat Building,	om
		Kohima – 797 001.	Om
		Tele: 0370-2241046,2241543	
28	Orissa	The Project Director,	
		Orissa State AIDS Cell,	
		2 nd Floor, Oil Orissa Building,	orissasacs@gmail.co
		Nayapalli, Bhubaneshwar – 751 012.	m
		Tele: 0674-2395134,2393235 / 415.	
29	Pondicherry	The Project Director,	
		Pondichery AIDS Control Society,	pondicherrysacs@gm
		No93, Perumal Koil Street (Up-Stairs),	ail.com
20		Pondicherry - 605001.Tele : 0413-234596,2337000	
30	Punjab	The Project Director,	
		Punjab State AIDS Control Society,	punjabsacs@gmail.co
		4th Floor, Prayaas Building, Sector-38 B, Chandigarh. Tele: 0172-2636795.	m
31	Rajasthan	The Project Director,	
31	Kajastiiaii	Rajasthan State AIDS Control Society,	
		Medical & Health Directorate,	rajasthansacs@gmail.
		Swasthya Bhawan, Tilak Marg, "C" Scheme, Jaipur-302	com
		005.	John The Control of t
		Tele: 0141-2225532, 2222452, 2221792	
32	Sikkim	The Project Director,	
		Sikkim State AIDS Control Society,	cildrimence@amail.co
		S.T.N.M. Hospital, Yangthang Building,	sikkimsacs@gmail.co
		Kazi Road, Gangtok, Sikkim -737 101	m
		Tele: 03592-205343,224481	
33	Tamil Nadu	The Project Director,	
		Tamil Nadu State AIDS Control Society,	tnsacs@tn.nic.in
		417 Pantheon Road, Egmore, Chennai – 600 008	and the second second
		Tele: 044-28190261, 28194917	

S.	Name of the State	Address of the SACS	Email Id
No			
34	Tripura	The Project Director,	
		Tripura State AIDS Control Society,	
		Health Directorate Building,	tripurasacs@gmail.co
		Pandit Nehru Complex, Gurkhabasti, 2 nd Floor,	m
		P.O. Kunjaban, Agartala, West Tripura – 799 006	
		Tele: 0381-2321614, 2221614	
35	Uttar Pradesh	The Project Director,	
		Uttar Pradesh State AIDS Control Society,	
		A -Block, 4 th Floor, P.IC.U.P. Bhawan,	upsacs@gmail.com
		Vibhuti Khand, Gomti Nagar, Lucknow – 226 010	
		Tele: 0522 – 2720360/61	
36	Uttarakhand	The Project Director,	
		Uttarakhand State AIDS Control Society,	
		Red Cross Bhawan, Near Directorate Medical Health,	uttranchalsacs@gmai
		Dandalakhound, Gujrada, (Opp, I.T. Park),	1.com
		Sahstradhara Road, DehradunTELE: 0135-27228144,	
		3107947	
37	West Bengal	The Project Director,	wbsacs@gmail.com,
		West Bengal State AIDS Prevention and Control Society,	
		Swasthya Bhawan, 1 st Floor, Wing –B,	
		GN-29, Sector-V, Salt Lake City, Kolkata – 700 091	
		Tele: 033-23574400, 23576000, Fax: 033-23570122.	

Appendix 'A

Schedule	Minimum value of completed contract	Similar Product
No.	(In Million Indian Rupees or	
	equivalent)	
I	13.0	Blood Bags
II	12.0	Blood Bags
III	11.0	Blood Bags
IV	2.0	Blood Bags
V	8.0	Blood Bags
VI	11.0	Blood Bags
VII	9.0	Blood Bags
VIII	5.0	Blood Bags
IX	4.0	Blood Bags
X	37.0	Blood Bags
XI	31.0	Blood Bags
XII	16.0	Blood Bags
XIII	14.0	Blood Bags
XIV	24.0	Blood Bags
XV	20.0	Blood Bags
XVI	11.0	Blood Bags
XVII	9.0	Blood Bags

Appendix 'B'

Schedule	Annual Turnover (in Million
No.	Indian Rupees or equivalent)
I	20.0
II	18.0
III	17.0
IV	3.0
V	12.0
VI	17.0
VII	14.0
VIII	8.0
IX	6.0
X	56.0
XI	47.0
XII	24.0
XIII	21.0
XIV	36.0
XV	30.0
XVI	17.0
XVII	14.0

SECTION IV. TECHNICAL SPECIFICATIONS

Section IV. Technical Specifications

INDEX FOR TECHNICAL SPECIFICATIONS

	Topic
Part A	Technical Specifications For SINGLE BLOOD BAGS (350ML), DOUBLE BLOOD BAGS 350ML & 450 ML, TRIPLE BLOOD BAGS 350ML & 450ML, QUADRUPLE BLOOD BAGS 350ML & 450ML
Part B	Technical Specifications - General
Part C	Special Instructions
Part D	Inspection & Tests
Part E	Bar coding requirements for all medical supplies

SECTION IV: TECHNICAL SPECIFICATIONS FOR SINGLE BLOOD BAGS (350ML), DOUBLE BLOOD BAGS 350ML & 450 ML, TRIPLE BLOOD BAGS 350ML & 450ML, QUADRUPLE BLOOD BAGS 350ML & 450ML

PART A

Bidders are required to mention "Comply"/ "Not comply" or specific information requested against each criteria of the following Technical Specification for the items being supplied.

General Specifications

Requirements	Your Offer (Please fill-in)
SINGLE BLOOD BAGS (350ML), DOUBLE BLOOD BAGS 350ML & 450 ML,	"Comply"/"Not
TRIPLE BLOOD BAGS 350ML & 450ML, QUADRUPLE BLOOD BAGS 350ML &	comply"
450ML General Specifications	
(a) Plastic Blood Bags should meet all the standards as laid down in ISO 3826, for the manufacturers have to produce documentary evidence from the laboratories approved by Government of India.	
(b) Bio-compatibility of the material of the plastics blood bags must be certified by the manufacturer and must be supplied by the test reports of the following:	
1. Cell culture cyto-toxicity	
2. Hemolysis	
3. Systemic infections (acute toxicity)	
4. Sensitization	
5. Intra-cutaneous injection (irritation)	
6. Pyrogen test	
7. Sterility	
(c) To assess quality of stored blood, manufacturer should provide documented evidence of following Bio-chemical parameters of blood stored in CPDA/CPDA-1/CPD-SAGM containing DEHP plasticized PVC blood bags manufactured by the company on $28^{th}/35^{th}/42^{nd}$ day of storage. The parameters are:	
1. Plasma pH	
2. ATP (% of initial volume)	
3. 2,3-DPG (% of initial volume)	
4. Plasma K+ (mEq/L)	
5. % of viable red cells (24 hours post transfusion)	
6. DEHP leaching (mg/100ml)	
7. DEHP should not be more than 0.01% w/v in the PVC	
(d) The plastic blood bag should have a shelf-life of minimum 2 years. Stability reports from a recognized laboratory must be produced	
(e) Slit present at the bottom of the bag should be "adequate to hang the blood bag during transfusion"	
(f) Packing size of good: Individual plastic blood bags should be packed in a plastic pack and such 5-10 bags should be packed in aluminum foil pack. The label of the aluminium foil pack should read as 'Aluminium foil pack once opened, the bags should be used within ten days. Ten such aluminium foiled packs should be packed in the corrugated boxes	

which should indicate clearly and legibly the name of the manufacturer, name of the product, batch number, quantity, date of manufacturing, date of expiry, gross and net weight and consignee's name and address and other particulars as required. It should also mention "To be stored at 20 to 25°C". There should be a temperature indicator on the carton to ensure cold chain maintenance.

(g) External sterility of the plastic blood bags should be ensured. The outer surface should be moisture free.

(h) Each carton should contain:

A copy of test reports.

A certificate mentioning "Blood may be collected up to the expiry date marked on the label which will be compatible with shelf life of components prepared as per required standards".

(i) Satisfactory Report from reputed Government users for last two years to be provided.

(j) At least five bags should be provided for the technical evaluation at the time of quotation.

Schedule – I to V: Single Blood Bags (350 ml)

Requirements	Your Offer (Please fill-in)
SINGLE BLOOD BAGS (350ml)	"Comply"/"Not
	comply"
Blood collection bag made up of DEHP (Di-2-ethylhexyl phthalate) plasticized	
PVC (Polyvinylchloride), collapsible non-vented sterile containers complete with	
collecting tube for completely closed system to avoid the chances of	
contamination.	
Capacity:	
Single blood bag – 350 ml	
Design and shapes:	
1. Flexible pre-sterilized	
2. Pyrogen free	
3. Non-toxic, non-haemolytic, biocompatible material	
4. No risk of contamination and air embolism (closed system) with all leak	
proof seals (Disposable Bags)	
5. Slit on the both sides of the bags should be enough to accommodate $5-10$	
ml volume test tubes	
6. The capacity of the bag should be enough to prevent any ballooning/rupture	
of the bag from the seam when it is filled up with the requisite volume of	
blood	
Tubing of bag:	
1. Flexible non-kinking	
2. Non-sticking	
3. Transparent	

-	Requirements	
	SINGLE BLOOD BAGS (350ml)	(Please fill-in) "Comply"/ "Not comply"
4.	Leak-proof	<u> </u>
5.	The tubing should have ID/Segment number as that on the bag	
6.	The tube should have multiple printed ID/Segment numbers. The numbers	
	should be legible and clear.	
7.	A clamp should be provided for closed system.	
Needle		
1.	16 gauge ultra thin walled and straight	
2.	Sharp regular margins and beveled tip	
3.	Rust proof	
4.	Tightly fixed with hub covered with sterile guard	
5.	Hermetically sealed	
	The needle should not separate from the tube at any point of time,	
	especially while removing it from the vein for donor safety.	
Exterr	nal Port:	
1.	Tamper proof and shouldn't be re-capped	
	Easily accessible	
Packa	•	
1.	Protective dual packaging (Individual & Aluminum) eliminating microbial	
	contamination on surface maintaining the contents of the bag	
2.	Easy to handle	
Antico	agulant and preservative solution:	
1.	CPDA-1 (49 ml i.e. 14 ml/100 ml of blood)	
	Clear & colourless	
3.	No discolouration on storage at room temperature	
4.	Manufacturer to supply anticoagulant quality check certificate	
Label:		
1.	Non-peal off	
	Heat sealed labels .	
	Remain attached between room temperature to 4°C with a transparent	
	adhesive	
4.	Date of manufacturing, date of expiry and lot number must be mentioned on	
	each bag	
5.	The expiry date should be at least 2 years from the date of supply of blood bags to the institute.	

<u>Schedule – VI to VII: Double Blood Bags (350 ml)</u> <u>Schedule – VIII to IX: Double Blood Bags (450 ml)</u>

Requir	ements	Your Offer
	D 11 DI 1D (250 1/9 450 1)	(Please fill-in)
	Double Blood Bags (350 ml/ & 450 ml)	"Comply"/
Blo	ood collection bag made up of DEHP (Di-2-ethylhexyl phthalate) plasticized PVC (Polyvinylchloride),	"Not comply"
	lapsible non-vented sterile containers complete with collecting tube for completely closed system to	
	oid the chances of contamination.	
Capacit	ty:	
	Double Blood bag	
•	Double Blood dag	
	Primary bag (350 ml/450 ml)	
	One Satellite bag (of 300 ml capacity)	
Design	and shapes:	
_		
7.	Flexible pre-sterilized	
8.	Pyrogen free	
9.	Non-toxic, non-haemolytic, biocompatible material	
10.	No risk of contamination and air embolism (closed system) with all leak proof seals (Disposable	
1.1	Bags)	
	Slits at both sides of the bags should be enough to accommodate 5 – 10 ml volume test tubes.	
12.	The capacity of the bag should be enough to prevent any ballooning/rupture of the bag from the seam	
	when it is filled up with the requisite volume of blood.	
Tubing	of bag:	
1.	Flexible non-kinking	
2.	Non-sticking State of the state	
3.	Transparent	
4.	Leak-proof	
5.	All the tubings should have same ID/Segment number as that on the bag	
6.	The tube should have multiple printed ID/Segment numbers. The numbers should be legible and	
	clear.	
7.	A clamp should be provided for closed system.	
Needle:		
1.	16 gauge ultra thin walled and straight	
2.	Sharp regular margins and beveled tip	
3.	Rust proof	
4.	Tightly fixed with hub covered with sterile guard	
5.	Hermetically sealed	
6.	The needle should not separate from the tube at any point of time, especially while	
0.	removing it from the vein for donor safety.	
	Temoving it from the vein for donor safety.	

Requirements		Your Offer (Please fill-in)
	Double Blood Bags (350 ml/ & 450 ml)	"Comply"/ "Not comply"
Extern	al Port:	
1.	Tamper proof and shouldn't be re-capped	
2.	Easily accessible	
Packag	e:	
1.	Protective dual packaging (Individual & Aluminum) eliminating microbial contamination on surface	
2.	maintaining the contents of the bag Easy to handle	
Anticoa	agulant and preservative solution:	
1.	CPDA-1 (14 ml/100 ml of blood) – Primary bag only	
2.	Clear & colourless	
3.	No discolouration on storage at room temperature	
4.	Manufacturer to supply anticoagulant quality check certificate	
Label:		
1.	Non-peal off	
2.	Heat sealed labels	
3.	Remain attached between room temperature to -80°C with a transparent adhesive	
4.	Date of manufacturing, date of expiry and lot number must be mentioned on each bag	
5.	The expiry date should be at least 2 years from the date of supply of blood bags to the	
	institute.	
Resista	nce to distortion:	
Filled to	o normal capacity	
•	Bag shall withstand an acceleration of 5000g for 30 min at temperature 4 ^o C to 24 ^o C without	
	becoming permanently distorted.	
•	Bag should be able to withstand temperature up to -80°C without breakage.	

<u>Schedule – X to XI: Triple Blood Bags (350 ml)</u> <u>Schedule – XII to XIII: Triple Blood Bags (450 ml)</u>

Requir	rements	Your Offer (Please fill-in)
	Triple Blood Bags (350 ml / 450 ml)	"Comply"/ "Not comply"
collaps	collection bag made up of DEHP (Di-2-ethylhexyl phthalate) plasticized PVC (Polyvinylchloride), ible non-vented sterile containers complete with collecting tube for completely closed system to avoid inces of contamination.	
Capaci	ity:	
•	Triple Blood bag	
	Primary bag - 350 ml / 450 ml First Satellite bag (of 300 ml capacity) Second Satellite bag (of 300 ml capacity) for platelet storage for 5 days	
Design	and shapes:	
1. 2. 3. 4. 5. 6.	Flexible pre-sterilized Pyrogen free Non-toxic, non-haemolytic, biocompatible material No risk of contamination and air embolism (closed system) with all leak proof seals (Disposable Bags) Slits at both sides of the bags should be enough to accommodate 5 – 10 ml volume test tubes The capacity of the bag should be enough to prevent any ballooning/rupture of the bag from the seam when it is filled up with the requisite volume of blood	
Tubing	g of bag:	
	 Flexible non-kinking Non-sticking Transparent Leak-proof All the tubings should have same ID/Segment number as that on the bag The tube should have multiple printed ID/Segment numbers. The numbers should be legible and clear. A clamp should be provided for closed system. 	
Needle	:	
1. 2. 3. 4. 5.	16 gauge ultra thin walled and straight Sharp regular margins and beveled tip Rust proof Tightly fixed with hub covered with sterile guard Hermetically sealed	

Triple Blood Bags (350 ml / 450 ml) 6. The needle should not separate from the tube at any point of time, especiall removing it from the vein for donor safety. External Port: 1. Tamper proof and shouldn't be re-capped 2. Easily accessible Package: 1. Protective dual packaging (Individual & Aluminum) eliminating microbial contamination or maintaining the contents of the bag. 2. Easy to handle Anticoagulant and preservative solution: 1. CPDA-1 (14 ml/100 ml of blood) – Primary bag only 2. Clear & colourless 3. No discolouration on storage at room temperature 4. Manufacturer to supply anticoagulant quality check certificate	"Comply"/ "Not comply" y while
removing it from the vein for donor safety. External Port: 1. Tamper proof and shouldn't be re-capped 2. Easily accessible Package: 1. Protective dual packaging (Individual & Aluminum) eliminating microbial contamination of maintaining the contents of the bag. 2. Easy to handle Anticoagulant and preservative solution: 1. CPDA-1 (14 ml/100 ml of blood) – Primary bag only 2. Clear & colourless 3. No discolouration on storage at room temperature	y while
External Port: 1. Tamper proof and shouldn't be re-capped 2. Easily accessible Package: 1. Protective dual packaging (Individual & Aluminum) eliminating microbial contamination of maintaining the contents of the bag. 2. Easy to handle Anticoagulant and preservative solution: 1. CPDA-1 (14 ml/100 ml of blood) – Primary bag only 2. Clear & colourless 3. No discolouration on storage at room temperature	
 Tamper proof and shouldn't be re-capped Easily accessible Package: Protective dual packaging (Individual & Aluminum) eliminating microbial contamination of maintaining the contents of the bag. Easy to handle Anticoagulant and preservative solution: CPDA-1 (14 ml/100 ml of blood) – Primary bag only Clear & colourless No discolouration on storage at room temperature 	
 Easily accessible Package: Protective dual packaging (Individual & Aluminum) eliminating microbial contamination of maintaining the contents of the bag. Easy to handle Anticoagulant and preservative solution: CPDA-1 (14 ml/100 ml of blood) – Primary bag only Clear & colourless No discolouration on storage at room temperature 	
 Easily accessible Package: Protective dual packaging (Individual & Aluminum) eliminating microbial contamination of maintaining the contents of the bag. Easy to handle Anticoagulant and preservative solution: CPDA-1 (14 ml/100 ml of blood) – Primary bag only Clear & colourless No discolouration on storage at room temperature 	
 Protective dual packaging (Individual & Aluminum) eliminating microbial contamination of maintaining the contents of the bag. Easy to handle Anticoagulant and preservative solution: CPDA-1 (14 ml/100 ml of blood) – Primary bag only Clear & colourless No discolouration on storage at room temperature 	
maintaining the contents of the bag. 2. Easy to handle Anticoagulant and preservative solution: 1. CPDA-1 (14 ml/100 ml of blood) – Primary bag only 2. Clear & colourless 3. No discolouration on storage at room temperature	
 Easy to handle Anticoagulant and preservative solution: CPDA-1 (14 ml/100 ml of blood) – Primary bag only Clear & colourless No discolouration on storage at room temperature 	n surface
Anticoagulant and preservative solution: 1. CPDA-1 (14 ml/100 ml of blood) – Primary bag only 2. Clear & colourless 3. No discolouration on storage at room temperature	
 CPDA-1 (14 ml/100 ml of blood) – Primary bag only Clear & colourless No discolouration on storage at room temperature 	
 Clear & colourless No discolouration on storage at room temperature 	
3. No discolouration on storage at room temperature	
E I	
4. Manufacturer to supply anticoagulant quality check certificate	
Label:	
1. Non-peal off	
2. Heat sealed labels	
3. Remain attached between room temperature to -80°C with a transparent adhesive	
4. Date of manufacturing, date of expiry and lot number must be mentioned on each bag	_
5. The expiry date should be at least 2 years from the date of supply of blood bags	to the
institute.	
Resistance to distortion:	
Filled to normal capacity	
• Bag shall withstand an acceleration of 5000g for 30 min at temperature 4°C to 24°C without	
becoming permanently distorted.	
• Bag should be able to withstand temperature up to -80°C without breakage.	

<u>Schedule – XIV & XV: Quadruple Blood Bags (350 ml)</u> <u>Schedule – XVI & XVII: Quadruple Blood Bags (450 ml)</u>

Requirements		Your Offer (Please fill-in)
	Quadruple Blood Bags (350 ml / 450 ml)	"Comply"/ "Not comply"
(Polyvinylchlo	on bag made up of DEHP (Di-2-ethylhexyl phthalate) plasticized PVC ride), collapsible non-vented sterile containers complete with collecting etely closed system to avoid the chances of contamination.	1 2
Capacity:		
Ovedmente blee	ad has	
Quadruple blo	-	
•	Primary bag – (350 ml/450ml) with top and top	
•	First satellite bag (of 100 ml capacity)- containing additive solution for 42 days red cell storage	
	•	
•	Second Satellite bag (of 300 ml capacity) for platelet storage for 5	
	days Third Satallita has (of 200 ml consoits)	
Dogian and ab	Third Satellite bag (of 300 ml capacity)	
Design and sh	apes:	
1.	Flexible pre-sterilized	
	Pyrogen free	
	Non-toxic, non-haemolytic, biocompatible material	
4.	· · · · · · · · · · · · · · · · · · ·	
	leak proof seals (Disposable Bags)	
5.	Slit on the both sides of the bags should be enough to accommodate 5	
	– 10 ml volume test tubes	
6.	The capacity of the bag should be enough to prevent any	
	ballooning/rupture of the bag from the seam when it is filled up with	
	the requisite volume of blood.	
Tubing of bag	:	
1	Flexible non-kinking	
2.		
3.	Transparent	
4.	Leak-proof	
5.	All the tubings should have ID/Segment number as on the bag	
6.	The tube should have multiple printed ID/Segment numbers. The	
	numbers should be legible and clear.	
7.	A clamp should be provided for closed system.	
Needle:		
1.	16 gauge ultra thin walled and straight	
2.	Sharp regular margins and beveled tip	
3.	Rust proof	
4.	Tightly fixed with hub covered with sterile guard	
5.	Hermetically sealed	
6.	The needle should not separate from the tube at any point of	

Requireme	nts	Your Offer (Please fill-in)
	Quadruple Blood Bags (350 ml / 450 ml)	"Comply"/ "Not comply"
	time, especially while removing it from the vein for donor safety.	
External P	Port:	
	 Tamper proof and shouldn't be re-capped Easily accessible 	
Package:		
	 Protective dual packaging (Individual & Aluminum) eliminating microbial contamination on surface maintaining the contents of the bag. Easy to handle 	
Anticoagu	lant and preservative solution:	
	 CPD (63ml i.e. 14 ml/100 ml of blood) – Primary bag only Additive solution (100 ml) - first satellite Bag. Clear & colourless No discolouration on storage at room temperature Manufacturer to supply anticoagulant quality check certificate 	
Label:		
2. 3.	Non-peal off Heat sealed labels Remain attached between room temperature to -80°C with a transparent adhesive Date of manufacturing, date of expiry and lot number must be mentioned on each bag The expiry date should be at least 2 years from the date of supply of blood	
	bags to the institute.	
Resistance	to distortion:	
• Ba	ormal capacity g shall withstand an acceleration of 5000g for 30 min at temperature 4°C to C without becoming permanently distorted. g should be able to withstand temperature up to -80°C without breakage.	

PART B

TECHNICAL SPECIFICATION – GENERAL

Our	Minimum Requirements	Your Offer
	TECHNICAL SPECIFICATION – GENERAL	(Please fill-in) "Comply"/
	TECHNICAL SPECIFICATION – GENERAL	"Not comply"
1.	Product and Package Specifications	1 vot compty
1.1.	The required packing standards and labeling must meet the requirements given in this Technical	
1.0	Specification and Part.	
1.2.	Not only the Goods but also the packaging components should also meet specifications suitable for use in a climate similar to that prevailing in the country of the Purchaser. All packaging must be proporty asseled and topposed proof.	
1.3.	be properly sealed and tampered-proof. All labeling and packaging inserts shall be in the language requested by the Purchaser or English if not otherwise stated	
1.4.	Goods requiring refrigeration or freezing for stability must specifically indicate storage	
1	requirements on labels and containers and be shipped in special containers to ensure stability in transit from point of shipment to port of entry.	
1.5.	Upon award, the successful Supplier shall, on demand, provide a translated version in the	
	language of the bid of the prescriber's information for any specific goods the Purchaser may	
	request	
2.	Product Information	
2.1. i	The following information will be required for each pharmaceutical product offered by the Bidder: International Non-Proprietary Name (INN), if applicable;	
i	i) Brand Name (if it appears on label);	
i	ii) Name and address of the manufacturer;	
i	v) Country of origin; and	
7	y)Compendia standards	
2.2.	Upon award, the supplier shall, on demand, provide a translated version in English, of the prescriber's information for any specific product, the Purchaser may request.	
2.3.	Failure to include any of this information, at the discretion of the Purchaser, may render the bid non-responsive.	
3.	Expiration Date	
3.1.	All products must indicate the dates of manufacture and expiry. In addition, unless otherwise stated in Part A of these Specifications, all products must arrive at the final consignee point with a remaining shelf life of at least five-sixths (5/6ths) of the total stipulated shelf life at the time of manufacture.	
4.	<u>Recalls</u>	
4.1.	If products must be recalled because of problems with product quality as a result of quality check carried out during the life span of the drug or adverse reactions to the pharmaceutical, the supplier will be obligated to notify the purchaser providing full details about the reason leading to the recall and shall take steps to replace the product in question at its own cost with a fresh batch of acceptable pharmaceuticals, or withdraw and give a full refund if the product has been take off the market due to safety problems.	
5.	Labeling Instructions	
5.1. 7	The label for each aluminum pack & each blood bags shall include: (a) the Purchaser's logo and code number and any specific color coding if required (b) content per pack (c) instructions for use	
	(c) instructions for use	

Our Minimum Requirements	Your Offer (Please fill-in)
TECHNICAL SPECIFICATION – GENERAL	"Comply"/ "Not comply"
(d) special storage requirements	
(e) batch number	
(f) date of manufacture and date of expiry (in clear language, not code)	
(g) name and address of manufacture with license number	
(h) any additional cautionary statement	
5.2. The outer case or carton should also display the above information	
6. Details of Packing/Cases	
6.1. All cases should prominently indicate the following:	
i) The generic name of the product;	
ii) date of manufacture and expiry (in clear language not code);	
iii) batch number; and	
iv) quantity per case.	
6.2. No case should contain drugs from more than one batch.	
7. <u>Unique Identifier</u>	
7.1. The Purchaser shall have the right to request the Supplier to imprint a logo on the containers used	
for packaging and in certain dosage forms such as tablets and this will be indicated in Part A of the	
Technical Specifications. The design of such logo shall be provided to the supplier at the time of	
Contract award.	
8. Qualifications of Manufacturer	
8.1. The bidder shall furnish a certificate from the competent FDRA that the manufacturer of the	
pharmaceutical or vaccine product covered by this Invitation for Bids is licensed to manufacture	
these products.	
9. Standards and Quality Assurance Requirements	
9.1. All products must:	
(a) Meet the requirements of manufacturing legislation and regulation of pharmaceuticals or	
vaccines in the country of origin;	
(b) Conform to all the specifications contained herein; and	
(c) Must undergo strict raw material inspection, in process checks, appropriate material handling to eliminate cross contamination (of molecules) and final product testing to ensure quality and consistency of the products.	
9.2. The Bidder is required to furnish to the Purchaser:	
(a) With each consignment, a certificate of quality assurance test results concerning quantitative assay, chemical	
analysis and other tests, as applicable to the product being supplied and Part A of these Specifications.	
(b) Assay methodology of any or all tests if requested.	
(c) Evidence of basis for expiration dating and other stability data on the offered package (as per climatic	
conditions prevalent in India) concerning the commercial final package upon request.	
(d) Package integrity test results.	
9.3. The Bidder will also be required to provide the purchaser with access to its manufacturing facilities to inspect its	
facilities, quality control procedures for raw materials, test methods, in-process tests, and finished Goods.	

THE PRODUCTS OFFERED ARE IN ACCORDANCE WITH THE SPECIFICATIONS AND REQUIREMENTS YES NO

ANY DEVIATION MUST BE LIST	

PART C

SPECIAL INSTRUCTIONS

Requirement									
SPECIAL INSTRUCTIONS									
1.	Each blood bag packing, inner carton and nested cartons to have the following words printed in red ink with bold letters.								
	"NACO SUPPLIES- NOT FOR SALE"								
2.	Life of the product, indicating the date of manufacture and date of expiry should be printed as per Drugs & Cosmetics Act-India								
3.	Equivalency of Standards & Codes								
	Wherever reference is made in the Technical Specifications to specific standards and codes to be met by the Product to be furnished or tested, the provisions of the latest current edition or revision of the relevant standards or codes in effect shall apply, unless otherwise expressly stated in the Contract. Where such standards and codes are national or authoritative standards that ensure substantial equivalence to the standards and codes specified will be acceptable								
4.	Packing (Clause 10 of GCC) Add as clause 10.3 of the GCC the following –								
	Packing Instruction: Each Aluminum pack will be marked on one side with proper paint/indelible ink, the following;								
i)Projec	t :Third National HIV/AIDS Control Project								
ii)RITE	S LTD. Purchase Order No. :								
	atry of origin of Goods :								
	lier's Name and :								
	ng list reference number :								
5.	Each outer packing containing the unit packing should have the following label printed in bold letters in large size.								
i)	Purchaser's Name : MINISTRY OF HEALTH & FAMILY WELFARE, Govt. of India, through RITES LTD.								
ii)	Project :Third National HIV/ AIDS Control Project								
,	iii) RITES LTD. Purchase Order No:								
iv) v)	Country of origin of Goods Supplier's Name								
6.	Any other labeling requirement which the purchaser may ask at the time of approving the labeling samples								

PART D

Inspection and Tests

The following inspection procedures and tests are required by the Purchaser

Our Minimum Requirements	Your Offer (Please fill-in)
Inspection and Tests	"Comply"/ "Not comply"
a. Two sets of samples each of 40 bags will be drawn at random from each batch by the Purchaser's Inspector at the manufacturer's premises & sealed before dispatch.	
b. One set of sealed sample will be sent to an independent laboratory selected by the Inspector for conducting the required test to confirm whether the samples conform to the prescribed specification. Another set of sealed sample will be retained with the testing lab as counter sample till the shelf life.	
c. A note will be issued by the Purchaser on the basis of test report, accepting or rejecting the batch as the case may be	
d. The Goods will be dispatched only after the above inspection and test procedure has been followed and dispatch clearance has been issued.	
e. The Purchaser/consignee shall have the right to draw samples at random from the consignment anytime during the shelf life of the Goods and get them retested to satisfy whether the lots conform to the laid down specifications. In the event of the product failing to conform to specifications, the consignee shall reject that batch of supply and inform the supplier for arranging replacement of the rejected batches at supplier's cost.	

PART E

Bar coding requirements for all medical supplies

Our Requirements	Your Offer
Bar coding requirements for all medical supplies	(Please fill-in) "Comply"/"Not comply"
Section A) Primary packaging (Item level and monocarton level)	T
At individual item level (strip of 10 tablets, syrup bottle, injections, vials etc) and/ or on its monocarton (wherever applicable), are required to have a pre printed barcode on its product packaging using either of the barcode symbologies mentioned below:	
a) GS1 linear barcode symbology (EAN-13/UPC-A/EAN-8) to encode GTIN (Global Trade Identification Number) within the barcode.	
b) GSI Data Matrix symbology to encode 14 digits product code (GTIN14) within the barcode and using (01) application identifier (to be used where ptinting space is extremely limited).	
Examples of the same are reproduced at Annexure 'A'.	
All other human readable information on product packaging shall be as required under existing Regulatory labeling & marking requirements.	
Section B) Secondary level Packaging (Intermediate packaging) At secondary level packaging (e.g. box of 10 strips containing 10 tabs each, pack of 10 vials, pack of 10 injections etc), barcode encoding following information to be stickered or preprinted on secondary packaging:	
 Product identification Code (GTIN-14 of secondary pack) using application identifier (01). Expiry date in YYMMDD format using application identifier (17) Batch/Lot Number using application identifier (10) 	
GSI-128 barcode symbology to be used to generate the barcode.	
Examples of the same are reproduced at Annexure 'B'.	
All other human readable information on product packaging shall be as required under existing Regulatory labeling & marking requirements.	
Section C) Tertiary level packaging (Shipper level packaging)	
At shipper level packaging, a single label containing two barcodes needs to be generated and stickered. The barcodes will encode following information:	

Our Requirements	Your Offer (Please fill-in)
Bar coding requirements for all medical supplies	"Comply"/ "Not comply"
The first barcode will contain the following information:	
 Product Identification Code (GTIN-14 of shipper level pack) using application identifier (01). Expiry Date in YYMMDD format using application identifier (17) Batch/Lot Number using application identifier (10) 	
The second barcode will contain the following information: 1) SSCC (Serial Shipping Container Code) using application identifier (00)	
Examples of the same are reproduced at annexure 'c'.	
All other human readable information on product packaging shall be as required under existing Regulatory labeling & marking requirements.	

Annexure "A"

Examples of Primary Level Packaging

For generation of GSI barcode at primary level packaging either of the mentioned symbologies can be used, following GSI General Specifications.

The following GSI barcode symbologies are available as options:-

1) The barcode sample for EAN-13 barcode symbology encoding GTIN-13



2) The barcode sample for UPC-A barcode symbology encoding GTIN-12



Note: Both GTIN-13 GTIN-12 are in extensive use worldwide

3) The barcode sample for EAN-8 barcode symbology encoding GTIN-8 (Used where printing space is a constraint)



4) The barcode sample for GSI Data Matrix barcode symbology encoding GTIN-14 (Used where printing space is extremely limited)



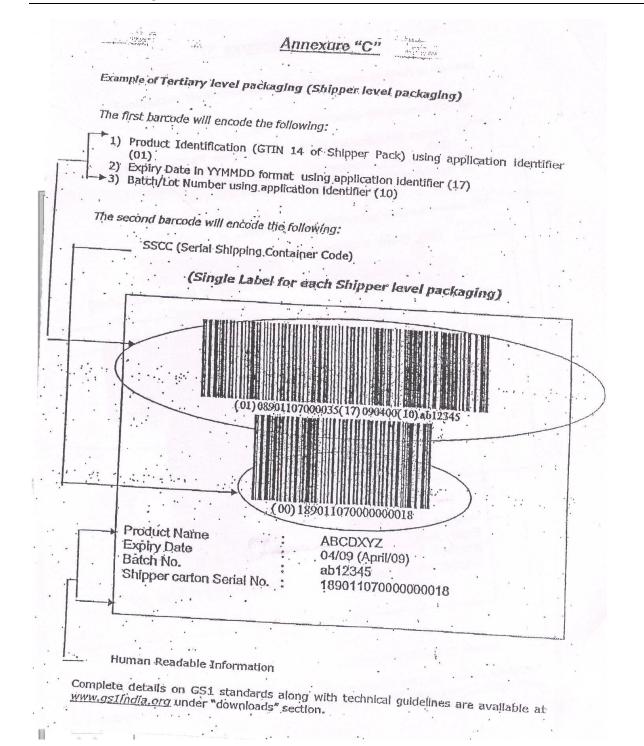
Annexure "B"

Example of Secondary level Packaging

The barcode will encode:

- 1) Product identification (GTIN 14 of secondary pack) using application identifier (01)
- 2) Expiry date in **YYMMDD** format using application identifier (17)
- 3) Batch/Lot Number using application identifier (10)





SECTION V. SAMPLE FORMS

SAMPLE FORMS

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1. Bid Form

	Date:	[insert: date of bid]
[Purchase	r specify: "IFB No.:	[number]"]
		[insert: name of Contract]
To: [Purchaser insert: Name and addre	ss of Purchaser]	
Dear Sir or Madam:		
which is hereby acknowledged, we, the unContract in full conformity with the said figures](Bid Price") or such other sums as may be	ndersigned, offer to su Bidding Documents fo) [inse e determined in accord	Addenda Nos. [insert numbers], the receipt of pply and deliver the Goods under the above-named or the sum of Rs[insert: amount in vert: amount in words] (hereinafter called "the Total ance with the terms and conditions of the Contract. ttached herewith and are made part of this bid.
We undertake, if our bid is accepted in the schedule of requirements.	cepted, to deliver the	Goods in accordance with the delivery schedule
If our bid is accepted, we under within the times specified in the Bidding I		formance security in the form, in the amounts, and
We agree to abide by this bid, for remain binding upon us and may be accept		iod specified in Clause 17.1 of the ITB and it shall before the expiration of that period.
	n of award, shall consti	ed between us, this bid, together with your written tute a binding Contract between us. We understand eceive.
		is made to us, in executing) the above contract, we ree in India namely "Prevention of Corruption Act
Commissions or gratuities, if an execution if we are awarded the Contract,		by us to agents relating to this bid, and to contract
Name and Address of Agent	Amount in Indian Rupees	Purpose of Commission or Gratuity
(if none, state "none")		
We confirm that we comply with the eligib	bility requirements as p	per ITB clause 4 of the bidding documents.
We understand that you are not bound to a	accept the lowest or any	y bid you may receive.
Dated this [insert: number] day of [insert:	ert: month],[insert:	vear].
Signed:		
In the capacity of [insert: title or position	<i>i</i>]	
Duly authorized to sign this bid for and or	behalf of [insert: na	me of Bidder]

Section V: Sample Forms 87

2a. Price Schedule for indigenous items

1	2	3	4		5					6	7	8	9
Schedule No	Product	Unit pack size	Quantity offered		Price for each unit					Total Unit price	Total price	Name of manufacturer	Country of origin
				Ex-factory Ex- warehouse Ex- showroom Off-the- shelf	Excis	e duty, if any In INR		ax/ VAT if any In INR	Other charges including inland transportatio n, incidental charges etc.				
				(a)		(b)		(c)	(d)	(a+b+c+d)	4 x 6		

Note:

(a) In case of discrepancy between unit price and total price, the unit price shall prevail.

(b) In case of discrepancy between price quoted in figure and words, price in words, shall prevail.

(c) "We hereby declare that in quoting the above price, we have taken into account the entire credit on inputs available under the CENVAT CREDIT scheme & VAT. Total Bid Price:

Currency:

In figures: In words:

Signed: _____
Dated:

In the capacity of: [insert: title or other appropriate designation]

2b. Price Schedule for imported items

1	2	3	4		5		6	7	8	9
Schedul e No	Product	Unit pack size	Quantity offered	Pri	Price for each unit		Total Unit price	Total price	Name of manufacturer	Country of origin
				CIF (Indian port) (in INR)	Custom duty (in INR)	Other charges including inland transportatio n etc. (in INR)				
				(a)	(b)	(c)	(a+b+c)	4 x 6		

T. T	- 4	

(a) (b)	In case of discrepancy between unit price and total price, the unit price shall prevail. In case of discrepancy between price quoted in figure and words, price in words, shall prevail.	Total Bid Price: Currency: In figures: In words:
	Signed: _	
	Datadi	

In the capacity of: [insert: title or other appropriate designation]

2c. Price Schedule for already imported items

1	2	3	4		5			6	7	8	9
Schedul e No	110000	Unit pack size	Quantity offered	P	rice for ea	ch unit		Total Unit price	Total price	Name of manufacturer	Country of origin
				Unit price including Custom/import Duties (in INR)		ax/ VAT any	Other charges including inland transportatio n etc. (in INR)				
				(a)		(b)	(c)	(a+b+c)	4 x 6		
	_										

a)	In case of discrepanc			

- (b) In case of discrepancy between price quoted in figure and words, price in words, shall
- (c) prevail.

The supplier should provide the details of custom duty already paid/payable separately

10	otai	Bia	Price:
Cι	ırre	ncy:	

In figures: In words:

Signed:		
Dated:		

In the capacity of: [insert: title or other appropriate designation]

3. Bid Security Form

Date: [insert: date]

IFB: [insert: name and number of IFB]
Contract: [insert: name and number of Contract]

To: [insert: name and address of Purchaser]

WHEREAS [insert: name of Bidder] (hereinafter called "the Bidder") has submitted its bid dated [insert: date of bid] for the performance of the above-named Contract (hereinafter called "the Bid")

KNOW ALL PERSONS by these present that WE [insert: name of bank] of [insert: address of bank] (hereinafter called "the Bank") are bound unto [insert: name of Purchaser] (hereinafter called "the Purchaser") in the sum of: [insert: amount], 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 [insert: number] day of [insert: month], [insert: year].

THE CONDITIONS of this obligation are the following:

- 1. If, after the bid submission deadline, the Bidder
 - (a) withdraws its bid during the period of bid validity specified by the Bidder in the Bid Form, or
 - (b) does not accept the Purchaser's corrections of arithmetic errors in accordance with the Instructions to Bidders; or
- 2. If the Bidder, having been notified of the acceptance of its bid by the Purchaser during the period of bid validity
 - (a) fails or refuses to sign the Contract Agreement when required; or
 - (b) fails or refuses to issue the performance security in accordance with the Instructions to Bidders.
 - (c) In case of any false, incorrect or misleading information provided in the bid.

We undertake to pay to 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 any one of the two above-named CONDITIONS, and specifying the occurred condition or conditions.

This guarantee will remain in full force up to and including [insert: the date that is 45 days after the period of bid validity], and any demand in respect thereof must reach the Bank not later than the above date.

Signed:	
Date:	
in the capacity of: [insert:	title or other appropriate designation

For and on behalf of the Bank

Common Seal of the Bank

4. Form of Contract Agreement

THIS CONTRACT AGREEMENT is made

the [insert: number] day of [insert: month], [insert: year].

BETWEEN

- [insert: Name of Purchaser], a [insert: description of type of legal entity, for example, an agency of the Ministry of of the Government of [insert: country of Purchaser], or corporation incorporated under the laws of [insert: country of Purchaser]] and having its principal place of business at [insert: address of Purchaser] (hereinafter called "the Purchaser"), and
- [insert: name of Supplier], a corporation incorporated under the laws of [insert: country of Supplier] and having its principal place of business at [insert: address of Supplier] (hereinafter called "the Supplier").

WHEREAS the Purchaser invited bids for certain goods and ancillary services, viz., [insert: brief description of goods and services] and has accepted a bid by the Supplier for the supply of those goods and services at a unit rate of [insert: contract price in words and figures] (hereinafter called "the Contract Price") during the period of contract i.e. _______

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.
- 2. The following documents shall constitute the Contract between the Purchaser and the Supplier, and each shall be read and construed as an integral part of the Contract:
 - (a) This Contract Agreement
 - (b) Instruction to bidder
 - (c) General Conditions of Contract
 - (d) Technical Requirements (including Technical Specifications, Functional Requirements and Implementation Schedule)
 - (e) The Supplier's bid and original Price Schedules
 - (f) The Schedule of Requirements
 - (g) The Purchaser's Notification of Award
 - (h) [Add here: any other documents]
- 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. **BRIEF DESCRIPTION** UNIT TOTAL **DELIVERY** NO. OF PHARMACEUTICALS **PRICE TERMS** PRICE & VACCINES **TOTAL VALUE: Delivery Schedule:** For and on behalf of the Purchaser in the capacity of [insert: title or other appropriate designation] Signed: in the presence of _____ For and on behalf of the Supplier Signed: in the capacity of [insert: title or other appropriate designation] in the presence of _____ CONTRACT AGREEMENT dated the [insert: number] day of [insert: month], [insert: year] **BETWEEN** [insert: name of Purchaser], "the Purchaser" and [insert: name of Supplier], "the Supplier"

5. Performance Security Bank Guarantee

(unconditional)

Date: [insert: date]

IFB: [insert: name or number of IFB]
Contract: [insert: name or number of Contract]

To: [insert: name and address of Purchaser]

Dear Sir or Madam:

We refer to the Contract Agreement ("the Contract") signed on [insert: date] between you and [insert: name of Supplier] ("the Supplier") concerning the supply and delivery of [insert: a brief description of the Goods]. By this letter we, the undersigned, [insert: name of bank], a bank (or company) organized under the laws of [insert: country of bank] and having its registered/principal office at [insert: address of bank], (hereinafter, "the Bank") do hereby jointly and severally with the Supplier irrevocably guarantee payment owed to you by the Supplier, pursuant to the Contract, up to the sum of [insert: amount in numbers and words]. This guarantee shall be reduced or expire as provided for by GCC Sub-Clause 8.4.

We undertake to make payment under this Letter of Guarantee upon receipt by us of your first written demand signed by your duly authorized officer declaring the Supplier to be in default under the Contract and without cavil or argument any sum or sums within the above-named limits, without your need to prove or show grounds or reasons for your demand and without the right of the Supplier to dispute or question such demand. Our liability under this Letter of Guarantee shall be to pay to you whichever is the lesser of the sum so requested or the amount then guaranteed under this Letter in respect of any demand duly made under this Letter prior to expiry of this Letter of Guarantee, without being entitled to inquire whether or not this payment is lawfully demanded.

This Letter of Guarantee shall be valid from the date of issue until the date of expiration of the guarantee, as governed by the Contract. Except for the documents herein specified, no other documents or other action shall be required, notwithstanding any applicable law or regulation. Our liability under this Letter of Guarantee shall become null and void immediately upon its expiry, whether it is returned or not, and no claim may be made under this Letter after such expiry or after the aggregate of the sums paid by us to you shall equal the sums guaranteed under this Letter, whichever is the earlier. All notices to be given under this Letter shall be given by registered (airmail) post to the addressee at the address herein set out or as otherwise advised by and between the parties hereto.

This guarantee shall expire no later than the ____ day of _____, 2____, and any demand for payment under it must be received by us at this office on or before that date.

We hereby agree that any part of the Contract may be amended, renewed, extended, modified, compromised, released, or discharged by mutual agreement between you and the Supplier, and this security may be exchanged or surrendered without in any way impairing or affecting our liabilities hereunder without notice to us and without the necessity for any additional endorsement, consent, or guarantee by us, provided, however, that the sum guaranteed shall not be increased or decreased.

No action, event, or condition that by any applicable law should operate to discharge us from liability hereunder shall have any effect, and we hereby waive any right we may have to apply such law, so that in all respects our liability hereunder shall be irrevocable and, except as stated herein, unconditional in all respects.

For and on behalf of the Bank	
Signed:	_
Date:	_
in the capacity of: [insert: title of	or other appropriate designation
Common Seal of the Bank	

Section V: Sample Forms 94

6. Proforma for Performance Statement (for a period of last five years)

		Bid No	Date	of opening	Time _	Но	ours
Name of t	he Firm						
Order placed by	Order No. and Date	Description and	Value of order		completion elivery	Remarks indicating	Was the supply of pharmaceuticals/
(full address of Purchaser)		quantity of ordered goods		As per contract	Actual	reasons for late delivery, if any	Consumables satisfactory*
1	2	3	4	5	6	7	8

Signature and seal of the Bidder	
C	

Countersigned by seal of Charted Accountant_____

^{*} The Bidder shall also furnish the following documents in connection with their past performance:

Section V: Sample Forms 95

For supplies within India & for Exports

- a. For supplies made to public sector units in India, an Affidavit confirming that the performance statement given is correct.
- b. However in case of supplies to private sector units, an affidavit confirming that the performance statement is correct along with following supporting evidence.
 - i. Copy of Purchase Orders
 - ii. Copy of Invoices
 - iii. Proof of Payment received from Purchasers
 - iv. Documentary evidence (Client's certificate) in support of satisfactory completion of contract

7. Manufacturer's Authorization

[The Bidder shall require the Manufacturer to fill in this Form in accordance with the instructions indicated. This letter of authorization should be on the letterhead of the Manufacturer and should be signed by a person with the proper authority to sign documents that are legally binding on the Manufacturer. The Bidder shall include it in its bid, if so indicated in the ITB.]

Date: [insert date (as day, month and year) of Bid Submission]

IFB No.: [insert number of bidding process]

Alternative No.: [insert identification No if this is a Bid for an alternative]

To: [insert complete name of Purchaser]

WHEREAS

We [insert complete name of Manufacturer], who are official manufacturers of [insert type of goods manufactured], having factories at [insert full address of Manufacturer's factories], do hereby authorize [insert complete name of Bidder] to submit a bid the purpose of which is to provide the following Goods, manufactured by us [insert name and or brief description of the Goods], and to subsequently negotiate and sign the Contract against the above IFB.

We hereby extend our full guarantee and warranty in accordance with Clause 15 of the General Conditions of Contract, with respect to the Goods offered by the above firm against this IFB.

No company or firm or individual other than M/s. ________ are authorized to bid, and conclude the contract for the above goods manufactured by us against this specific IFB.

Signed: [insert signature(s) of authorized representative(s) of the Manufacturer]

Name: [insert complete name(s) of authorized representative(s) of the Manufacturer]

Title: [insert title]

Duly authorized to sign this Authorization on behalf of: [insert complete name of Bidder]

Dated on _______ day of _______, ____ [insert date of signing]

Note – Modify this format suitably in cases where manufacturer's warranty and guarantee are not applicable for the items for which bids are invited.

8. Acknowledgement of Receipt of Goods (for 90% Payment)

(This certificate is to be issued to RITES and copy to Supplier and NACO. All the three copies "should be signed in ORIGINAL".)

To

MSM Division, RITES Ltd., RITES Office Complex, Annex Building, 4th Floor,

Plot No.144, Sector 44, Gurgaon - 122003, Haryana.

Fax: 91(124)2571659/2571660, Tel: 91(124) 2728-408/405/403

Email: rites_naco@rediffmail.com, rites_naco@rites.com

This is to certify that the Goods as detailed below have been received duly inspected in good condition in accordance with the conditions of the contract and amendment if any.

Purchaser :RITES Ltd., Gurgaon, Haryana on behalf MoH&FW (NACO) Contract i.e. NOA No. & Date : Description of Goods (Schedule No.) : Delivery Lot No. : Overtity supplied in Numbers	of
Contract i.e. NOA No. & Date : Description of Goods (Schedule No.) : Delivery Lot No. :	
Description of Goods (Schedule No.) : Delivery Lot No. :	
Delivery Lot No. :	
•	
Quantity supplied in Numbers : Quantity supplied in Words :	
- V 11	
Name of Supplier :	
Batch No(s).	
Manufacturing Date(s) :	
Expiry Date(s) :	
Invoice No. and Date :	
Date of delivery at Consignee :	
destination site	
Outstanding/dues with the supplier as :	
per NOA & amendment, if any	
Consignee full Address:	
Signature of Designated Consignee :	
Name :	
Designation :	
Seal :	
Contact No. :	
Fax No. :	

Note: In addition to sending this document through post, it is requested to send a scanned copy by email to rites_naco@rediffmail.com also.

Copy To:

- (1) To Supplier
- (2) Under Secretary (Admn. P&C, Proc), National AIDS Control Organization, Ministry of Health & Family Welfare, 9th Floor, Chanderlok Building, 36, Janpath, New Delhi 110001, Fax: 011-23731746

9. Final Acceptance Certificate (for Balance 10% Payment)

(This certificate is to be issued to RITES and copy to Supplier and NACO. All the three copies "should be signed in ORIGINAL".)

1	No.	Date

To

MSM Division, RITES Ltd., RITES Office Complex, Annex Building, 4th Floor,

Plot No.144, Sector 44, Gurgaon - 122003, Haryana.

Fax: 91(124)2571659/2571660, Tel: 91(124) 2728-408/405/403

Email: rites_naco@rediffmail.com, rites_naco@rites.com

Project Name	:National HIV/AIDS Control Programme
Purchaser	:RITES Ltd., Gurgaon, Haryana on behalf of
	MoH&FW (NACO)
Contract i.e. NOA No. & Date	:
Description of Goods (Schedule No.)	:
Delivery Lot No.	:
Quantity supplied in Numbers	:
Quantity supplied in Words	:
Name of Supplier	:
Batch No(s).	:
Manufacturing Date(s)	:
Expiry Date(s)	:
Invoice No. and Date	:
Date of Final Acceptance	:
	CERTIFICATE
We confirm having received material a accordance with the contract and entered in	as detailed above in good condition on in the Stock ledger.
Consignee full Address:	
	Signature of Designated Consignee :
	Name :
	Designation :
	Seal :
	Contact No. :
	Fax No. :

Note: In addition to sending this document through post, it is requested to send a scanned copy by email to rites_naco@rediffmail.com also.

Copy To:

- (1) To Supplier
- (2) Under Secretary (Admn. P&C, Proc), National AIDS Control Organization, Ministry of Health & Family Welfare, 9th Floor, Chanderlok Building, 36, Janpath, New Delhi 110001, Fax: 011-23731746

10. AFFIDAVIT(On Stamp Paper)

I	son/daughter of	resident of	solemnly
undertake that	I am an authorized sig	natory of M/s	(insert name of
the company w	vith full address) and I	hereby undertake that the supplie	es for which payments are
being made ha	ve been correctly made	e to the respective consignees. I	take full responsibility for
the correctness	s of the documents sub-	mitted for which the payment ha	as been claimed. I further
undertake that	without prejudice to the	ne rights of purchaser as per the	contract, I shall be solely
responsible if	any of the document is	found to be fake even to make	good any loss suffered by
the purchaser	due to incorrectness o	of the documents submitted by	us for claiming payment
against invoice	e(s) no(s)	(insert details of inv	voices for which payments
are being clain	ned) amounting to	-	
		Name:	
		Address: _	
			(Supplier full address)
Witness 1			
Address:			
Witness 2			
Address	·		

Note:

- 1. The affidavit is to be submitted on a non judicial stamp paper of Rs 100 /-(Rupee hundred) duly notorised and to be signed by the authorized signatory of the firm.
- 2. This affidavit is to be submitted along with the invoices at the time of claiming 80% payment.

11. PROFORMA FOR OTHER DETAILS OF BIDDER, MANUFACTURER AND ITS BANK

1.	Name	& full	address	of the	Manufacturer:
т.	1 tuille	or ruii	uuui Cbb	OI UIC	munulucturor.

- 2. (a) Telephone & Fax No
 - (b) Telex No.
 - (c) Telegraphic address:
 - (d) Email
- 3. Location of the manufacturing factory.
- 4. Name & full address of the Bidder
- 5. (a) Telephone/Mobile & Fax No
 - (b) Telex No.
 - (c) Telegraphic address:
 - (d) Email

Office/Works
Office/Works

Office/Factory/Works
Office/Works

6. Details of two Persons that RITES Ltd. may contact for requests for clarification during bid evaluation:

	1 st	2 nd
(i) Name:		
(ii) Tel number (direct):		
(iii)Mobile No.		
(iv) Email address		

- 7. Bank details from where the Bank Guarantee for Bid Security has been issued:
- (i) Name and address of the Bank:
- (ii) For a foreign bank, name of correspondent Bank in India:
- (iii) Name of the contact Person
- (iv) Phone number/Mobile
- (v) Fax Number
- (vi) Email address

Signature and seal of the Bidder

12. INTEGRITY PACT

Between

RITES LTD. acting for and on behalf of and as an Agent / Power of Attorney Holde	r of
hereinafter called the "Purchaser" AND	
hereinafter referred to as "The Bidder/Suplier"	

Preamble

The Purchaser intends to award, under laid down organizational procedures, contract/s for ______. The Purchaser values full compliance with all relevant laws and regulations, and economic use of resources, and of fairness and transparency in his relations with the Bidder/s and/or Supplier/s.

In order to achieve these goals, the Purchaser will appoint an Independent External Monitor (IEM) who will monitor the Tender process and execution of the contract for compliance with the principles mentioned above.

Section 1 – Commitments of the Purchaser

- (1) The Purchaser commits himself to take all measures necessary to prevent corruption and to observe the following principles:-
 - 1. No employee of the Purchaser, personally or through family members, will in connection with the tender or for the execution of the contract, demand, take a promise for or accept, for self or third person, any material or immaterial benefit which the person is not legally entitled to.
 - 2. The Purchaser will, during the tender process, treat all Bidders with equity and reason. The Purchaser will in particular, before and during the tender process, provide to all Bidders the same information and will not provide to any Bidder confidential/additional information through which the Bidder could obtain an advantage in relation to the tender process or the contract execution.
 - 3. The Purchaser will exclude from the process all known prejudiced persons.
- (2) If the Purchaser obtains information on the conduct of any of his employees which is a criminal offence under the IPC (Indian Penal Code) /PC (Prevention of Corruption) Act, or if there be a substantive suspicion in this regard, the Purchaser will inform its Chief Vigilance Officer and in addition can initiate disciplinary action.

Section 2 – Commitments of the Bidder/Supplier

- (1) The Bidder/Supplier commits himself to take all measures necessary to prevent corruption. He commits himself to observe the following principles during his participation in the tender process and during the contract execution.
 - 1. The Bidder/Supplier will not directly or through any other person or firm, offer, promise or give to any of the Purchaser's employees involved in the tender process or the execution of the contract or to any third person any material or other benefit which he is not legally entitled to, in order to obtain in exchange any advantage of any kind whatsoever during the tender process or during the execution of the contract.
 - 2. The Bidder/Supplier will not enter with other Bidders into any undisclosed agreement or understanding, whether formal or informal. This applies in particular to prices, specifications, certifications, subsidiary contracts, submission or non-submission of bids or any other actions, to restrict competitiveness or to introduce cartelization in the bidding process.

- 3. The Bidder/Supplier will not commit any offence under the relevant IPC/PC Act; further the Bidder/ Supplier will not use improperly, for purposes of competition or personal gain, or pass on to others, any information or document provided by the Purchaser as part of the business relationship, regarding plans, technical proposals and business details, including information contained or transmitted electronically.
- 4. The Bidder/Supplier will, when presenting his bid, disclose any and all payments he has made, is committed to or intends to make to agents, brokers or any other intermediaries in connection with the award of the contract.
- (2) The Bidder/ Supplier will not instigate third persons to commit offences outlined above or be an accessory to such offences.

Section 3-Disqualification from tender process and exclusion from future contracts

If the Bidder/Supplier, before award or during execution has committed a transgression through a violation of Section 2 above, or in any other form such as to put his reliability or credibility in question, the Purchaser is entitled to disqualify the Bidder/Supplier from the tender process or take action as per the procedure mentioned in the "Guideline on banning of business dealing" annexed and marked as **Annexure** "A".

Section 4- Compensation for Damages

- (1) If the Purchaser has disqualified in terms of the provisions in Section 3, the Bidder/Supplier from the tender process prior to the award of contract, the Purchaser is entitled to demand and recover the damages equivalent to Earnest Money Deposit/ Bid Security.
- (2) If the Purchaser has terminated the contract during execution in terms of the provisions under Section 3, the Purchaser shall be entitled to demand and recover from the Supplier the damages equivalent to Performance Security.

Section -5 Previous transgression

- (1) The Bidder/ Supplier declares that no previous transgression occurred in the last 3 years with any other Company in any country conforming to the Anti-Corruption approach or with any other Public Sector Enterprise in India that could justify his exclusion from the tender process.
- (2) If the Bidder/Supplier makes incorrect statement on this subject, he can be disqualified from the tender process or action can be taken as per the procedure mentioned in "Guideline on banning of business dealing".

Section -6 Equal treatment of all Bidders/Suppliers

- (1) The Bidder/Supplier undertakes to demand from all partners (if permitted under the conditions/ clauses of the contract) a commitment to act in conformity with this Integrity Pact and to submit it to the Purchaser before signing the contract.
- (2) The Bidder/ Supplier confirms that any violation by any of his partners to act in conformity with the provisions of this Integrity Pact can be construed as a violation by the Bidder/Supplier himself, leading to possible Termination of Contract in terms of Section 4.
- (3) The Purchaser will disqualify from the tender process all bidders who do not sign this Pact or violate its provisions.

Section 7- Criminal charges against violating Bidders/Suppliers

If the Purchaser obtains knowledge of conduct of a Bidder, Supplier or Partners, or of an employee or a representative or an associate of a Bidder, Supplier, which constitutes corruption, or if the Purchaser has substantive suspicion in this regard, the Purchaser will inform the same to its Chief Vigilance Officer.

Section -8 Independent External Monitor/Monitors

- (1) The Purchaser shall appoint competent and credible Independent External Monitor for this Pact. The task of the Monitor is to review independently and objectively, whether and to what extent the parties comply with the obligations under this agreement.
- (2) The Monitor is not subject to instructions by the representatives of the parties and will perform his functions neutrally and independently. He will report to the MD/RITES Ltd.
- (3) The Bidder/Supplier accepts that the Monitor has the right of access without restriction to all Project documentation of the Purchaser including that provided by the Supplier. The Supplier will also grant the Monitor, upon his request and demonstration of a valid interest, unrestricted and unconditional access to his project documentation. The same is applicable to Partners. The Monitor is under contractual obligation to treat the information and documents of the Bidder/Supplier/Partners with confidentiality.
- (4) The Purchaser will provide to the Monitor sufficient information about all meetings among the parties related to the Project provided such meetings could have an impact on the contractual relations between the Purchaser and the Supplier. The parties offer to the Monitor the option to participate in such meetings.
- (5) As soon as the Monitor notices or has reason to believe that violation of the agreement by the Purchaser or the Bidder/ Supplier, has taken place, he will request the Party concerned to discontinue or take corrective action, or to take any other relevant action. The Monitor can in this regard submit non-binding recommendations. Beyond this, the Monitor has no right to demand from the parties that they act in a specific manner or refrain from action or tolerate action.
- (6) The Monitor will submit a written report to the MD/RITES Ltd. within 8-10 weeks from the date of reference or intimation to him by the Purchaser and should the occasion arise, submit proposal for correcting problematic situations.
- (7) If the Monitor has reported to the MD/RITES Ltd. of a substantiated suspicion of an offence under relevant IPC/PC Act, and the MD/RITES Ltd. has not, within reasonable time, taken visible action to proceed against such offender or reported it to the Chief Vigilance Officer, the Monitor may also transmit this information directly to the Central Vigilance Commissioner.
- (8) The word Monitor would include both singular and plural.

Section – 9 Pact Duration

This pact begins when both parties have legally signed it. It expires for the Supplier when his Security Deposit is released on completion of the contractual obligation.

If any claim is made/lodged during this time the same shall be binding and continue to be valid despite the lapse of this pact specified above, unless it is discharged/determined by MD/RITES Ltd.

Section 10 Other Provisions

- (1) This agreement is subject to Indian Law. Place of performance and jurisdiction shall be as stated in the Contract Agreement.
- (2) Changes and supplements as well as termination notices need to be made in writing.
- (3) If the Supplier is a partnership or a consortium, this agreement must be signed by the Partner in charge/ Lead Member nominated as being incharge and who holds the Power of Attorney signed by legally authorised signatories of all the partners/Members. The Memorandum of Understanding /Joint Venture Agreement will incorporate a provision to the effect that all Members of the Consortium will comply with

the provisions in the Integrity Pact to be signed by the Lead Member on behalf of the Consortium. Any violation of Section 2 above by any of the Partners/Members will be construed as a violation by the consortium leading to possible Termination of Contract in terms of Section 3.

(4) Should one or several provisions of this agreement turn out to be invalid, the remainder of this agreement remains valid. In this case, the parties will strive to come to an agreement to their original intentions.

RITES Ltd. Agent / Power of Attorney	Holder		
(For & on behalf of the Pu	rchaser)	(For the Bio	lder/Supplier)
(Office Seal)			(Office Seal)
Place: Date: Witness 1:			
(Name & Address)			
Witness 2			
(Name & Address)			

Annexure - 1

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12 CHECK LIST

(All the pages of the bid should be Serial Numbered & signed/initialled)

SI. N	lo.	Activity	Yes/No/ NA	Page No. in the Bid
1	(a)	Bid Security for required amount		
	(b)	Bid Security in the form of		
	(i)	Bank Guarantee as per format in Bidding document		
ĺ	(ii)	Draft or Banker's cheque issued by Nationalised bank		
	(c)	Validity Date of Bid Security (Valid upto45-days beyond the bids validity) as specified in ITB Data Sheet clause18.2)		
ŀ	(d)	Amendment in Bid Security (if any)		
2	, ,	The Bank details from where the Bank Guarantee has been issued along with Phone, fax numbers and email Ids. For Banks from outside India the details of the correspondent Bank in India.		
3	(a)	Bid Form duly signed		
	(b)	Power of Attorney in favour of the signatory		
4	(a)	Availing Deemed Export benefits?		
	(b)	Form of Declaration regarding Deemed Export		
5	(0)	The manufacturer's authorization form in Form 7 of Section V		
6		Documents establishing post qualification (ITB 6)		
	a)	Certificate of incorporation of Manufacturer		
	b)	Manufacturing Licence of the good(s) quoted in bid		
,	c)	Proof of Exp in manufacturing & marketing of specific goods for at least one		
	ŕ	year, Indicate Serial No. in performance statement		
((d)	Proof of experience in manufacturing & marketing of similar goods for at least		
		three years, Indicate Serial Nos. in performance statement		
(6	e)	Performance statement as per required Proforma, along with supporting documents		
		viz. (i) Copy of Purchase Orders, (ii) Copy of Invoices, (iii) Proof of Payment		
		received from Purchasers & (iv) Documentary evidence (Client's certificate) in		
		support of satisfactory completion of contract.		
(1	f)	ISO 13485:2003 (or FDA 21 CFR 820) by way of accreditation by an independent recognized certification body		
(!	g)	Indicate Sr. No. in performance statement which establishes the post qualification		
	<i>O</i> ,	criteria of completing one similar contract in last three years		
(l	h)	Certificate of having achieved Annual production rate of equivalent product for last three years by CA.		
(i)	Copies complete set of audited financial statements of accounts (including		
(-	′	balance sheet, profit and loss account, auditor's reports and IT returns) certified by		
		the auditor of the Company for last three financial years		
7		Documents to establish that product is registered in India as per ITB clause 6.4 if applicable		
8		Details of onsite quality control laboratory facilities and services and range of		
~		test conducted.		
9		Affidavit to disclosure about any instance of debarment/blacklisting by state or		
		central Govt. Health organisation		
10		Statement of installed manufacturing capacity certified by appropriate authority		
11		No deviation statement on technical specification		
12		Check list of technical specification. Please give compliance (Yes/No) of each		
		clause of technical specification in tabular form.		

Sl. No.		Activity	Yes/No/ NA	Page No. in the Bid
13	(a)	Agreement with all terms and condition of the bid document		
	(b)	If no, have you indicated deviations		
14	(a)	Mentioned Price in the appropriate Proforma		
	(b)	Conditional or unconditional discount mentioned in the bid (if any)		
15		Copies of original documents defining the constitution or legal status, place of		
		registration, and principal place of business; for both manufacturer & non		
		manufacturer		
16		Undertaking as per clause ITB 6 (C)(6) {The bidder and the manufacturer whose		
		product is offered by the bidder shall disclose instance of previous past performance		
		of his and the manufacturer whose product is procured by the bidder, that may have		
		resulted into adverse actions taken against the bidder during the last three years.		
		Such adverse actions taken against the bidder or manufacturer may be treated as		
		unsatisfactory performance history while deciding the award of contract. If no		
		adverse action has been taken against the Bidder, the Bidder must provide a		
		statement in its bid saying that there has been no such previous past performance		
	1	resulting in adverse actions being taken against him.}		
17	(a)	The bidder shall provide an undertaking that:		
		The proprietor/promoter/director of the firm, its employee, partner or		
		representative is not convicted by a court of law following prosecution for		
		offence involving moral turpitude in relation to business dealings including		
		malpractices such as bribery, corruption, fraud, substitution of bids, interpolation,		
		misrepresentation, evasion, or habitual default in payment of tax levied by law; etc.		
	(b)	The firm does not employ a government servant, who has been dismissed or		
		removed on account of corruption.		
18		Form 11: Proforma for other details of Bidder, Manufacturer and its Bank		
19		Form 12: Integrity Pact		
20		Is your company a Micro or Small Enterprises as per Micro, Small and Medium		
		Enterprises Development (MSMED) Act 2006? If yes, submit the copy of relevant		
		registration certificate.		