# E-TENDER DOCUMENT

**FOR**

## Supply of Pharmaceutical Products for Onward Supplies to Foreign Country

**Tender No: HLL/PSD/RBD/2025-26/TENDER/06 Dt: 25.04.2025**

**E – Tendering**


# HLL Lifecare Limited

(A Govt. Of India Enterprise)
**CIN : U25193KL1966GOI002621**
**HLL Bhavan, Poojappura,**
**Thiruvananthapuram -695012**
**Kerala, India**
**Tel: 0471 2775500, 0471 2350959**
(EXTN – 606 /531)
**Website – www.lifecarehll.com**

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**HLL LIFECARE LIMITED**
(A Government of India Enterprise)
Procurement Services Division
Corporate Head Office, Poojappura.P.O,
Thiruvananthapuram – 695012, Kerala, India
Tel: 0471 2775500, 0471 2350959 (EXTN – 606 /531)

## NOTICE INVITING TENDER (NIT)

**IFB No: HLL/PSD/RBD/2025-26/TENDER/06** **25.04.2025**

HLL Lifecare Limited (HLL), a Government of India Enterprise, invites an e-tender from eligible, competent and experienced parties who are capable of executing the following item/work meeting the requirements as per our tender.

| Sl No | Particulars | Description |
|---|---|---|
| 1 | Name of Item/Work | **Supply of Pharmaceutical Products for Onward Supplies to Foreign Country** |
| 2 | Location of Delivery/Work | **HLL Lifecare Limited, 1 st floor,145, Punjab Expeller Compound, Sihani Chungi,Meerut Road, Ghaziabad, UP-201001, GSTIN: 09AAACH5598K1ZZ ,DL.NO:UP1421B002349, UP1420G000037, UP1420B002371** |
| 3 | Brief description of Item/Work | Supply of Pharmaceutical Products for Onward Supplies to Foreign Country |
| 4 | Bid Security/EMD | Rs.2,00,000.00 |
| 5 | Bid submission fee/Tender fee | Rs.1,500.00 |
| 6 | Period of completion | **Considering the Urgency 7 Days from the date of the Letter of Intent /Notification of Award/ Purchase order, failing which the POs will stand Cancelled.** |
| 7 | Price Validity | 180 days from the date of opening of bid |
| 8 | Eligibility criteria for Bidders | As per Tender document |
| 9 | HLL A/c Details for payment of Tender Fees and EMD (Payment mode: NEFT/RTGS) | Name of Bank: HDFC BANK<br>A/c number: 00630330000605<br>IFSC Code: HDFC0000063<br>Branch name: Vazhuthacaud, Thiruvananthapuram |
| 10 | Last date and time for online submission of online bids | **28-04-2025 at 14:00 hrs** |
| 11 | Date and time of opening of e-tender | **29-04-2025 at 14:00 hrs** |
| 12 | Address for Communication at HLL regarding the tender | Vice President (PS) and GH (HCS)<br>Procurement Services Division<br>HLL Lifecare Limited<br>Corporate & Regd. Office<br>HLL Bhavan, Poojappura,<br>Thiruvananthapuram-695012<br>E-mail: sdrbdsouth@lifecarehll.com |

**एचएलएल लाइफ़केयर लिमिटेड**
(भारत सरकार का उद्यम)
सोर्सिंग प्रभाग
कॉर्पोरेट मुख्यालय, पूजप्पुरा.पी.ओ.,
तिरुवनंतपुरम - 695012, केरल, भारत
दूरभाष: 0471 2775500, 0471 2350959 (एक्स्ट - 606/531)

## निविदा आमंत्रण सूचना (एनआईटी)

आईएफबी संख्या: एचएलएल/पएसडी/आरबीडी/2025-26/निविदा/06 25.04.2025

एचएलएल लाइफकेयर लिमिटेड (एचएलएल), भारत सरकार का उद्यम, योग्य, सक्षम और अनुभवी पार्टियों से एक ई-निविदा आमंत्रित करता है, जो हमारी निविदा के अनुसार आवश्यकताओं को पूरा करने के लिए निम्नलिखित मद/कार्य को निष्पादित करने में सक्षम हैं।

| क्र.सं. | ब्यौरा | विवरण |
|---|---|---|
| 1 | मद/कार्य का नाम | **Supply of Pharmaceutical Products for Onward Supplies to Foreign Country** |
| 2 | सुपुर्दगी/कार्य का स्थान | **HLL Lifecare Limited, 1 st floor,145, Punjab Expeller Compound, Sihani Chungi,Meerut Road, Ghaziabad, UP-201001,GSTIN: 09AAACH5598K1ZZ ,DL.NO:UP1421B002349, UP1420G000037, UP1420B002371** |
| 3 | मद/कार्य का संक्षिप्त विवरण | Supply of Pharmaceutical Products for Onward Supplies to Foreign Country |
| 4 | बोली प्रतिभूति/ईएमडी | Rs.2,00,000.00 |
| 5 | बोली प्रस्तुतीकरण शुल्क / निविदा शुल्क | Rs.1,500.00 |
| 6 | पूरा करने की अवधि | **Considering the urgency,7 Days from the date of the Letter of Intent /Notification of Award/ Purchase order, failing which the POs will stand Cancelled.** |
| 7 | मूल्य वैधता | 180 days from the date of opening of bid |
| 8 | बोलीदाताओं के लिए पात्रता मानदंड | As per Tender document |
| 9 | निविदा शुल्क और ईएमडी के भुगतान के लिए एचएलएल खाते का विवरण (भुगतान मोड: एनईएफटी/आरटीजीएस) | Name of Bank: HDFC BANK<br>A/c number: 00630330000605<br>IFSC Code: HDFC0000063<br>Branch name: Vazhuthacaud, Thiruvananthapuram |
| 10 | ऑनलाइन बोलियों के ऑनलाइन प्रस्तुतीकरण की अंतिम तारीख और समय | **28-04-2025 at 14:00 hrs** |
| 11 | ई-निविदा खोलने की तिथि और समय | **29-04-2025 at 14:00 hrs** |
| 12 | एचएलएल में निविदा के संबंध में पत्र व्यवहार के लिए पता | Vice President (PS) and GH (HCS)<br>Procurement Services Division<br>HLL Lifecare Limited<br>Corporate & Regd. Office<br>HLL Bhavan, Poojappura,<br>Thiruvananthapuram-695012<br>E-mail: sdrbdsouth@lifecarehll.com |

## GENERAL INSTRUCTIONS TO BIDDERS

1. This tender is an e-Tender and is being published online in Government eProcurement portal, https://etenders.gov.in/eprocure/app
2. Bid documents including the Bill of Quantities (BoQ) can be downloaded free of cost from the Central Public Procurement Portal of Government of India (e-portal). All Corrigendum/extension regarding this e-tender shall be uploaded on this website i.e. https://etenders.gov.in/eprocure/app.
3. The tender and its corrigendum/extension will also be published in our company website, URL address: http://www.lifecarehll.com/tender.
4. The tendering process is done online only at Government eProcurement portal (URL address: https://etenders.gov.in/eprocure/app). Aspiring bidders may download and go through the tender document.
5. All bid documents are to be submitted online only and in the designated cover(s)/envelope(s) on the Government eProcurement website. Tenders/bids shall be accepted only through online mode on the Government eProcurement website and no manual submission of the same shall be entertained. Late tenders will not be accepted.
6. The complete bidding process is online. Bidders should be in possession of valid Digital Signature Certificate (DSC) of class II or above for online submission of bids. Prior to bidding DSC need to be registered on the website mentioned above. If the envelope is not digitally signed & encrypted the Purchaser shall not accept such open Bids for evaluation purpose and shall be treated as non-responsive and shall be rejected.
7. Bidders are advised to go through “Bidder Manual Kit”, “System Settings” & “FAQ” links available on the login page of the e-Tender portal for guidelines, procedures & system requirements. In case of any technical difficulty, Bidders may contact the help desk numbers & email ids mentioned at the e-tender portal.
8. Bidders are advised to visit CPPP website https://etenders.gov.in regularly to keep themselves updated, for any changes/modifications/any corrigendum in the Tender Enquiry Document.
9. The bidders are required to submit soft copies of their bids electronically on the CPP Portal, using valid Digital Signature Certificates. The instructions given below are meant to assist the bidders in registering on the CPP Portal, prepare their bids in accordance with the requirements and submitting their bids online on the Government eProcurement Portal.

   9.1 Registration

   a) Bidders are required to register in the Government e-procurement portal, obtain ‘Login ID’ & ‘Password’ and go through the instructions available in the Home page after log in to the CPP Portal (URL: https://etenders.gov.in/eprocure/app), by clicking on the link “Online bidder Enrolment” on the CPP Portal which is free of charge.

   b) As part of the enrolment process, the bidders will be required to choose a unique user name and assign a password for their accounts.

   c) Bidders are advised to register their valid email address and mobile numbers as part of the registration process. These would be used for any communication from the CPP Portal.

   d) They should also obtain Digital Signature Certificate (DSC) in parallel which is essentially required for submission of their application. The process normally takes 03 days’ time. The bidders are required to have Class II or above digital certificate or above with both signing and encryption from the authorized digital signature Issuance

Company. Please refer online portal i.e. - https://etenders.gov.in/eprocure/app for more details.

e) Upon enrolment, the bidders will be required to register their valid Digital Signature Certificate (Class II or above Certificates with signing key usage) issued by any Certifying Authority recognized by CCA India (e.g. Sify /nCode / eMudhra etc.), with their profile.

f) Bidder then logs in to the site through the secured log-in by entering their user ID/password and the password of the DSC / e-Token.

g) The Bidder intending to participate in the bid is required to register in the e-tenders portal using his/her Login ID and attach his/her valid Digital Signature Certificate (DSC) to his/her unique Login ID. He/She have to submit the relevant information as asked for about the firm/contractor. The bidders, who submit their bids for this tender after digitally signing using their Digital Signature Certificate (DSC), accept that they have clearly understood and agreed the terms and conditions including all the Forms/Annexure of this tender.

h) Only those bidders having a valid and active registration, on the date of bid submission, shall submit bids online on the e-procurement portal.

i) Only one valid DSC should be registered by a bidder. Please note that the bidders are responsible to ensure that they do not lend their DSC's to others which may lead to misuse.

j) Ineligible bidder or bidders who do not possess valid & active registration, on the date of bid submission, are strictly advised to refrain themselves from participating in this tender.

9.2 **Searching for Tender Documents**

a) There are various search options built in the CPP Portal, to facilitate bidders to search active tenders by several parameters. These parameters could include Tender ID, Organization Name, Form of Contract, Location, Date, Value etc. There is also an option of advanced search for tenders, wherein the bidders may combine a number of search parameters such as Organization

b) Once the bidders have selected the tenders they are interested in, they may download the required documents/tender schedules. These tenders can be moved to the respective 'My Tenders' folder. This would enable the CPP Portal to intimate the bidders through SMS/ e-mail in case there is any corrigendum issued to the tender document.

c) The bidder should make a note of the unique Tender ID assigned to each tender, in case they want to obtain any clarification/help from the Helpdesk

9.3 **Preparation of Bid**

a) Bidder should take into account any corrigendum published on the tender document before submitting their bids.

b) Please go through the tender document carefully to understand the documents required to be submitted as part of the bid. Please note the number of covers in which the bid documents have to be submitted, the number of documents - including the names and content of each of the document that need to be submitted. Any deviations from these may lead to rejection of the bid.

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

d) To avoid the time and effort required in uploading the same set of standard documents which are required to be submitted as a part of every bid, a provision of uploading such standard documents (e.g. PAN card copy, annual reports, auditor certificates etc.) has been provided to the bidders. Bidders can use "My Space" or ''Other Important Documents'' area available to them to upload such documents. These documents may be directly submitted from the "My Space" area while submitting a bid, and need not be uploaded again and again. This will lead to a reduction in the time required for bid submission process.

e) Note: My Documents space is only a repository given to the Bidders to ease the uploading process. If Bidder has uploaded his Documents in My Documents space, this does not automatically ensure these Documents being part of Technical Bid.

**10.** More information useful for submitting online bids on the CPP Portal may be obtained at https://etenders.gov.in/eprocure/app

**11.** Tenderer are required to upload the digitally signed file of scanned documents. Bid documents may be scanned with 100 dpi with black and white option which helps in reducing size of the scanned document. Uploading application in location other than specified above shall not be considered. Hard copy of application shall not be entertained.

**12.** Any queries relating to the process of online bid submission or queries relating to CPP Portal in general may be directed to the 24x7 CPP Portal Helpdesk. The 24x7 Help Desk details are as below: -

For any technical related queries please call at 24 x 7 Help Desk Number:
0120-4001 062, 0120-4001 002, 0120-4001 005, 0120-6277 787

Note:- International Bidders are requested to prefix +91 as country code

E-Mail Support: For any Issues or Clarifications relating to the published tenders, bidders are requested to contact the respective Tender Inviting Authority
Technical - support-eproc@nic.in, Policy Related - cppp-doe@nic.in

**13.** Bidders are requested to kindly mention the URL of the portal and Tender ID in the subject while emailing any issue along with the contact details.

**14.** Any queries relating to the tender document and the terms and conditions contained therein should be addressed to the Tender Inviting Authority for a tender or the relevant contact person indicated in the tender. Address for communication and place of opening of bids:

**Vice President (PS) and GH (HCS)**
**Procurement Services Division**
**HLL Lifecare Limited**
**Corporate & Regd. Office**
**HLL Bhavan, Poojappura,**
**Thiruvananthapuram-695012**

**15.** E-mail: sdrbdsouth@lifecarehll.com The bids shall be opened online at the **Office of the Vice President (PS) and GH (HCS)** in the presence of the Bidders/their authorized representatives who wish to attend at the above address. If the tender opening date happens to be on a holiday or non-working day due to any other valid reason, the tender opening process will be done on the next working day at same time and place.

**16.** More details can be had from the Office of the Vice President (PS) and GH (HCS) during working hours. The Tender Inviting Authority shall not be responsible for any failure, malfunction or breakdown of the electronic system while downloading or uploading the documents by the Bidder during the e-procurement process.

**17.** A firm/bidder shall submit only one bid in the same bidding process. A Bidder (either as a firm or as an individual or as a partner of a firm) who submits or participates in more than one bid will cause all the proposals in which the Bidder has participated to be disqualified.

**18. Online Tender Process:**

The tender process shall consist of the following stages:

i. Downloading of tender document: Tender document will be available for free download on Government e-procurement portal (URL: https://etenders.gov.in/eprocure/app).

ii. Pre-bid meeting: Not Applicable for this tender

iii. Publishing of Corrigendum: All corrigenda shall be published on Government e-procurement portal (URL: https://etenders.gov.in/eprocure/app) and HLL website (URL address: http://www.lifecarehll.com/tender) and shall not be available elsewhere.

iv. Bid submission: Bidders have to submit their bids along with supporting documents to support their eligibility, as required in this tender document on Government e-procurement portal. No manual submission of bid is allowed and manual bids shall not be accepted under any circumstances.

v. Opening of Technical Bid and Bidder short-listing: The technical bids will be opened, evaluated and shortlisted as per the eligibility and technical qualifications. All documents in support of technical qualifications shall be submitted (online). Failure to submit the documents online will attract disqualification. Bids shortlisted by this process will be taken up for opening the financial bid.

vi. Opening of Financial Bids: Bids of the qualified bidders shall only be considered for opening and evaluation of the financial bid on the date and time mentioned in critical date's section.

**19. Tender Processing Fees and Bid Security (EMD):**

Tender fee (Non-refundable) and EMD as per the tender conditions shall be paid separately, thru RTGS/NEFT transfer in the following HLL A/c details:

| | | |
|---|---|---|
| Name of Bank | : | HDFC BANK |
| A/c number | : | 00630330000605 |
| IFSC Code | : | HDFC0000063 |
| Branch name | : | VAZHUTHACAUD, Thiruvananthapuram |

Document of the above transactions (UTR NUMBER and DATE OF UTR) completed successfully by the bidder, shall be uploaded at the locations separately while submitting the bids online.

Note: Any transaction charges levied while using any of the above modes of payment has to be borne by the bidder. The supplier / contractor's bid will be evaluated only if payment is effective on the date and time of bid opening.

**20.** HLL Lifecare Limited does not bind themselves to accept the lowest or any bid or to give any reasons for their decisions which shall be final and binding on the bidders.

**21.** HLL Lifecare Limited reserves to themselves the right of accepting the whole or any part of the tender and bidder shall be bound to perform the same at his quoted rates.

**22.** In case, it is found during the evaluation or at any time before placing of PO or after its execution and during the period of subsistence thereof, that one or more of the eligibility conditions have not been met by the bidder or the applicant has made material misrepresentation or has given any materially incorrect or false information, appropriate legal/penal etc., action shall be taken by HLL Lifecare as deemed fit.

**23.** Conditional bids and bids not uploaded with appropriate/desired documents may be rejected out rightly and decision of HLL Lifecare Limited in this regard shall be final and binding.

**24.** The technical bids should be uploaded as per the requirements of NIT and should not contain price information otherwise the bid will be rejected.

**25.** HLL Lifecare Limited Ltd. reserves the right to verify the claims made by the bidders and to carry out the capability assessment of the bidders and the HLL Lifecare Limited's decision shall be final in this regard.

**26.** Submission Process:

For submission of bids, all interested bidders have to register online as explained above in this document. After registration, bidders shall submit their Technical bid and Financial bid online on Government e-procurement portal (URL: https://etenders.gov.in/eprocure/app).

**Note:- It is necessary to click on "Freeze bid" link / icon to complete the process of bid submission otherwise the bid will not get submitted online and the same shall not be available for viewing/ opening during bid opening process**.

**VICE PRESIDENT (PS) AND GH (HCS)**

# INSTRUCTIONS TO THE BIDDERS (ITB)

## Section 1

### I. COMPANY BACKGROUND:

HLL Lifecare Limited (HLL) is a public sector undertaking under the administrative control of the Ministry of Health & Family Welfare, Government of India. HLL's purpose of business is to provide quality healthcare products and services at affordable rates. In its quest to become a comprehensive healthcare solutions provider, HLL had diversified into hospital products and healthcare services, while nurturing its core business of providing quality contraceptives. HLL Vending Business Division is offering solution for retailing and making available range of HLL's - quality healthcare products / Sanitary Napkins / Condoms etc., products through state-of-art Vending machines. HLL has also forayed into the Service sectors of Healthcare Diagnostics and Pharmaceutical retail business for more than 10 years.

### TENDER DETAILS

HLL Lifecare Limited (HLL), a Government of India Enterprise, invites online bids from the eligible, competent and experienced Suppliers/Dealers/Manufacturers for:

a) Supply of Pharmaceutical Products for Onward Supplies to Foreign Country as per the below said items. Supplies to be effected and deliveries to be made to HLL Ghaziabad Depot.

b) Supply to be made on Door delivery basis to our warehouse at **HLL Lifecare Limited, 1 st floor,145, Punjab Expeller Compound, Sihani Chungi,Meerut Road, Ghaziabad, UP-201001, GSTIN: 09AAACH5598K1ZZ ,DL.NO:UP1421B002349, UP1420G000037, UP1420B002371**

c) The total quantity mentioned is only an indicative quantity and may change depending on actual requirement.

d) Suppliers must ensure strict compliance to all statutory regulations, quality standards and packing material specifications (as applicable ) detailed in Annexure – 5

### II. Product List

***In BOQ, Bidders are requested to quote for per piece rate only and GST amount (not in percentage)***

| Sl.No | Item | UOM | Required quantity |
|---|---|---|---|
| 1 | Cefotaxime for Inj. 1g Vial | VIAL | 6,60,000 |
| 2 | Ceftazidime for Inj. 1g Vial | VIAL | 1,80,000 |
| 3 | Ceftriaxone for Inj. 1g Vial | VIAL | 15,00,000 |
| 4 | Imipenem+Cilastatin for Inj.500mg | VIAL | 60,000 |
| 5 | Doxycycline Cap. 100mg | CAP | 68,00,000 |
| 6 | Amikacin Inj. 500mg/2mL Vial | VIAL | 2,56,000 |
| 7 | Vancomycin Inj. 500mg Vial | VIAL | 1,40,000 |
| 8 | Teicoplanin Inj. 400mg Vial | VIAL | 78,000 |
| 9 | Sulbactam + Cefoperazone Inj.2g Vial | VIAL | 12,000 |
| 10 | Sodium Stibogluconte Inj.10g/100mL Vial | VIAL | 600 |
| 11 | Fluconazole Inj. 200mg in100mL Vial | VIAL | 30,000 |
| 12 | Fluconazole Cap. 200mg | CAP | 90,000 |
| 13 | Amphotericin Inj. 50mg Vial | VIAL | 3,750 |
| 14 | Nystatin Tab. 500,000IU | TAB | 6,00,000 |
| 15 | Colistimethate for Injection 1,000,000IU Vial | VIAL | 1,09,000 |
| 16 | Ketoconazole Tab 200mg | TAB | 30,000 |

| | | | |
|---|---|---|---|
| 17 | Adenosine Inj. 6mg/2mL Vial | AMP | 7,500 |
| 18 | Amiodarone Tab. 100mg | TAB | 10,50,000 |
| 19 | Amiodarone Inj. 150mg/3mL Amp. | AMP | 35,000 |
| 20 | Metoprolol Inj. 5mg/5mL Amp. | AMP | 1,000 |
| 21 | Sodium Nitroprusside Inj. 50mgVial | VIAL | 250 |
| 22 | Glyceryl Trinitrate(Nitroglycerin) Inj. 50mg/10mL Vial or Amp. | VIAL | 27,000 |
| 23 | Verapamil Injection 5mg in 2mL Ampoule | AMP | 10,500 |
| 24 | Isoprenaline Inj. 2mg/2mL Amp. | AMP | 5,400 |
| 25 | Noradrenaline Inj. 4mg/2mL Amp | AMP | 7,50,000 |
| 26 | Phenylephrine Inj. 10mg/1mLVial | VIAL | 2,100 |
| 27 | Protamine Sulphate Inj.50mg/5mL | VIAL | 6,000 |
| 28 | Tranexamic Acid Tab/Cap 500mg | TAB | 12,00,000 |
| 29 | Factor VIII 200-350IU Vial with vW Factor | VIAL | 4,500 |
| 30 | Dried Recombinant Factor VIII Fraction 200IU-350IU Vial | Vial | 9,996 |
| 31 | Dried Recombinant Factor VIII Fraction 500IU Vial | Vial | 15,000 |
| 32 | Papaverine HCl Inj.60mg/2mL Amp. | AMP | 3,200 |
| 33 | Triclofos Oral Solution 500 mg/5 mL in 30 mL Bottle | Bot | 8,000 |
| 34 | Paracetamol Infu. 10mg/mL,100mL | VIAL | 76,000 |
| 35 | Ethosuximide Cap 250mg | CAP | 85,000 |
| 36 | Flunarizine HClTab. 5mg | TAB | 75,00,000 |
| 37 | Disulfiram Tab. 250mg | TAB | 1,00,000 |
| 38 | Levetiracetam Inj. 500mg/5mLVial/Amp. | VIAL | 31,500 |
| 39 | Fat emulsion Inj. 20%,100mLBot. | BOT | 5,100 |
| 40 | Amino acid solution for intravenous infusion, 100mL Bottle | BOT | 4,800 |
| 41 | Thiamine Tab.100mg | TAB | 6,90,000 |
| 42 | Epoetin Inj. 4000IU PFS | PFSY | 8,00,000 |
| 43 | Desferrioxamine Inj. 500mg | VIAL | 2,70,000 |
| 44 | Human Albumin Sol. 20%, 50mLBot. | BOT | 2,00,000 |
| 45 | Phosphate Tab. 500 mg | TAB | 1,56,000 |
| 46 | Total Parenteral Nutrition in500mL-1,500mL Collap. Bag | BAG | 15,000 |
| 47 | Salbutamol Respiratory Sol.0.5%,15mL Vial | VIAL | 2,75,000 |
| 48 | Pirfenidone Tablets 200mg | TAB | 5,50,000 |
| 49 | Rabies Vaccine (Human use)0.5mL/1mL-Inactivated | DOSE | 15,00,000 |
| 50 | Pneumococcal Vaccine SingleDose Vial (PPSV-23) | VIAL | 4,500 |
| 51 | Varicella Vaccine 0.5mL Vial | VIAL | 450 |
| 52 | Anti Rabies Serum Inj. 1,000I.U./5mL Amp. | VIAL | 24,000 |
| 53 | Anti Rabies HumanImmunoglobulin 300IU Vial | VIAL | 1,200 |
| 54 | Biphasic Isophane Insulin(Human) Inj. 30/70 Vial | VIAL | 22,00,000 |
| 55 | InsulinIsophane (human) 1,000IU/10mLVial | VIAL | 36,000 |
| 56 | Insulin Soluble (Human) Inj.1,000IU/10mL Vial | VIAL | 72,000 |
| 57 | Glucagon HCl Inj. 1 IU Vial | VIAL | 1,200 |

| | | | |
|---|---|---|---|
| 58 | Dexamethasone Tab. 4 mg | TAB | 2,70,000 |
| 59 | Hydroxyprogesterone Inj.250mg/1mL Amp. | AMP | 9,000 |
| 60 | Tetracosactrin Inj.250mcg/1mL Amp. | AMP | 1,300 |
| 61 | Somatropin for Inj. 2IU-30IU | IU | 2,24,000 |
| 62 | Desmopresin Tab. 100mcg | TAB | 60,000 |
| 63 | Desmopressin acetate NasalSpray 10mcg/metered spray( 50 MS) | VIAL | 4,200 |
| 64 | Triamcinolone Acetonide Inj (preser. free) 40mg/mLVial | VIAL | 1,500 |
| 65 | Diazoxide tab. 50mg | TAB | 12,000 |
| 66 | Dutasteride Cap. 0.5 mg | CAP | 30,000 |
| 67 | Estradiol Valerate Tab. 1mg | TAB | 18,000 |
| 68 | Metoclopramide Tab. 10mg | TAB | 11,00,000 |
| 69 | Metoclopramide Inj.10mg/2mLAmp. | AMP | 11,00,000 |
| 70 | Ranitidine Inj. 50mg/2mL Amp. | AMP | 6,50,000 |
| 71 | Mesalazine tab. 400mg | TAB | 1,50,000 |
| 72 | Macrogol 3350/4000 (10g-13.125g of PEG) Oral powder sachets (with or without electrolytes) | Sach | 9,498 |
| 73 | Terlipressin Acetate Inj.1mg Vial/Amp. | VIAL | 21,000 |
| 74 | Fluorescein Sodium Inj.10%, 2mL-5mL Vial | VIAL | 2,900 |
| 75 | Olopatadine HCl Eyedrops 1mg/mL, 5mL Dropper Bot. | BOT | 18,000 |
| 76 | Chlorambucil Tab. 2mg | TAB | 15,000 |
| 77 | Cyclophosphamide Tab. 50mg | TAB | 1,05,000 |
| 78 | Epirubicin HCl Inj.50mg/25mL Vial | VIAL | 4,500 |
| 79 | Mitomycin for Inj. 2mg Vial | VIAL | 6,000 |
| 80 | Cytarabine Inj. 100mg/5mL Vial(Not for Intrathecal use) | VIAL | 1,020 |
| 81 | Cytarabine Inj. 100mg/mLVial ,Preservative free | VIAL | 1,100 |
| 82 | Fluorouraci Inj. 250mg/10mLVial | VIAL | 600 |
| 83 | Gemcitabine HClInj. 200mg Vial | VIAL | 3,000 |
| 84 | Methotrexate Inj. 1g/10mL Vial | VIAL | 13,200 |
| 85 | Etoposide Cap.100mg | CAP | 2,400 |
| 86 | Etoposide Cap. 50mg | CAP | 900 |
| 87 | Temozolomide Cap. 100mg | CAP | 9,600 |
| 88 | Carboplatin inj.150mg/15mLVial with Diluent | VIAL | 2,250 |
| 89 | Carboplatin Inj. 450mg/45mLVial | VIAL | 12,600 |
| 90 | Procarbazine Cap. 50mg | CAP | 2,400 |
| 91 | Imatinib Mesilate Tab/Cap100mg | CAP | 3,60,000 |
| 92 | Filgrastim Inj 300mcg in0.5mL-1mL, PFS/Vial | VIAL | 1,35,000 |
| 93 | Tamoxifen Tab. 20mg | TAB | 9,00,000 |
| 94 | Goserelin Acetate Implant3.6mg | IMPL | 11,400 |
| 95 | Abiraterone Acetate Tab.250mg | TAB | 3,00,000 |
| 96 | Mesna Inj. 200mgin 2mL Vial/Ampoule | VIAL | 18,000 |
| 97 | Clotrimazole Pessaries 100mg | PESS | 1,02,000 |
| 98 | Tolterodine SR Cap. 2mg | CAP | 60,000 |
| 99 | Solifenacin Tab. 5mg | TAB | 24,000 |
| 100 | Oxybutynine Tab. 2.5mg | TAB | 1,80,000 |

| | | | |
|---|---|---|---|
| 101 | Etomidate Inj. 20mg/10mLVial/Amp. | VIAL | 1,800 |
| 102 | GlycopyrrolateInj. 200mcg/mL Vial | AMP | 24,000 |
| 103 | Flumazenil Inj. 500mcg/5mLVial/Amp. | VIAL | 660 |
| 104 | Naloxone Inj. 400mcg/mL Amp. | AMP | 6,000 |
| 105 | Dantroline Sodium Inj. 20mg Vial | VIAL | 350 |
| 106 | Lidocaine Topical aerosol10%, 50mL Bot. | BOT | 2,700 |
| 107 | Lidocaine HCl Gel 2%,30g Tube | TUBE | 2,40,000 |
| 108 | Lidocaine 2% + Adrenalin1:80,000 Inj. 30mL Vial | VIAL | 2,10,000 |
| 109 | Dexmedetomidine HCl Inj200mcg/2mL Vial | VIAL | 2,400 |
| 110 | Morphine Tab. 30mg | TAB | 1,50,000 |
| 111 | Morphine Sulphate Oral solu.2mg/1mL, 100mL Bott. | BOT | 750 |

## III. TECHNICAL SPECIFICATION

All Pharmaceutical products should have relevant & valid WHO GMP certificate.

## IV. PACKING SPECIFICATION

| Packing Material Specification | |
|---|---|
| BABY CARTON | 350Gsm foreign art card with 4 Colour printing, single side printing,Tuck in flap system,finishing with outer gloss lamination with dye punching and pasting |
| DISPLAY CARTON | 350Gsm ITC saffaire graphic 4 Colour printing, single side printing,Tuck in flap system with locked bottom,finishing with outer gloss lamination with dye punching and pasting |
| MASTER CARTON (CORRUGATED BOX) | Narrow Flute 7 Ply Corrugated Card Board Box Total Gsm = >1147 inner & outer ply virgin kraft paper of which outer ply to be alkali resistant with bitumen. The box shall be single piece with double stapling using flat wire of MS or GI material as per ISI 10066, 1981. Gsm: - (outer Line bituminised) 160, Inner lining 120x3 flute= 150x3 (@35% extra for 3 ply corrugating). Direction of flute: Vertical, nature of flute: Narrow. Punch Resistance - Not less than 45deg. C 0Zs per tear inch. Bursting strength: 18 kg/cm2 (min.) (bursting factor not less than 20, Gum -Nature: Starch Based.). |
| ALUMINIUM FOIL | Thickness- 0.021mm to 0.022 mm (21 to 22 micron), Gsm - 59 (54 to 56 aluminium + HSL 3 min.) |
| PVC/PVDC | Food Grade Thermo formable transparent blister foil. Thickness= 0.35 mm max. Gsm= 320 to 330, Sealing= Proper sealing, PVC= Non Toxic - PVC food grade, Yield= 3.125 to 3.03 mt2/ kg |
| LEAFLET | Maplitho Deluxe Paper Of 70 Gsm Min. Two Folds Printed In Single Colour (Black) |

Bidders may adopt appropriate packing mode, however shall ensure that the packing must be suitable for cargo handling/export by air.

There may also be a branding requirement in the tertiary packing.

Product literature must be printed in English

SPECIMEN LABEL FOR OUTER CARTON
Product Name:
Batch No. :
Mfg. Date:
Exp. Date:

Total Quantity:
Net Weight of the Carton:
Manufactured By:

## V. SUPPLY LOCATION

Supply to be made on Door delivery basis to our below warehouse;

**HLL Lifecare Limited,**
**1 st floor, 145, Punjab Expeller Compound,**
**Sihani Chungi,Meerut Road, Ghaziabad,UP-201001,**
**GSTIN: 09AAACH5598K1ZZ,**
**DL.NO:UP1421B002349, UP1420G000037, UP1420B002371**

# Section 2:

## 1. ELIGIBLE BIDDERS

Bidders are requested to submit the Tender processing fee and EMD online on or before the due date as mentioned in the NIT. The bidders who failed to submit the tender fee and EMD before the submission deadline will be considered as technically non responsive.

A Bidder should have following eligibility criteria as of the date of bid submission and should continue to meet these till the award of the contract.

1.1. Valid manufacturing license/Factory License (Self–attested Copy) along with the list of products manufactured in this facility wherever applicable. The quoted products should be highlighted for ready reference.

1.2. All Pharmaceutical products should have relevant & valid WHO GMP certificate.

1.3. Original Manufacturers having a minimum average annual turnover of Rs.1 Crore (Rupees One Crore only) during the last three years i.e., 2021-2022, 2022-2023 and 2023-2024 (original/ provisional) will only be eligible for participation.

Authorized agents are also eligible to bid provided their minimum average turnover in the last three years i.e., 2021-2022, 2022-2023 and 2023-2024 (original/ provisional) is Rs.50 lakh (Rupees Fifty lakhs only) and their Principal manufacturers meets the eligibility criteria for principal manufacturer as specified above.

In case of bid by authorized agents, manufacturers authorization form must be attached with the bid submitted. If an Original Manufacture is participating in the tender but wishes to make the supplies through its authorised agent, the manufacturer has to ensure that the Authorised agent meets all the eligible criteria mentioned, including minimum average turnover in the last three years i.e., 2021-2022, 2022-2023 and 2023-2024 (original/ provisional) (original / provisional) is Rs.50 lakh (Rupees Fifty lakhs only) and documentary proof for the same has to be attached along with original authorization letter.

1.4. The offered supply should comply with the provisions of the relevant standards for the product as applicable as amended up to date.

1.5. The Bidder must submit an in-house batch wise COA pass test report (hard copy ), manufacturing license with product permission, WHO GMP certificate for every batch of items along with consignment and soft copies of COA's shall be send to sdrbdsouth@lifecarehll.com.

1.6. The bidder should submit MSDS certificate for all dangerous goods & all necessary documents as specified against each item in the technical specification along with the consignment, if asked

1.7. Firm should submit a non-conviction certificate issued by the State Drug Controller, to the effect that the manufacturer has not been convicted under the Drugs and Cosmetics Act, 1940 and rules thereunder during the preceding three years and that no case / proceedings is pending against the manufacturer in any Court of Law in India under the Drugs & Cosmetics Act and for Non-Pharma items self-declared non conviction from bidder/manufacture to be submitted.

1.8. Primary manufacturers/authorized agents are allowed to participate in the Tender. Manufacturer's authorization form in original may be submitted by participating authorized agents.

1.9. The bidders who are able to supply the items within 7 days from the date of PO are only eligible to bid.

1.10. Suppliers must ensure strict compliance to all statutory regulations, quality standards and Packing material specifications (as applicable) detailed in Annexure 5.

1.11. A firm/bidder shall submit only one bid in the same bidding process. A Bidder (either as a firm or as an individual or as a partner of a firm) who submits or participates in more than one bid will cause all the proposals in which the Bidder has participated to be disqualified.

1.12. Bidders who are eligible as per the Provisions of Public Procurement –Preference to Make in India Order No.P-45021/12/2017PP (BE-II), 2017 (published by Department for Promotion of Industry and Internal Trade) inclusive of the latest amendments are eligible to participate in the tender. A self-declaration as per Annexure 14 with respect to this order must be submitted.

1.13. Any bidder from a country which shares a land border with India will be eligible to bid in this tender only if the bidder is registered with Competent Authority, as per order no F.No.6/18/2019-PPD dated 23-July-2020 (Rule 144 (xi) of the GFR, 2017 and any amendments issued thereafter) inclusive of the latest amendments issued by Ministry of Finance, GOI at Annexure 13 of this bidding document. The bidder must comply with all provisions mentioned in this order. A self-declaration as per Annexure 13 with respect to this order must be submitted.

1.14. Purchase preference to Micro and Small Enterprises (MSEs): Purchase preference will be given to MSEs as defined in Public Procurement Policy for Micro and Small Enterprises (MSEs) Order, 2012 dated 23.03.2012 issued by Ministry of Micro, Small and Medium Enterprises and its subsequent Orders/Notifications issued by concerned Ministry.

1.15. (a) Bidder/ manufacturer who has been de-recognized/debarred/banned/blacklisted for the product mentioned in the tender by any other State Government / Central Govt. Organization /State Medical Corporations/ Director Health Services and or convicted by any court of law due to (i) quality failure of the drug(s) supplied (NSQ/ Spurious/ Adulterated/ Misbranded etc.) (ii) Submission of fake or forged documents (iii) Submission of incorrect information / Suppression of vital information & facts can't participate for that product in the tender during the period of de-recognition / debarment/ Banned/blacklisted. Bidder / manufacturing unit which has been de-recognized/ debarred/banned/blacklisted by State Medical Corporation for any reasons can't participate for that product in the tender during the period of de-recognition/debarment/banned. If we, at a later date, are found guilty of suppressing facts in this regard, such act on our part shall be considered a fraudulent practice in accordance with the Instructions to Bidders and the Purchaser shall be entitled to reject our BID for the product quoted, submitted by us against this Tender.

(b) Any bidder who has been convicted by a competent court of law for supplying (NSQ/ Spurious/ Adulterated/ Misbranded etc.) drugs within a period of last 3 years from the date of floating of tender shall not be eligible to participate in the tender for that product..

(c) Any bidder who is a distributor/ authorized agent then they should ensure that their Principal manufacturer is not been de-recognized/debarred/banned/blacklisted for the quoted product by any other State Government / Central Govt. Organization /State Medical Corporations/ Director Health Services and or convicted by any court of law due to (i) quality failure of the drug(s) supplied (NSQ/ Spurious/ Adulterated/ Misbranded etc.) (ii) Submission of fake or forged documents (iii) Submission of incorrect information / Suppression of vital information & facts can't participate for that product in the tender during the period of de-recognition / debarment/ Banned/blacklisted. Bidder / manufacturing unit which has been de-recognized/ debarred/banned/blacklisted by State Medical Corporation for any reasons can't participate for that product in the tender

during the period of de-recognition/debarment/banned. If we, at a later date, are found guilty of suppressing facts in this regard, such act on our part shall be considered a fraudulent practice in accordance with the Instructions to Bidders and the Purchaser shall be entitled to reject our BID for the product quoted, submitted by us against this Tender.

1.15. The products offered in the tender must be only manufactured in INDIA

1.16. For the Items quoted in the tender enquiry, firm will have to submit the samples on demand. If firm fails to submit the samples, the tender will be rejected.

1.17. MRP should not be printed on any package.

1.18 All the cold chain items to be transported to HLL warehouse with Data Logger for Cold Chain Maintenance

## 2. COST OF BIDDING

2.1 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 these costs, regardless of the conduct or outcome of the bidding process.

2.2 Tender documents may be downloaded free of cost from the Government e-procurement portal (URL: https://etenders.gov.in/eprocure/app). However, tender document fees, as mentioned in the NIT, is required to be submitted along with the online bid (Not applicable currently due to GO)

## 3. GETTING INFORMATION FROM WEB PORTAL

3.1. All prospective bidders are expected to see all information regarding submission of bid for the Work published in the e tender website during the period from the date of publication of NIT for the Work and up to the last date and time for submission of bid. Non observance of information published in the website shall not be entertained as a reason for any claim or dispute regarding a tender at any stage.

3.2. All bids shall be submitted online on the Government e-procurement portal only in the relevant envelope(s)/ cover(s), as per the type of tender. No manual submission of bids shall be entertained for the tenders published through Government e-procurement portal under any circumstances.

3.3. The Government e-procurement portal shall not allow submission of bids online after the stipulated date & time. The bidder is advised to submit the bids well before the stipulated date & time to avoid any kind of network issues, traffic congestion, etc. In this regard, the department shall not be responsible for any kind of such issues faced by bidder.

## 4. BIDDING DOCUMENTS

4.1. Content of Bidding Documents

The bidding documents shall consists of the following unless otherwise specified

a. Notice Inviting Tender (NIT)
b. General Instruction to Bidders
c. Instructions to Bidders
d. General Conditions of Contract (GCC)
e. Special Conditions of Contract (SCC)
f. Annexures to Bid
g. Product List

4.2. The Bidder is required to login to the e-procurement portal and download the listed documents from the website as mentioned in NIT. He shall save it in his system and

undertake the necessary preparatory work off-line and upload the completed bid at his convenience before the closing date and time of submission.

4.3. The bidder is expected to examine carefully all instructions, Conditions of Contract, Annexures, Terms, Product List in the Bid Document. Failure to comply with the requirements of Bid Document shall be at the Bidder's own risk.

## 5. CLARIFICATION OF BIDDING DOCUMENTS

5.1. A prospective bidder requiring any clarification of the bidding documents shall contact the office of the Tender Inviting Authority on any working day between 10 AM and 5 PM.

5.2 In case the clarification sought necessitates modification of the bid documents, being unavoidable, the Tender Inviting Authority may effect the required modification and publish them in the website through corrigendum.

## 6. AMENDMENT TO BIDDING DOCUMENTS

6.1. Before the deadline for submission of bids, the Tender Inviting Authority may modify the bidding document by issuing addenda.

6.2. Any addendum thus issued shall be a part of the bidding documents which will be published in the e-tender website. The Tender Inviting Authority will not be responsible for the prospective bidders not viewing the website in time.

6.3. If the addendum thus published does involves major changes in the scope of work, the Tender Inviting Authority may at his own discretion, extend the deadline for submission of bids for a suitable period to enable prospective bidders to take reasonable time for bid preparation taking into account the addendum published.

## 7. PREPARATION OF BIDS

### 7.1 Language of the Bid

All documents relating to the bid shall be in the English language.

### 7.2 Documents to be submitted along with the Technical Bid

The online bid submitted by the bidder shall comprise the following:

a) Self-Declaration as per Annexure 1

b) Bid form as per Annexure-2

c) Valid manufacturing license/Factory License/Drug License issued by state drug controller (Self–attested Copy) along with the list of products manufactured in this facility wherever applicable. The quoted products should be highlighted for ready reference.

d) All Pharmaceutical products should have relevant & valid WHO GMP certificate.

e) Firm should submit a non-conviction certificate issued by the State Drug Controller to the effect that the manufacturer has not been convicted under the Drugs and Cosmetics Act, 1940 and rules thereunder during the preceding three years and that no case / proceedings is pending against the manufacturer in any Court of Law in India under the Drugs & Cosmetics Act.

f) Power of attorney for signatory of bid in Rs 200/- stamp paper duly notarized.

g) Copy of GST Certificate (self-attested copy)

h) Copy of Permanent Account Number (Self–attested Copy)

i) Certificate of incorporation and associated documents like Article of Association and Memorandum of Association/Partnership deed/HUF etc as applicable. (Self–attested Copy).

j) Under taking letter for replacement of complaint/defective goods as per Annexure-3

k) List of quoted products with specification Compliance– Annexure 4

l) Suppliers must ensure strict compliance to all statutory regulations, quality standards and packing material specifications (as applicable ) detailed in Annexure – 5

m) Authorization letter from manufacturer (Original) must be submitted as per Annexure 6.

n) Documentary proof attested by Charted Accountant for establishing the average annual turnover of Original Manufacturers having a minimum average annual turnover of Rs.1 Crore (Rupees One Crore only) during the last three years i.e. 2021-2022, 2022-2023 and 2023-2024 (original/ provisional). In case of Authorized agents they must submit the documentary proof for minimum average turnover in the last three years i.e., 2021-2022, 2022-2023 and 2023-2024 (original/ provisional) is Rs.50 lakhs (Rupees Fifty lakhs only) and documentary proof attested by Charted Accountant for establishing their Principal manufacturers meets the eligibility criteria for original manufacturer as specified above. In case of bid by authorized agents, manufacturers authorization form must be attached with the bid submitted.

   If an Original Manufacture is participating in the tender but wishes to make the supplies through its authorised agent, the manufacturer has to ensure that the Authorised minimum average turnover in the last three years i.e., 2021-2022, 2022-2023 and 2023-2024 (original/ provisional) is Rs.50 lakhs (Rupees Fifty lakhs only) and documentary proof attested by Charted Accountant for the same has to be attached.

o) Annexure 7 - Category details of organization, in case of MSE, If the bidder is a MSE, it shall declare in the bid document the Udyog Aadhar Memorandum Number issued to it under the MSMED Act, 2006. If a MSE bidder do not furnish the UAM Number along with bid documents, such MSE unit will not be eligible for the benefits available under Public Procurement Policy for MSEs Order 2012."

p) Duly filled, signed and sealed Annexure 8 - Indemnity Certificate

q) Annexure 10 - Check List

r) Annexure 11 – Compliance To Rule 144 (XI) of GFR 2017 (Self Declaration)

s) Annexure 12 - Make In India Preference (Self Declaration)

t) Annexure 13 - Pre Contract Integrity Pact

u) Annexure 14- Fall Clause Declaration

v) Annexure 15- Bidder Info

w) Annxeue 16- List of Banks integrated with NeSL

x) Annexure 17- Indicative Challan

Note: If any of the above document are not applicable for eligible bidders then they shall attach a "NOT APPLICABLE "statement mentioning the justification for the same.

All Annexures must be dully signed and sealed while submitting the same.

Bidders shall not make any addition, deletion or correction in any of the bid documents. If tampering of documents is noticed during tender evaluation, the bid will be rejected and the bidder will be blacklisted.

**8. Bid Prices**

8.1 The Bidder shall bid as described in the Bill of Quantities.

8.2 The rates quoted by the Bidder shall include cost of the material, freight charges, Insurance or any other charges and applicable GST on **Door delivery basis.**

8.3 The rates and prices quoted by the bidder shall remain firm during the entire period of contract and may be renewed on mutually agreed terms & conditions for a further period.

8.4 Price comparison during evaluation will be done on the Unit basic price of the product excluding GST. The unit basic price of the product shall include cost of the material, freight charges, Insurance or any other charges excluding GST for door delivery basis at **HLL Lifecare Limited, 1 st floor,145, Punjab Expeller Compound, Sihani Chungi,Meerut Road, Ghaziabad, UP-201001,GSTIN: 09AAACH5598K1ZZ ,DL.NO:UP1421B002349, UP1420G000037, UP1420B002371** If a firm quotes NIL Charges/ consideration, the bid for that item(s) shall be treated as unresponsive and will not be considered.

8.5 Rate shall be offered separately for each item as per price schedule. Selection of bidder will be based on the lowest price quoted for each item.

**9. Currencies of Bid and Payment**

9.1. The currency of bid and payment shall be quoted by the bidder entirely in Indian Rupees.

All payments shall be made in Indian Rupees only.

**10. SUBMISSION OF BIDS**

The Bidder shall submit their bid online only through the Government eProcurement portal (URL: https://etenders.gov.in/eprocure/app) as per the procedure laid down for e-submission as detailed in the web site. For e tenders, the bidders shall download the tender documents including the Bill of Quantity (BoQ) file from the portal. The Bidder shall fill up the documents and submit the same online using their Digital Signature Certificate. On successful submission of bids, a system generated receipt can be downloaded by the bidder for future reference. Copies of all certificates and documents shall be uploaded while submitting the tender online.

The tender is invited in 3 **Envelope system** from the registered and eligible firms at CPP Portal.

**a) Envelope - I (Tender Fee and EMD):**

Tender fee (Non-refundable) and EMD as per the tender conditions shall be paid separately, thru RTGS/NEFT transfer in the following HLL A/c details:

| | | |
|---|---|---|
| Name of Bank | : | HDFC BANK |
| A/c number | : | 00630330000605 |
| IFSC Code | : | HDFC0000063 |
| Branch name | : | Vazhuthacaud, Thiruvananthapuram |

Document of the above transactions completed successfully by the bidder, shall be uploaded separately while submitting the bids online.

**NOTE**

- SSI/MSE units interested in availing exemption from payment of Tender Fee and EMD should submit a valid copy of their registration certificate issued by the concerned DIC or NSIC / Udyog Aadhaar.
- If the bidder is a MSE, it shall declare in the bid document the Udyog Aadhar Memorandum Number issued to it under the MSMED Act, 2006.

- If a MSE bidder do not furnish the UAM Number along with bid documents, such MSE unit will not be eligible for the benefits available under Public Procurement Policy for MSEs Order 2012.
- The Party has to provide Performance Security/Security Deposit if Tender is awarded to them.

**b) Envelope - II (Technical bid):** Technical Bid should contain dully filled, signed and scanned soft copy documents as mentioned in Instructions to Bid (ITB) - Documents to be submitted along with the Technical Bid - Section 7.2.

**c) Envelope – III (Financial Bid): The Financial e-Bid through CPP portal:**

All rates shall be quoted in the format provided and no other format is acceptable. If the price bid has been given as a standard format with the tender document, then the same is to be downloaded and to be filled by all the bidders. Bidders are required to download the file, open it and complete the colored (Unprotected) cells with their respective financial quotes and other details (such as name of the bidder). No other cells should be changed. Once the details have been completed, the bidder should save it and submit it online, without changing the filename. If the file is found to be modified by the bidder, the bid will be rejected.

Prices indicated on the Price Schedule shall be entered separately in the following manner:

(i) The Unit basic price of the product shall include cost of the material, freight charges, Insurance or any other charges excluding GST for door delivery basis to our delivery location(s) and the same has to be entered in the Basic Unit rate column of BOQ.

(ii) HSN Code and GST amount as applicable in appropriate column of BOQ.

(iii) The total unit cost in figure and words.

(iv) Prices shall be quoted in Indian Rupees.

(v) If a firm quotes NIL Charges/ consideration, the bid for that item(s) shall be treated as unresponsive and will not be considered.

(vi) If the Tenderer desires to ask for GST to be paid extra, the same must be specifically stated in the allotted column of BoQ. In the absence of any such stipulation or mentioned as zero then the price will be taken inclusive of GST and no claim for the same will be entertained later

(vii) Price comparison during evaluation will be done on the Unit basic price of the product.

(viii) In case bidders quoted different GST amount or percentage for the same item, in such case GST amount ascertained/ decided by the purchaser shall be final

(ix) The need for indication of all such price components by the tenderers, as required in BoQ is for the purpose of comparison of the tenders by the purchaser and will no way restrict the purchaser's right to award the contract on the selected tenderer on any of the terms offered.

(x) Example for illustration purpose

**An example is illustrated below for entering of rate**

*In case of the product number 01 – as per Annexure 4, assume the UOM is Tab for which the rate is applicable. Assume an illustrative value of INR 100 per 10 Tablets*

*which the bidder is planning to quote for the tender. In that case the rate has to be calculated and updated in the BOQ for price bid as INR 10 per tab.*

*Prices indicated on the Price Schedule shall be entered separately in the following manner:*

*The Unit basic price of the product (Rs 10 as per the above example) including freight Charges for inland transportation to HLL Lifecare Limited, 1 st floor,145, Punjab Expeller Compound, Sihani Chungi,Meerut Road, Ghaziabad, UP-201001,GSTIN: 09AAACH5598K1ZZ ,DL.NO:UP1421B002349, UP1420G000037, UP1420B002371.*

*The GST amount of any particular item quoted per piece, as applicable in Value. Note that in the BOQ format, only the value of applicable GST can be entered (percentage of GST cannot be entered).*

*The total unit cost in figure and words. The total unit price will be the basis for evaluation. Note that this will be automatically updated in the BOQ*

Note:-

1. HLL Lifecare Limited reserves the right to verify the credential submitted by the agency at any stage (before or after the award the work). If at any stage, any information / documents submitted by the applicant is found to be incorrect / false or have some discrepancy which disqualifies the firm then HLL shall take the following action:

   a) The agency shall be liable for debarment from tendering in HLL Lifecare Limited, apart from any other appropriate contractual /legal action.

2. On demand of the Tender Inviting Authority, this whole set of certificates and documents shall be send to the Tender Inviting Authority's office address (as given in the NIT) by registered post/Speed post of India Post in such a way that it shall be delivered to the Tender Inviting Authority before the deadline mentioned. The Tender Inviting Authority reserves the right to reject any bid, for which the above details are not received before the deadline.

3. The Tender Inviting Authority shall not be responsible for any failure, malfunction or breakdown of the electronic system while downloading or uploading the documents by the Bidder during the e-procurement process.

**11. Deadline for Submission of the Bids**

11.1 Bid shall be received only online on or before the date and time as notified in NIT.

11.2 The Tender Inviting Authority, in exceptional circumstances and at its own discretion, may extend the last date for submission of bids, in which case all rights and obligations previously subject to the original date will then be subject to the new date of submission. The Bidder will not be able to submit his bid after expiry of the date and time of submission of bid (server time).

Modification, Resubmission and Withdrawal of Bids

11.3. Re submission or modification of bid by the bidders for any number of times before the date and time of submission is allowed. Resubmission of bid shall require uploading of all documents including price bid afresh.

11.4. If the bidder fails to submit his modified bids within the pre-defined time of receipt, the system shall consider only the last bid submitted.

11.5. The Bidder can withdraw his/her bid before the date and time of receipt of the bid. The system shall not allow any withdrawal after the date and time of submission.

## 12. BID OPENING AND EVALUATION

Bids shall be opened on the specified date & time, by the tender inviting authority or his authorized representative in the presence of bidders or their designated representatives who choose to attend.

### 12.1. Bid Opening Process

12.1.1 Opening of bids shall be carried out in the same order as it is occurring in invitation of bids or as in order of receipt of bids in the portal. The bidders & guest users can view the summary of opening of bids from any system. Bidders are not required to be present during the bid opening at the opening location if they so desire.

**Envelope - I:** Envelope- I Opening date shall be as mentioned in NIT Document. (Envelop – I shall contain scanned copy of Tender Fees and EMD).

**Envelope - II:** Opening date shall be as mentioned in NIT. The intimation regarding acceptance / rejection of their bids will be intimated to the contractors/firms through e-tendering portal.

If any clarification is needed from bidder about the deficiency in his uploaded documents in Envelope- I, he will be asked to provide it through CPP portal. The bidder shall upload the requisite clarification/documents within time specified by HLL Lifecare Limited, failing which tender will be liable for rejection. In extraordinary circumstances the bidders may be requested to submit the deficient documents intimated through the e-tendering portal additionally by e-mail (As mentioned in the NIT).

**Envelope - III:** The technically qualified bidders, financial bids shall be opened as per Eligibility Criteria. (Depending on evaluation of Envelop I, the date shall be intimated through CPP Portal)

12.1.2. In the event of the specified date of bid opening being declared a holiday for HLL, the bids will be opened at the same time on the next working day.

### 12.2. Confidentiality

12.2.1. 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 award has been announced in favour of the successful bidder.

12.2.2. Any effort by a Bidder to influence the Purchaser during processing of bids, evaluation, bid comparison or award decisions shall be treated as Corrupt & Fraudulent Practices and may result in the rejection of the Bidders' bid.

### 12.3 Clarification of Bids

12.3.1. To assist in the examination, evaluation, and comparison of bids, the Tender Inviting Authority may ask the bidder for required clarification on the information submitted with the bid. The request for clarification and the response shall be in writing or by e-mail, but no change in the price or substance of the Bid shall be sought, offered, or permitted.

12.3.2. No Bidder shall contact the Tender Inviting Authority on any matter relating to the submitted bid from the time of the bid opening to the time the contract is awarded. If the Bidder wishes to bring additional information to the notice of the Tender Inviting Authority, he shall do so in writing.

**12.4. Examination of Bids, and Determination of Responsiveness**

12.4.1. During the bid opening, the Tender Inviting Authority will determine for each Bid whether it meets the required eligibility as specified in the NIT and the required documents and certificates.

12.4.2. A substantially responsive bid is one which conforms to all the terms, conditions, and requirements of the bidding documents, without material deviation or reservation.

A material deviation or reservation is one:-

- which affects in any substantial way the scope, quality, or performance of the Works;
- which limits in any substantial way, inconsistent with the bidding documents, the Purchaser's rights or the Bidder's obligations under the Contract;

or

- Whose rectification would affect unfairly the competitive position of other Bidders presenting substantially responsive Bids.

12.4.3. If a Bid is not substantially responsive, it may be rejected by the Tender Inviting Authority, and may not subsequently be made responsive by correction or withdrawal of the nonconforming material deviation or reservation.

12.4.4. Non submission of legible or required documents or evidences may render the bid non-responsive.

12.4.5. Bidder can witness the principal activities and view the documents/summary reports for that particular work by logging on to the portal with his DSC from anywhere.

12.4.6. In case only single bid is received, then the purchaser reserves the right to accept/reject the bid as per prevailing norms of GFR and CPP portal, or to go for retender.

**12.5. Negotiation on Bids**

The Tender Inviting Authority reserves the right to negotiate with the lowest evaluated responsive bidder.

**13. BID VALIDITY**

13.1. Bids shall remain valid for the period of **180 (One Hundred And Eighty)** days from the date of opening of the technical bid as specified in the NIT. A bid valid for a shorter period shall be rejected by HLL as non-responsive.

13.2. In exceptional circumstances, prior to expiry of the original bid validity period, the Tendering Authority may request the bidders to extend the period of validity for a specified additional period. The request and the responses thereto shall be made in writing or by email. A bidder may refuse the request without forfeiting its bid security (if applicable). A bidder agreeing to the request will not be required or permitted to modify its bid, but will be required to extend the validity of its bid security (if applicable) for the period of the extension.

**14. STATUTORY EXEMPTIONS:**

- **MSE** - Statutory exemptions as per relevant guidelines shall be applicable for MSE vendors. However, the preferences with respect to MSE shall not be applicable who are only involved the trading of the product under the scope of this tender.
- **PPP MII** - Preferences for Make in India products / services shall be applicable in line with Government Order No.P-45021/12/2017PP (BE-II), 2017 (published by Department for Promotion of Industry and Internal Trade) inclusive of the latest amendments. Self declaration to be submitted to claim MAKE IN INDIA preference.

## 15. BID SECURITY (EMD)

**15(a)**

i) The Bidder shall furnish, as part of his Bid, a Bid Security for an amount as detailed in the Notice Inviting Tender (NIT). For e-tenders, Bidders shall remit the Bid Security using the payment options given in e-tender under Government e-Procurement system only.

ii) Each bid must be accompanied by E.M.D. Any Bid not accompanied by an acceptable Bid Security (EMD) shall be rejected as non-responsive.

iii) The Bid Security (EMD) of the unsuccessful Bidder shall become refundable as promptly as possible after opening of Price Bid and finalization of the tender.

iv) The Bid Security (EMD) of the successful Bidder will be discharged when the Bidder has furnished the required Security Deposit and acceptance of LOI/Work order.

v) SSI/MSE units interested in availing exemption from payment of Bid Security should submit a valid copy of their registration certificate issued by the concerned DIC or NSIC/Udyog Aadhaar. But the Party has to provide Security Deposit, if work is awarded to them.

vi) The bid security may be forfeited/ blacklisted/ de-barred from participating in HLL tenders for a period of 2 years.

vii) The Bid Security may be forfeited:

(a) If a Bidder:

- Changes its offer/bid during the period of bid validity or during the validity of the contract.
- Does not accept the correction of errors

(b) In the case of the successful Bidder, if the Bidder fails:

- To sign the Agreement
- To deliver the material within stipulated time frame as per PO.
- To accept the Notification of award/Letter of Indent/ Purchase order and/or submit the security deposit.
- To acknowledge the Notification of award/Letter of Indent/ Purchase order within 5 days from the date of issue by sending the signed copy of the same.

viii) In such cases the work shall be rearranged at the risk and cost of the selected bidder

ix) The Bid Security deposited will not carry any interest.

## 16. TENDER PROCESSING FEE

16.1. For e-tenders, the mode of remittance of Tender processing Fee shall be the same as detailed for remitting Bid Security (EMD). For e-tenders, Bidders shall remit the Tender fee using the payment options as mentioned in the e-tender

16.2. Any bid not accompanied by the Tender Fee as notified, shall be rejected as nonresponsive.

16.3. Tender Fee remitted will not be refunded.

## 17. ALTERATIONS AND ADDITIONS

17.1 The bid shall contain no alterations or additions, except those to comply with instructions, or as necessary to correct errors made by the bidder, in which case such corrections shall be initialed by the person or persons signing the bid.

17.2 The bidder shall not attach any conditions of his own to the Bid. The Bid price must be based on the tender documents. Any bidder who fails to comply with this clause will be disqualified.

## 18. INDEMNIFICATION CLAUSE

In case of any Adverse Drug Reaction / untoward side effects occurred due to the administration of the product supplied by your organization, the manufacture/ supplier shall be held liable for any legal or any other proceedings initiated by the Government of India / State Government Authorities. The Bidder shall indemnify, defend and hold harmless Government of India and HLL, its Affiliates, officers, directors, employees, agents, and their respective successors and assigns, from and against any and all loss, damage, claim, injury, cost or expenses (including without limitation reasonable attorney's fees), incurred in connection with third Party claims of any kind that arise out of or are attributable to (i) Manufacturer's/Bidders breach of any of its warranties, representations, covenants or obligations set forth herein or (ii) the negligent act or omission of the Manufacturer /Bidders.(iii) any product liability claim arising from the gross negligence or bad faith of, or intentional misconduct or intentional breach of this Contract by bidder or its affiliate. The Bidder has to submit the indemnity certificate duly signed and sealed in the format provided in Annexure 09

## 19. SECURITY DEPOSIT

19.1 Within 3 days of the receipt of notification of award from the purchaser/owner; "The successful Bidder on whom the purchase order / Letter of Intent is placed shall be required to furnish a Performance Guarantee mandatorily only in the form of e-Bank Guarantee (e-BGs) from the Banks which are integrated with NeSL any other mode of Performance Bank Guarantees from such Banks will not be accepted. The details of Banks which are integrated with NeSL is listed at Annexure 16 and Indicative Challan for obtaining e-BG from Bank is at Annexure 17".

19.2 The EMD submitted by the successful bidder shall be converted to Security Deposit and the bidder shall be allowed to remit the balance amount.

19.3 Failure of the successful Bidder to accept the notification of award or submission of security deposit within the time frame shall constitute sufficient grounds for the annulment of the award and forfeiture of the EMD, in which even the purchaser/owner may make the award to the next lowest evaluated bidder or call for new bids.

## 20. PERFORMANCE SECURITY

20.1 An amount of 3% of Basic Price (less GST) shall be deducted from the Invoices submitted by the successful bidder as performance security to be utilized in case of default or defective materials, supplies, work or service not rectified by the bidder. The performance security, less any sums charged by the purchaser, shall be paid over to the bidder after 365 days from the date of receipt of material and acceptance at designated HLL CFA / Depot anywhere in India. The bidder can submit Bank Guarantee (e-BGs) from the Banks which are integrated with NeSL any other mode of Performance Bank Guarantees from such Banks will not be accepted. The details of Banks which are integrated with NeSL is listed at Annexure 16 and Indicative Challan for obtaining e-BG

from Bank is at Annexure 17" , towards the 3% performance security against which the same shall be released.

20.2 After the submission of Performance Guarantee and its acceptance, the Bid Security will be refunded to the successful bidder.

**21. FORFEITURE OF SECURITY DEPOSIT**

If the successful bidder / Contractor fails to supply the ordered material at the rate finalized or execute the work and / or supplies only part quantity / partially execute the work or fails to comply with the terms and conditions of the purchase order / work order the security deposit furnished will be forfeited / Bank Guarantee encashed.

**22. PAYMENT TERMS**

22.1 No Advance payment shall be given.

a. **97% of the payable amount will be released within 120 days** of delivery and acceptance of consignment by HLL

b. **Remaining 3% will be released after 365 days** from the date of receipt of material and acceptance at designated HLL CFA / Depot anywhere in India. The bidder can submit Bank Guarantee (e-BGs) from the Banks which are integrated with NeSL any other mode of Performance Bank Guarantees from such Banks will not be accepted. The details of Banks which are integrated with NeSL is listed at Annexure 16 and Indicative Challan for obtaining e-BG from Bank is at Annexure 17", towards the 3% performance security against which the same shall be released.

22.2 After the submission of Performance Guarantee and its acceptance, the Bid Security will be refunded to the successful bidder.

22.3. The amount shall be paid by HLL in Indian Rupees.

22.4. Acceptance of the payment terms without any qualification shall form part of the technical bid. In case the payment terms are not accepted, the bid is likely to be rejected.

22.5 HLL will make payment to supplier towards the GST amount only after the invoice is uploaded by supplier in GST outward return i.e. GSTR-1 and credit of GST is available (reflected in GSTR-2A) to HLL.

**23. DELIVERY TERMS**

**Considering the Urgency, Goods must be delivered within 7 days from the issue of Notification of Award /Letter of Intent / Purchase order by HLL.**

23.1. **DELAY IN DELIVERY OF GOODS**Delivery of the Goods shall be made by the Supplier in accordance with the time schedule prescribed by the Purchaser in the Notice of award/ Letter of Indent / Purchase order. If at any time during performance of the Contract, the Supplier should encounter conditions impeding timely delivery of the Goods , the Supplier shall promptly notify the Purchaser in writing of the fact of the delay, its likely duration and its cause(s). As soon as practicable after receipt of the Supplier's notice, the Purchaser shall evaluate the situation and may at its discretion extend the Supplier's time for performance, with or without penalty.

If the vendor fails to deliver the full ordered quantity even during extended delivery period then the Notice of award/ Letter of Indent / Purchase order shall be short-closed.

23.2. A delay by the Supplier in the performance of its delivery obligations shall render the Supplier liable to the imposition of penalty pursuant to agreement, unless an extension of time is agreed upon pursuant to agreement without the application of liquidated damages. Levying of penalty shall be on a case to case basis.

23.3. In case of delay in supply the clause number 18 in GCC (Liquidated Damage) will be applicable.

23.4. If L1 defaults (fails to deliver goods on time) then the purchaser reserves the right to purchase the goods from an alternate supplier or from market at the risk and cost of supplier and if the purchase happens at a price higher than the ordered rates, the purchaser shall have the right to claim the difference upon whom order was originally placed and supplier will be under obligation to pay the same. The purchaser has the right to forfeit the performance security / Security Deposit in the event of default. In addition the purchaser is entitled to recover the business loss suffered by the purchaser consequent to default for supplying the product.

**24. TAXES AND DUTIES**

The Bidder shall bear and pay all taxes, duties, levies, GST and charges assessed on the bidder by all municipal, state, or national government authorities, loading & unloading charges etc in connection with the Goods and Services supplied under the Contract. Income Tax and Other Taxes as applicable at the time of execution of job or any other government-imposed liabilities would be deducted from each bill submitted by the bidder

**25. INSPECTION AND TESTS**

26.1 The Purchaser or its representative shall have the right to inspect and/or to test the Goods to confirm their conformity to the Contract at no extra cost of the Purchaser. The Special conditions of Contract and/or the Technical Specifications shall specify what inspections and tests the Purchaser requires and where they are to be conducted. The Purchaser shall notify the Supplier in writing of the identity of any representatives retained for these purposes.

26.2 The inspections and test may be conducted on the premises of the Supplier or at the Goods final destination. Where conducted on the premises of the Supplier or its subcontractor(s), all reasonable facilities and assistance including access to drawings and production data - shall be furnished to the inspectors at no charge to the Purchaser.

26.3 Should any inspected or tested Goods fail to conform to the specifications, the Purchaser may reject them and the Supplier shall either replace the rejected Goods or make all alternations necessary to meet specification requirements free of cost to the Purchaser.

26.4 The Purchasers right to inspect, test and, where necessary, reject the Goods' arrival in at any site shall in no way be limited or waived by reason of the Goods having previously been inspected, tested and passed by the Purchaser or its representative prior to the Goods dispatched.

26.5 HLL reserves the right to seek samples of the product being offered before placement of order and based on approval of samples by HLL/Ultimate customer the order shall be placed. If the sample is rejected due to quality/technical reasons, HLL reserves the right to approach the next higher bidder for samples and if approved, HLL shall proceed with order placement with the next higher bidders. The samples approved only be accepted against the order placed and any deviation would result in the rejection of the product supplied.

26.6 The supplier should submit the internal lab reports for the supplies made to HLL. The purchaser reserves the right to sample check the consignment at the time of delivery for which cost shall be borne by the supplier (pre-dispatch inspection). HLL may analyse the sample drawn from the goods received at depots/C&FAs. In case of sample testing failure at third party lab/ HLL's lab or quality related market complaints, the supplier shall take sole responsibility to replace the entire batch free of cost including the freight charges for collecting back the rejected items from HLL warehouses & resupply or

refund the payment for such rejected quantity equal to its Door delivery value if the payment is already made.

**26. INDEMNITY**

The Bidder shall indemnify, defend and hold harmless Government of India and HLL, its Affiliates, officers, directors, employees, agents, and their respective successors and assigns, from and against any and all loss, damage, claim, injury, cost or expenses (including without limitation reasonable attorney's fees), incurred in connection with third Party claims of any kind that arise out of or are attributable to (i) Manufacturer's/Bidders breach of any of its warranties, representations, covenants or obligations set forth herein or (ii) the negligent act or omission of the Manufacturer /Bidders.(iii) any product liability claim arising from the gross negligence or bad faith of, or intentional misconduct or intentional breach of this Contract by bidder or its affiliate. The Bidder has to submit the indemnity certificate duly signed and sealed in the format provided in Annexure 9

**27. SHORT SUPPLY:**

If any shortages in sealed boxes are detected, then supplier should be held responsible. In such a case, the supplier will have to make good of the loss or refund the payment for such quantity equal to its purchase value if the payment is already made. If the payment is not made, purchaser will have right to deduct the payment for the equivalent purchase value corresponding to quantity found short**.**

**28. PARALLEL RATE CONTRACTS:**

HLL reserves the right to enter into the rate contract / parallel rate contracts with one or more parties or to place adhoc contracts simultaneously or at any time during the currency of contract, with one or more suppliers.

The purchaser also reserves the rights (1) to enter into parallel Price Agreement(s) / Contract(s) simultaneously or at any time during the period of the Price Agreement/Rate Contract with one or more bidder(s) as he/they think fit and (2) to place adhoc contract or contracts simultaneously or at any time during the period of this Rate contract with one or more supplier(s) / bidder(s) for such quantity of such item or items as the purchaser (whose decision shall be final) may determine.

**29. IN CASE OF DEFAULT**

The purchaser is not bound to accept the L1 offer only and circumstances warranting where L1 shows its disinterest, L2 or higher offer may be considered for acceptance.

**30. RISK PURCHASE**

If L1 or any other parties' defaults (fails to deliver goods on time) then the purchaser reserves the right to purchase the goods an alternate supplier or from market at the risk and cost of L1 supplier and if the purchase happens at a price higher than the ordered rates, the purchaser shall have the right to claim the difference upon whom order was originally placed and L1 supplier will be under obligation to pay the same. In addition, the purchaser is entitled to recover the business loss suffered by the purchaser consequent to default for supplying the product.

**31. FORCE MAJEURE**

32.1 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 limited to, acts of the Purchaser either in its sovereign or contractual capacity, wars or revolutions, fires, floods, epidemics, quarantine restrictions and freight embargoes.

32.2 If a Force Majeure situation arises, the Supplier shall promptly notify the Purchaser in

writing within Seven days from the date of such conditions 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.

**32. GOODS REPLACEMENT:**

If goods are found to be defective during the sample testing by HLL or Quality related market complaint, on arrival of the material at designated HLL delivery point, supplier must replace the quantity free of cost with fresh batch upon demand by HLL. However replacement of goods will be accepted by HLL subject to the concurrence from the ordering institute else the purchase order will be cancelled and Clause 24 (Delay in delivery of goods) will be applied under the discretion of HLL.

**33. CLARIFICATIONS ON BIDS**

During the bid evaluation, HLL 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 price or substance of the bid shall be sought, offered, or permitted

**34. CONTACTING HLL**

a) From the time of bid opening to the time of Contract award, if any Bidder wishes to contact HLL on any matter related to the bid, he shall do so in writing by sending email to sdrbdsouth@lifecarehll.com.

b) If a Bidder tries to influence HLL directly or otherwise, interfere in the bid evaluation process and the Contract award decision, his bid will be rejected.

**35. HLL'S RIGHT TO ACCEPT OR REJECT ANY OR ALL BIDS**

The Purchaser reserves the right to accept or reject any bid, and to annul the bidding process and reject all bids at any time prior to award Contract award, without thereby incurring any liability to the affected bidder or bidders.

The purchaser does not bind itself to accept the lowest or any bid and reserves the right to reject any or all bids at any point of time prior to the issuance of the Notice of award/Letter of intent/Purchase order without reason whatsoever.

The purchaser reserves the right to resort to retendering without providing any reasons whatsoever. The purchaser shall not incur any liability on account of such rejection.

The purchaser reserves the right to modify any terms, conditions or specifications for submission of offer and to obtain revised bids from the bidders due to such changes, if any.

Canvassing of any kind will be a disqualification and the purchaser may decide to cancel the bidder from its empanelment.

The purchaser reserves the right to accept or reject any bid and annul the bidding process and reject all bids at any time prior to award of contract without thereby incurring any liability to the affected bidder or bidders or any obligation to inform the affected bidder or bidders of the ground for the purchaser's action.

**36. PURCHASER'S RIGHT TO VARY QUANTITIES AT TIME OF AWARD**

The Purchaser reserves the right at the time of award of contract to increase or decrease the quantity of goods and services originally specified in the bid document without any change in unit price or other terms and conditions.

**37. EVALUATION AND COMPARISON OF BIDS**

38.1 The Purchaser will evaluate and compare bids previously determined to be substantially

responsive.

38.2 Price comparison during evaluation will be done on the Unit basic price of the product excluding GST. The unit basic price of the product shall include cost of the material, freight charges, Insurance or any other charges excluding GST for door delivery basis to our delivery location(s)

38.3 Rate shall be offered separately for each item as per price schedule. Selection of bidder will be based on the lowest price quoted for each item.

**38. SETTLEMENT OF DISPUTES**

Arbitration shall not be a means of settlement of any dispute or claim arising out of the contract relating to the work. Any disputes or difference arising between the parties with respect to the performance of any part of this agreement or anything connected therewith, etc shall as far as possible be mutually settled by the process of dialog and negotiation. Any disputes or differences or questions or claims arising under or relating to a concerning or touching this agreement shall be referred for arbitration in accordance with the provisions of the Arbitration and Conciliation Act 1996.

The arbitration proceedings shall be held at Thiruvananthapuram. The award passed by the arbitrator shall be final and binding on the parties hereto. The conduct of such arbitration shall be in English. Subject to arbitration, the Courts at Thiruvananthapuram alone shall have jurisdiction in respect of settlement of any matter arising out or in connection with the contract.

**39. MAJOR RESPONSIBILITIES OF SUPPLIER**

a. The suppliers have to supply the goods as per the delivery schedules and quantity mentioned in the Notification of award/ Letter of Indent/ Purchase order. Supplies made shall be in strict conformance with the stipulations of tender specification and the respective Notification of award/ Letter of Indent/ Purchase orders.

b. The successful bidder shall acquire in its name all permits, approvals, and/or licenses from all local, state, or national government authorities or public service undertakings that are necessary for the performance of the Notification of award/ Letter of Indent/ Purchase order.

c. The Supplier shall comply with all laws in force in India. The laws will include all national, provincial, municipal, or other laws that affect the performance of the Contract and are binding upon the bidder. The Bidders shall indemnify and hold harmless HLL from and against any and all liabilities, damages, claims, fines, penalties, and expenses of whatever nature arising or resulting from the violation of such laws by the bidder or its personnel except that caused by HLL.

d. Any product related legal issues shall be handled and connected expenses therewith shall be borne by the bidder/ manufacturer only.

e. Any product related cases shall be handled and connected expenses therewith shall be borne by the contract manufacturer only

f. The bidder must undertake to provide the purchaser the consignment number (s) by which the items ordered had been dispatched from their sites, so as to have online/web access to the tracking system of physical movements of the consignments sent through the courier.

g. The supplier should submit the internal lab reports for the supplies made to HLL. The purchaser reserves the right to sample check the consignment at the time of delivery for which cost shall be borne by the supplier (pre-dispatch inspection). HLL may analyse the sample drawn from the goods received at depots/C&FAs. In case of sample testing failure

at third party lab/ HLL's lab or quality related market complaints, the supplier shall take sole responsibility to replace the entire batch free of cost including the freight charges for collecting back the rejected items from HLL warehouses & resupply or refund the payment for such rejected quantity equal to its Door delivery value if the payment is already made.

**40. The final quantities mentioned in Annexure 4 may vary as per the final requirement and the order may be placed in single or multiple lots during the bid validity period.**

**41. GOVERNING LANGUAGE**

The contract shall be written in English language. English language version of the Contract shall govern its interpretation. All correspondence and documents pertaining to the Contract which are exchanged by the parties shall be written in the same language.

**42. AWARD CRITERIA**

The Purchaser will award the contract with the successful bidders whose bid has been determined to be substantially responsive and has been determined as the lowest evaluated bid in the respective price slabs, provided further that the bidder is determined to be qualified to perform the contract satisfactorily.

**43. NOTIFICATION OF AWARD**

44.1 Prior to the expiration of the period of bid validity, the Purchaser will notify the successful bidder in writing by registered letter or by email, to be confirmed, that its bid had been accepted.

44.2 The notification of award will constitute the formation of the contract.

44.3 The notification of award/ Letter of Intent/ Purchase order will constitute the formation of the Contract. The supplier shall give acceptance of the Notification of award/Letter of Intent/ Purchase order within 5 days from the date of issue by sending the signed copy of the same failing which ,the purchaser shall have the right to cancel the order. The conditions mentioned in the Notification of award/Rate contract agreement/Letter of Intent/ Purchase order will be mutually binding for both the parties and the bidder and the purchaser shall abide by the same. In case of any default in any of the condition of the Notification of award/Letter of Intent/ Purchase order, the purchaser reserves the rights to invoke Bid Securing clause.

44.4 The Purchase order (PO) / Notice of award is liable to be cancelled, if the supplier is unable to comply with or violates any of the terms and conditions laid down in the Purchase order/ Notice of Award. Therefore, up on such cancellation of PO/ Notice of award by HLL, the Supplier will be liable to refund the outstanding advance amount forthwith.

44.5 The successful bidder shall confirm the acceptance of the Notice of award/Purchase order as per the terms & conditions of the tender by signing and returning the duplicate copy of Purchase order (PO)/Notice of award within 5 days from the date of issue of the of purchase order/ Notice of award, failing which HLL shall have the right to reject the purchase order/ Notice of award.

**44. TERMINATION**

HLL reserve right to terminate/ cancel the Notification of award/ Letter of Indent/ Purchase order at any time for any reason without any liability on HLL.

**45. FALL CLAUSE**

The BIDDER undertakes that it has not supplied/is not supplying similar product/systems or subsystems OR providing similar services at a price/ charge lower than that offered in the present bid in respect of any other Ministry/Department of the Government of India or

PSU and if it is found any stage that similar product/systems or sub systems was supplied by the BIDDER to any to the Ministry/Department of the Government of India or a PSU at a lower price, then that very price, with due allowance for elapsed time will be applicable to the present case and the difference in the cost would be refunded by the BIDDER to HLL, if the contract has already been concluded.

## 46. CORRUPT OR FRAUDULENT PRACTICES

The purchaser requires that the bidders, suppliers and contractors observe the highest standard of ethics during the procurement and execution of such contracts. In pursuit of this policy, the following are defined:

| Sl. No | Term | Meaning |
|---|---|---|
| (a) | Corrupt practice | The offering, giving, receiving, or soliciting, directly or indirectly, of anything of value to influence the action of a public official in the procurement process or in contract execution. |
| (b) | Fraudulent practice | A misrepresentation or omission of facts in order to influence a procurement process or the execution of a contract. |
| (c) | Collusive practice | Means a scheme or arrangement between two or more bidders, with or without the knowledge of the purchaser, designed to establish bid prices at artificial, non-competitive levels. |
| (d) | Coercive practice | Means harming or threatening to harm, directly or indirectly, persons or their property to influence their participation in the procurement process or affect the execution of a contract. |

The Purchaser will reject the proposal for award if it determines that the Bidder recommended for award has, directly or through an agent, engaged in corrupt, fraudulent, collusive or coercive practices in competing for the Contract in question.

## 47. SHELF LIFE

The supplies of all products should be from fresh stock only. **At the time of receipt of medicines at HLL depot, the products offered should have products should have minimum 70% of remaining shelf life with latest manufacturing date** Products to be supplied should be of standard quality/quantity as per specification and must be as per the formulations/standard approved/specified by the Drug Control Act and Food & Drug Control Administration Regulation or as per the regulation of any such statutory authorities..

## 48. FLEXIBILITY OF PRICES

The purchaser has option to re-negotiate with rate contract holder to bring down the rate contract prices whenever market fluctuations affect the prices abnormally.

## 49. LICENSE AND PERMITS

The Supplier shall acquire in its name all permits, approvals, and/or licenses from all local, state, or national government authorities or public service undertakings that are necessary for the performance of the Contract.

The Supplier shall comply with all laws in force in India. The laws will include all national, provincial, municipal, or other laws that affect the performance of the Contract and are binding upon the Supplier The Supplier shall indemnify and hold harmless Purchaser from and against any and all liabilities, damages, claims, fines, penalties, and expenses of whatever nature arising or resulting from the violation of such laws by the Supplier or its personnel.

## 50. INTEGRITY PACT

Pre-Contract Integrity Pact and Independent External Monitor

The Integrity pact annexed shall be part and parcel of this document, and has to be signed by bidder(s) at the pre-tendering stage itself, as a pre bid obligation and should be submitted along with the financial and technical bids. All the bidders are bound to comply with the Integrity Pact clauses. Bids submitted without signing Integrity Pact will be ab initio rejected without assigning any reason.

The Integrity pact annexed shall be part and parcel of this document, and has to be signed by bidder(s) at the pre-tendering stage itself, as a pre-bid obligation and should be submitted along with the financial and technical bids. All the bidders are bound to comply with the Integrity Pact clauses. Bids submitted without signing Integrity Pact will be ab initio rejected without assigning any reason.

The email id of the Independent External Monitor for HLL is given below.

Email id: iemhll@lifecarehll.com

## 51. RESTRICTIONS UNDER RULE 144 (XI) OF GFR 2017 FOR BIDDERS FROM A COUNTRY SHARING LAND BORDER WITH INDIA.

Any bidder from a country which shares a land border with India will be eligible to bid in this tender only if the bidder is registered with Competent Authority, as per order no F.No.6/18/2019-PPD dated 23-July-2020 (Rule 144 (xi) of GFR) inclusive of the latest amendments issued by Ministry of Finance, GOI at Appendix of this bidding document. The bidder must comply with all provisions mentioned in this order. A self-declaration (as per format provided in Annexure 12) with respect to this order must be submitted.

## 52. PURCHASE PREFERENCE TO MICRO AND SMALL ENTERPRISES (MSE's)

Purchase preference will be given to MSEs as defined in Public Procurement Policy for Micro and Small Enterprises (MSEs) Order, 2012 dated 23.03.2012 issued by Ministry of Micro, Small and Medium Enterprises and its subsequent Orders/Notifications issued by concerned Ministry. However, the preferences with respect to MSE shall not be applicable who are only involved the trading of the product under the scope of this tender.

## 53. PROVISIONS OF PUBLIC PROCUREMENT (PREFERENCE TO MAKE IN INDIA) ORDER 2017

Statutory exemptions as per relevant guidelines shall be applicable for MSE vendors. Preferences for Make in India products / services shall be applicable in line with Government Order No.P-45021/12/2017PP (BE-II), 2017 (published by Department for Promotion of Industry and Internal Trade) inclusive of the latest amendments. Self-declaration to be submitted to claim MAKE IN INDIA preference as per Annexure 15.

## 54. SPLITTING OF ORDER

In case of critical/vital/safety/security nature of the item, large quantity under procurement, urgent delivery requirements and inadequate vendor capacity, HLL reserves the right to split the contract quantity between the bidders. The splitting ratio shall be at the discretion of HLL. The lowest rate accepted would be counter offered to the L2 party. On acceptance of the counter offer, the order will be placed on L2 for the respective percentage. In case of non-acceptance of the counter offer by the L2 party, a similar offer shall be made to L3 and L4, and so on.

## 55. MRP should not be printed in any package

**56. Goods and Services Tax (GST) :**

a. If a tenderer asks for Goods and Services Tax to be paid extra, the rate and nature with HSN code of Goods and Services Tax applicable should be correctly shown separately. The Goods and Services Tax will be paid as per the rate at which it is liable to be assessed or has actually been assessed provided the transaction is legally liable to Goods and Services Tax and is payable as per the terms of the contract. If any refund of Tax is received at a later date, the Supplier must return the amount forth-with to the purchaser.

b. In case within the delivery period stipulated in the contract, there is an increase in the statutory taxes like GST or fresh imposition of taxes which may be levied in respect of the goods and services specified in the contract, reimbursement of these statutory variation shall be allowed to the extent of actual quantum of taxes paid by the supplier. This benefit, however, cannot be availed by the supplier in case the period of delivery is extended due to unexcused delay by the supplier.

c. But nevertheless, the Purchaser shall be entitled to the benefit of any decrease in price on account of reduction in or remission of GST or any other duty or tax or levy or on account of any other grounds. In case of downward revision in taxes/duties, the actual quantum of reduction of taxes/duties must be reimbursed to the purchaser by the supplier. All such adjustments shall include all reliefs, exemptions, rebates, concession etc. if any obtained by the supplier.

# GENERAL CONDITIONS OF CONTRACT (GCC)

## 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 the attachments and appendices thereto and all documents incorporated by reference therein;

(b) "The Contract Price" means the price payable to the Supplier under the Contract for the full and proper performance of its contractual obligations;

(c) "The Goods" means all the products, and/or other materials which the Supplier is required to supply to the Purchaser under the Contract;

(d) "Services" means services ancillary to the supply of the Goods, such as transportation and insurance, and other incidental services, covered under the contract;

(e) "GCC" means the General Conditions of Contract contained in this section.

(f) "SCC" means the Special Conditions of Contract.

(g) "The Purchaser" means the Organisation purchasing the Goods, as named in SCC;

(h) "The Supplier" means the individual or firm supplying the Goods under this Contract;

(i) "Day" means calendar day.

(j) "Delivery period" means the period applicable upto completion of supply of goods by the supplier at the required site mentioned in Notification of award/ Letter of Indent/ Purchase order and accepted by the Purchaser.

## 2. APPLICATION

2.1 These General Conditions shall apply to the extent that they are not superseded by provisions in other parts of the Contract.

## 3. STANDARDS

3.1 The Goods supplied under this Contract shall conform to the standards mentioned in the Technical Specifications.

## 4. USE OF CONTRACT DOCUMENTS AND INFORMATION

4.1 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 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.

4.2 The Supplier shall not, without the Purchaser's prior written consent, make use of any document or information enumerated in GCC Clause 4.1 except for purposes of performing the Contract.

4.3 Any document, other than the Contract itself, enumerated in GCC clause 4.1 shall remain the property of the Purchaser and shall be returned (in all copies) to the Purchaser on completion of the supplier's performance under the Contract if so required by the Purchaser.

## 5. SUBCONTRACTS

The supplier shall notify the Purchaser in writing of all subcontracts awarded under the contract if not already specified in his bid. Such notification, in his original bid or later, shall not relieve the Supplier from any liability or obligation under the contract.

**6. CONTRACT AMENDMENTS**

6.1 Subject to GCC Clauses, no variation in or modification of the terms of the Contract shall be made except by written amendment signed by the parties.

**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.

7.2 Any product related cases shall be handled and connected expenses therewith shall be borne by the Supplier only.

**8. INSURANCE**

For delivery of goods at site, the insurance shall be obtained by the Supplier in an amount equal to 110% of the value of the goods from "Warehouse to Warehouse" (Final destinations) on "All Risks" basis including War Risks and Strike.

**9. CHANGE ORDERS**

9.1 The Purchaser may at any time by written order given to the Supplier, make changes within the general scope of the Contract in any one or more of the following:

(a) The method of shipping or packing
(b) The place of delivery; or
(c) The services to be provided by the Supplier.

**10. ASSIGNMENT**

10.1 The Supplier shall not assign, in whole or in part, its obligations to perform under the contract, except with the Purchaser's prior written consent.

**11. TERMINATION BY DEFAULT**

11.1 The Purchaser may, without prejudice to any other remedy for breach of contract, by written notice of default sent to the supplier, terminate the Contract in whole or part;

(a) if the Supplier fails to deliver any or all of the goods within the time period(s) specified in the Contract, or within any extension thereof granted by the Purchaser, or

(b) If the Supplier fails to perform any other obligation(s) under the contract.

11.2 In the event the Purchaser terminates the Contract in whole or in part, 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. However, the Supplier shall continue the performance of the Contract till such time.

**12. TERMINATION FOR INSOLVENCY**

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 which has accrued or will accrue thereafter to the Purchaser.

**13. APPLICABLE LAW**

The Contract shall be interpreted in accordance with the laws of the Union of India.

## 14. NOTICES

14.1 Any notice given by one party to the other pursuant to this Contract shall be sent to other party in writing or by cable, telex or facsimile and confirmed in writing to the other Party's address specified in Special Conditions of Contract.

14.2 A notice shall be effective when delivered or on the notice's effective date, whichever is later.

## 15. TAXES AND DUTIES

Supplier shall be entirely responsible for all taxes, duties, license fees, octroi etc., incurred until delivery of the contracted Goods to the Purchaser.

## 16. PACKING

16.1 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. Packing shall adhere to conditions stipulated in Technical specification.

16.2 The packing, marking and documentation within and outside the packages shall comply strictly with such special requirements as shall be provided for in the Contract including additional requirements, if any, specified in SCC and in any subsequent instructions ordered by the Purchaser

## 17. DELIVERY AND DOCUMENTS

Delivery of the Goods shall be made by the Supplier in accordance with the terms specified by the Purchaser in the Letter of Indent / Notification of Award / Purchase order. The details of dispatching and/or other documents to be furnished by the Supplier are specified in SCC, if any.

## 18. LIQUIDATED DAMAGES

If the Supplier fails to deliver any or all of the Goods or perform of services within the time period(s) specified in the Contract, the Purchaser shall without prejudice to its other remedies under the Contract, deduct from the Contract Price, as liquidated damages, a sum equivalent to 0.5 percent of the delivered price of the delayed Goods or unperformed Services for each week of delay or part thereof until actual delivery or performance, up to a maximum deduction of 10 percent of the delayed Goods or Services contract price. Service tax as applicable will also be recovered in addition to the liquidated damages. However, H.L.L at its sole discretion reserves the right to accept or reject the delivery of materials which are supplied beyond the delivery date as mentioned in the purchase order. In the event of H.L.L accepting the delivery of the materials beyond the stipulated delivery date as per the Purchase order, penalty as mentioned above would apply. In the event of H.L.L rejecting the delivery of the materials beyond the stipulated delivery date as per the Purchase order, then the party is liable to repay HLL any advance amount which was paid by HLL, failing which HLL will have the right to initiate legal proceedings against such party/ successful bidder. Once the maximum is reached, the Purchaser may consider termination of the Contract. If the Supplier fail to comply with specific packing descriptions or instructions, the loss incurred by the purchaser on this account shall be indemnified by the supplier.

## 19. RESOLUTION OF DISPUTES

19.1 The Purchaser and the Supplier shall make every effort to resolve amicably by direct informal negotiation any disagreement or dispute arising between them under or in connection with the Contract.

19.2 If, after thirty (30) days from the commencement of such informal negotiations, the Purchaser and the Supplier have been unable to resolve amicably a Contract dispute, either party may require that the dispute be referred for resolution to the formal mechanisms specified in the Special Conditions of Contract. These mechanisms may include, but or not limited to, conciliation mediated by a third Party, adjudication in an agreed national forum, and national arbitration.

## Special Conditions of Contract (SCC)

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

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

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

**There are no special conditions or contract for this tender and all other conditions mentioned in other sections stands valid.**

**Annexure-01**

**SELF - DECLARATION**

**Supply of Pharmaceutical Products for Onward Supplies to Foreign Country**

**Tender No. HLL/PSD/RBD/2025-26/TENDER/06**

To,
Vice President (PS) and GH (HCS)
Procurement Services Division
HLL Lifecare Limited
Corporate & Regd. Office
HLL Bhavan, Poojappura,
Thiruvananthapuram-695012
E-mail: sdrbdsouth@lifecarehll.com

Dear Sir,

We certify that We or our Principal Manufacturer (if applicable), have not been de-registered or debarred or blacklisted or banned / suspended for business for any product or constituent of the product we have quoted, by State Government / Central Govt. Organization /State Medical Corporations/ Director Health Services and or convicted by any court of law, till the due date of submission of BID as specified in the subject BID. If we, at a later date, are found guilty of suppressing facts in this regard, such act on our part shall be considered a fraudulent practice in accordance with the Instructions to Bidders and the Purchaser shall be entitled to reject our BID for the de-registered or debarred or blacklisted or banned / suspended product quoted, submitted by us against this Tender.

Also certify that the quoted products possess relevant quality assurance certification issued by the concerned authorities for all the offered products.

We hereby guarantee that the drugs supplied by our company are not spurious and we further guarantee not to supply any sub-standard or spurious drugs. We assure that the drugs/medicines to be supplied shall be as per the formulations / standard approved / specified by the Drug Control Act and Food & Drug Control Administration Regulation or as per the regulation of any such statutory authorities.

We have also noted that after submission of BID and before award contract, if we are deregistered or debarred or blacklisted by State Government or Government of India / Drug Controller, our BID will be considered as Non-responsive.

We hereby declare that the facts furnished for the purpose of this tender are correct and true to the best of our knowledge. We are well aware that any discrepancy in the same makes us liable for disqualification / debarment / appropriate action by the tenderer.

Date:
Place:

Signature:
Name:

Designation:

Seal:

**BID FORM** **Annexure-02**

Ref: Date:

To,

Vice President (PS) and GH (HCS)
HLL Lifecare Limited,
HLL Bhavan, Poojappura,
Thiruvananthapuram -695012 Kerala, India
Tel: 0471 2775500, 0471 2350959 (EXTN - 606 /531)
Website – www.lifecarehll.com

Dear Sir,

**Supply of Pharmaceutical Products for Onward Supplies to Foreign Country**

**Tender No. HLL/PSD/RBD/2025-26/TENDER/06**

Having examined the Bidding Documents, including Addenda Nos. [insert numbers], the receipt of which is hereby acknowledged, we, the undersigned, offer our services in full conformity with the Bidding Documents for the total amount against the Product as indicated in the price Schedule.

We undertake that in case our bid is accepted, we shall commence work and shall make all reasonable endeavor to achieve contract acceptance.
We agree to abide by this bid, which, in accordance with consists of this letter, the Price Schedule, letter of authorization, documents establishing conformity, and Attachments through [specify: the number of attachments] to this Bid Form, up to 12 months from the date of opening of financial bids and it shall remain binding upon us and may be accepted by you at any time before the expiration of that period.

We declare that the above quoted price for product is firm and shall not be subject to any variation for the entire period of the assignment. We further declare that the above quoted prices include all taxes as on the date of bid submission, duties and levies payable by us under aforesaid assignment.

We declare that price/ rate offered is for pharmaceutical products at HLL Depot Ghaziabad and all other related activities.

The costs of withdrawals of these deviations / exclusions are enclosed with the Price Schedule. In case a formal final Contract is not prepared and executed between us, this bid, together with your written acceptance of the bid and your notification of award, shall constitute a binding contract between us. We understand that you are not bound to accept the lowest or any bid you may receive.

We, the Bidder shall indemnify, defend and hold harmless Government of India, HLL,its Affiliates, officers, directors, employees, agents, and their respective successors and assigns, from and against any and all loss, damage, claim, injury, cost or expenses (including without limitation reasonable attorney's fees), incurred in connection with third Party claims of any kind that arise out of or are attributable to (i) Manufacturer's/Bidders breach of any of its warranties, representations, covenants or obligations set forth herein or (ii) the negligent act or omission of the Manufacturer/Bidders. (iii) any product liability claim arising from the gross negligence or bad faith of, or intentional misconduct or intentional breach of this Contract by bidder or any affiliate.

We agree to all terms and conditions of the Bid Document and subsequent amendments.

Dated this [insert: number] day of [insert: month], [insert: year].
Signature.....
Name.....
Full Address with contact person Name, Phone number and Email
Designation and Common Seal...

**Annexure-03**

**UNDERTAKING LETTER FOR REPLACEMENT OF COMPLAINT/DEFECTIVE GOODS**

**Supply of Pharmaceutical Products for Onward Supplies to Foreign Country**

**Tender No. HLL/PSD/RBD/2025-26/TENDER/06**

To,
Vice President (PS) and GH (HCS)
HLL Lifecare Limited,
HLL Bhavan, Poojappura,
Thiruvananthapuram -695012 Kerala, India
Tel: 0471 2775500, 0471 2350959 (EXTN - 606 /531)
Website – www.lifecarehll.com

Dear Sir,

We hereby confirm and assure you, that the products supplied by us will meet all the quality standards and even if any quality complaint arises, we (name-----) take the responsibility to take back the complaint batches and replace and deliver fresh batch to HLL sores/ warehouse free of cost within 30 days, subject to approval from HLL. We (name----) shall also bear the transportation charges for collecting back the compliant/rejected batches or goods and the transportation charges incurred for making the replacement.

Signature_____

Name_____

Designation and Common Seal

Station_____
Date_____

**Annexure-04**

## List of quoted products with specification Compliance

## Supply of Pharmaceutical Products for Onward Supplies to Foreign Country

**TENDER No – HLL/PSD/RBD/2025-26/TENDER/06 Dated 25.04.2025**

| Sl.No | Item | Specification | UOM | Required quantity | Quoted (Yes/No) | Manufactured by | Offered Pack Size | 100% Technically Complied (Yes / No) |
|---|---|---|---|---|---|---|---|---|
| 1 | Cefotaxime for Inj. 1g Vial | Cefotaxime for Injection BP 1g OR Cefotaxime for Injection USP 1g. Each 1g vial to contain sterile Cefotaxime Sodium BP/USP equivalent to 1g of Cefotaxime as sterile dry powder for reconstitution with Water for Injection BP/USP for intramuscular and intravenous use. Note: 1.This injection should be stable for minimum of 24 months when stored at a temperature range of 28'C-32'C 2.Each vial should be labelled accordingly. | VIAL | 6,60,000 | | | | |
| 2 | Ceftazidime for Inj. 1g Vial | Ceftazidime for injection USP/BP 1g. Each vial to contain Ceftazidime Pentahydrate equivalent to 1g of Ceftazidine USP/BP as a sterile dry powder for reconstitution with suitable diluent for intramuscular and intravenous use. Note : 1.This injection should be stable for a minimum of 02 years when stored at a temperature not exceeding 25'C. 2.Each vial should be Labelled accordingly. | VIAL | 1,80,000 | | | | |
| 3 | Ceftriaxone for Inj. 1g Vial | Ceftriaxone for injection BP/USP 1g Each vial to contain sterile Ceftriaxone Sodium BP/USP equivalent to 1g of Ceftriaxone as sterile dry powder for reconstitution with water for injection BP/USP for intravenous injection Note: 1.This injection should be stable for a minimum of 02 years when stored under temperature not exceeding 30'C 2.Each vial should be labelled accordingly | VIAL | 15,00,000 | | | | |

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|---|---|---|---|---|---|---|---|---|
| 4 | Imipenem+ Cilastatin for Inj.500mg | Imipenem 500mg and Cilastatin 500mg Injection USP/IP Each vial to contain mixture of 500 mg of Imipenam USP/IP, 500 mg of Cilastatin Sodium USP/IP and Sodium Bicarbonate USP/IP as sterile dry powder for reconstitution for intramuscular and intravenous injection or intravenous infusion. Note: 1.The product should be stable for 02 years when stored under controlled room temperature. 2.Each vial should be labelled accordingly. | VIAL | 60,000 | | | | |
| 5 | Doxycycline Cap. 100mg | Doxycycline Capsule BP/USP 100mg Each capsule to contain Doxycycline Hydrochloride BP/USP equivalent to 100mg of Doxycycline. Note: The shelf life of the product should be minimum of 24 months. | CAP | 68,00,000 | | | | |
| 6 | Amikacin Inj. 500mg/2ml Vial | Amikacin Sulphate Injection USP 500mg in 2ml vial OR Amikacin Injection BP 500mg in 2ml vial Each vial to contain 500mg of Amikacin Sulphate USP/BP in water for injection BP/USP for Intravenous or Intramuscular Injection. Note: 1.This injection should be stable for 02 years when stored at a temperature range of 28'C-32'C 2.Each vial should be labelled accordingly | VIAL | 2,56,000 | | | | |
| 7 | Vancomycin Inj. 500mg Vial | Vancomycin Injection USP 500mg/ Vancomycin Hydrochloride for Injection USP 500mg Each 10ml vial to contain 500mg of Vancomycin Hydrochloride USP as sterile dry powder for reconstitution with water for injection BP/USP, before use for intravenous infusion. OR Vancomycin Injection/Infusion BP 500mg/ Vancomycin Hydrochloride for Infusion BP 500mg Each 10ml vial to contain 500mg of Vancomycin Hydrochloride BP as sterile dry powder for reconstitution with water for injection BP/USP, for intravenous infusion. Note: 1.This injection should be stable for a minimum of 02 years when stored below 25'C 2.Each vial should be labeled accordingly. | VIAL | 1,40,000 | | | | |

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|---|---|---|---|---|---|---|---|---|
| 8 | Teicoplanin Inj. 400mg Vial | Teicoplanin Injection 400mg vial Each vial to contain 400mg of Teicoplanin as sterile dry powder for reconstitution for intramuscular or intravenous use Note: 1.This injection should be stable for minimum of 24 months when stored at a temperature which is mentioned by the manufacturer. 2.Each vial should be labelled accordingly. | VIAL | 78,000 | | | | |
| 9 | Sulbactam + Cefoperazone Inj.2g Vial | Sulbactam sodium and Cefoperazone sodium injection(1:1) 2g vial. Each vial to contain Sulbactam sodium BP/USP equivalant to 1g of Sulbactam and Cefoperazone sodium BP/USP equivalant to 1g of Cefoperazone as a sterile dry powder for reconstitution with suitable diluent for intramuscular or intravenous use. Note: 01.The product should be stable for 02 years when stored at a temperature range of 28'C-32'C. 02.Each vial should be labeled accordingly. | VIAL | 12,000 | | | | |
| 10 | Sodium Stibogluconte Inj.10g/100 mL Vial | Sodium stibogluconte Injection BP,100ml Vial Each 100ml vial to contain sterile solution of Sodium Stibogluconate in water for injection BP/USP equivalent to pentavalent Antimony 100mg/ml for intramuscular and intravenous injection. Note: 1.The product should be stable for 02 years when at a temperature range of 28'C-32'C. The product should be protected from light. 2.Each vial should be labelled accordingly | VIAL | 600 | | | | |
| 11 | Fluconazole Inj. 200mg in100mL Vial | Fluconazole Infusion BP 200mg in 100ml vial OR Fluconazole Injection USP 200mg in 100ml Vial Each 100ml vial to contain 200mg of Fluconazole for intravenous infusion Note. 1.This injection should be stable for minimum of 24 months when stored at a temperature range of 28'C-32'C 2.Each vial should be labelled accordingly. | VIAL | 30,000 | | | | |
| 12 | Fluconazole Cap. 200mg | Fluconazole Capsule BP 200mg Each capsule to contain 200mg of Fluconazole. Note: 01.The shelf life of the product should be minimum of 24 months. | CAP | 90,000 | | | | |

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|---|---|---|---|---|---|---|---|---|
| 13 | Amphotericin Inj. 50mg Vial | Amphotericin B for Injection USP,50mg Each vial to contain 50mg of serile complex of Amhotericin B USP, and Deoxycholate Sodium as dry powder for reconstitution for intravenous infusion Note: 1.This injection should be stable for a minimum of 02 years when stored within a temperature range 2'C- 8'C 2.Each vial should be labelled accordingly | VIAL | 3,750 | | | | |
| 14 | Nystatin Tab. 500,000IU | Nystatin Tablet BP/USP 500,000 IU Each coated tablet to contain 500,000IU of Nystatin BP/USP for oral use. Note: 1. Should be packed in air-tight containers. 2. The tablet should be stable for a minimum of 24 months when stored under the prescribed temperature. | TAB | 6,00,000 | | | | |
| 15 | Colistimethate for Injection 1,000,000IU Vial | Colistimethate Sodium Injection 1,000,000IU Powder for solution for injection, infusion or inhalation. Colistimethate Sodium for injection BP/IP 1,000,000IU Vial Each vial to contain 1,000,000IU of Colistimethate Sodium BP as dry powder for reconstitution. OR Colistimethate for Injection USP 1,000,000IU Vial Each vial to contain 1,000,000IU of Colistimethate Sodium USP as dry powder for reconstitution. Note: 1. This injection should be stable for 24 months when stored at a temperature below 25'C. 2. The product should be protected from light. 3. Each vial should be labelled accordingly. | VIAL | 1,09,000 | | | | |
| 16 | Ketoconazole Tab 200mg | Ketoconazole Tablet 200mg Each tablet to contains 200mg of ketoconazole BP/USP Note: 01.The shelf life of the product should be minimum of 24 months | TAB | 30,000 | | | | |
| 17 | Adenosine Inj. 6mg/2mL Vial | Adenosine Injection 6mg in 2ml Vial Each 2ml vial to contain 6mg of Adenosine USP for Intravenous Injection Note: 1.This injection should be stable for 02 years when stored at a temperature range of 28'C-32'C 2.Each vial should be labeled accordingly. | AMP | 7,500 | | | | |
| 18 | Amiodarone Tab. 100mg | Amiodarone Tablet 100mg Each tablet to contain 100mg of Amiodarone Hydrochloride BP Note: 01.The shelf life of the product should be minimum of 24 months | TAB | 10,50,000 | | | | |

| | | | | | | | | |
|---|---|---|---|---|---|---|---|---|
| 19 | Amiodarone Inj. 150mg/3mL Amp. | Amiodarone Intravenous infusion BP,150mg/3ml Each 3ml ampoule to contain sterile, concentrated solution of, 50mg /1ml of Amiodarone Hydrochloride in water for injection BP/USP, suitable for intravenous infusion after dilution. Note; 1.The preparation should be stable for minimum of 24 months when stored under storage condition specified by the manufacturer. 2.Each ampoule should be lablled accordingly | AMP | 35,000 | | | | |
| 20 | Metoprolol Inj. 5mg/5mL Amp. | Metoprolol Tartrate injection 1mg/ml,5ml ampoule Metoprolol Tartrate injection USP 1mg/ml,5ml ampoule Each 5ml ampoule to contain a sterile solution of Metoprolol Tartrate USP 5mg and Sodium chloride USP 45mg as a tonicity adjusting agent in Water for injection BP/USP OR Metoprolol injection BP,1mg/ml,5ml ampoule Each 5ml ampoule to contain a sterile solution of Metoprolol Tartrate BP 5mg in Water for Injection BP/USP. Note: 1. The injection should comply with the requirements stated under Parenteral Preparations and it should be preserve in single-dose, light - resistant ampoules. 2.The injection should be stable for 24 months when stored at a temperature range of 28'C-32'C | AMP | 1,000 | | | | |
| 21 | Sodium Nitroprussiide Inj. 50mgVial | Sodium Nitroprusside for Injection BP/USP 50mg Each vial to contain Sodium Nitroprusside BP/USP equivalent to 50mg of dehydrated Sodium Nitroprusside as powder for reconstitution with suitable solvent for intravenous injection. Note: 1. This injection should be stable for a minimum of 24 months when stored at a temperature range of 28'C-32'C 2. Each vial should be labelled accordingly | VIAL | 250 | | | | |
| 22 | Glyceryl Trinitrate(Nitroglycerin) Inj. 50mg/10mL Vial or Amp. | Nitroglycerin Injection USP 50mg/10ml Each 10ml vial/ampoule to contain 50mg of Nitroglycerin for preparing Intravenous infusion. Note: 1.This injection should be stable for a minimum of 24 months when stored at a temperature range of 28'C-32'C 2.Each vial/ampoule should be labelled accordingly. | VIAL | 27,000 | | | | |

| | | | | | | | | |
|---|---|---|---|---|---|---|---|---|
| 23 | Verapamil Injection 5mg in 2mL Ampoule | Verapamil Injection BP 5mg in 2ml OR Verapamil Hydrochloride Injection USP 5mg in 2ml. Each 2ml ampoule to contain 5mg of Verapamil Hydrochloride BP/USP in Water for injection BP/USP for slow intravenous injection. Note: 1. This injection should be stable for a minimum of 02 years when stored at a temperature range of 28'C-32'C 2. Each ampoule should be labelled accordingly. | AMP | 10,500 | | | | |
| 24 | Isoprenaline Inj. 2mg/2mL Amp. | Isoprenaline Injection BP 2mg/2ml Each 2ml amber coloured ampoules to contain 2 mg of Isoprenaline Hydrochloride BP for intravenous infusion. OR Isoproterenol Hydrochloride injection USP 2mg/2ml Each 2ml amber coloured ampoule to contain 2mg of Isoproterenol Hydrochloride USP for intravenous infusion. Note: 1.The injection should be stable for a minimun of 24 months when stored at a temperature range of 28'C-32'C 2.Each ampoule should be labelled accordingly. | AMP | 5,400 | | | | |
| 25 | Noradrenaline Inj. 4mg/2mL Amp | Noradrenaline Acid Tartrate Injection BP,4mg in 2ml ampoule Each 2ml ampoule to contain Noradrenaline Acid Tartrate BP, 4mg equivalent to Noradrenaline base 1mg/ml, for intravenous infusion OR Norephinephrine Bitartrate USP,4mg in 2ml ampoule Each 2ml ampoule to contain Norephinephrine Bitartrate USP,4mg equivalent to Norephinephrine base 1mg/ml,for intravenous infusion. Note: 1.The product should be protected from light. 2.The product should be stable for minmum of 24 months when stored at a temperature range of 28'C-32'C 3.Each ampoule to be labeled accordingly. | AMP | 7,50,000 | | | | |
| 26 | Phenylephrine Inj. 10mg/1mL Vial | Phenylephrine injection BP 10mg/1ml OR Phenylephrine Hydrochloride Injection USP 10mg/1ml Each 1ml vial to contain 10mg of Phenylephrine Hydrochloride BP/USP 10mg for subcutaeneous injection, intramuscular injection slow intravenous injection or intravenous infusion. Note: 01.The shelf life of the product should be minimum of 24 months | VIAL | 2,100 | | | | |

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| 27 | Protamine Sulphate Inj.50mg/5 mL | Protamine Sulphate injection BP 50mg/5ml. Each 5ml ampoule to contain 50mg of Protamine Sulphate BP in water for injection BP/USP for intravenous injection OR Protamine Sulfate injection USP 50mg/5ml Each 5ml ampoule to contain 50mg of Protamine Sulphate USP in water for injection BP/USP for intravenous injection OR Protamine Sulfate for injection USP 50mg/5ml. Each 5ml ampoule to contain 50mg of Protamine Sulfate USP as sterile dry powder with suitable dry diluents for reconstitution with suitable solvent before use for intravenous injection. Note: 1.This injection should be stable for a minimum of 24 months when stored within the temperature range of 2C - 8C. 2.Sufficient overage to be added to the adequate shelf life. 3.Each ampoule should be labelled accordingly. | VIAL | 6,000 | | | | |
| 28 | Tranexamic Acid Tab/Cap 500mg | Tranexamic Acid Tablet BP/USP/IP 500mg OR Tranexamic Acid Capsule 500mg Each tablet/capsule to contain 500mg of Tranexamic Acid BP/USP/IP. Note: 01.Tablets should be scored 02. The shelf life of the product should be minimum of 24 months. | TAB | 12,00,000 | | | | |

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| 29 | Factor VIII 200-350IU Vial with vW Factor | Dried factor VIII Fraction BP (Dried Human Antihaemophilic Fraction) OR Human Coagulation Factor VIII Ph Eur. OR Antihemophilic Factor USP OR Dried Human Antihaemophilic Fraction IP Each vial to contain 200 - 350 IU of concentrated, intermediate purity and detergent treated dried factor VIII Fraction BP, Ph Eur,USP or IP. Note: 1. The product should contain von Willebrand factor and factor content should be stated on the label. General Note: 1. The item should be stable at temperature 2'C - 8'C. 2. The product should have minimum 24 months shelf life at the time of delivery to MSD. 3. Tenderer should submit detailed specifications of the product offered. 4. The product should ensure, at least two steps on virus inactivation as recommended by WHO/US.FDA 5. The donor selection process should be specified by the manufacturer. 6. Each batch should be certified as free from HIV and hepatitis viruses. 7. Anti viral test methods used for screening for HIV and Hepatitis viruses should be declared by the manufacturer. The test methods used should be approved by WHO/US.FDA. 8. Each vial to be supplied with suitable diluent. 9. The product should be protected from light. | VIAL | 4,500 | | | | |

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| 30 | Dried Recombinant Factor VIII Fraction 200IU-350IU Vial | Dried Recombinant Factor VIII Fraction 200IU-350IU vial<br>Dried factor VIII Fraction BP (Dried Recombinant Antihaemophilic Fraction) 200IU-350IU vial<br>OR<br>Recombinant Coagulation Factor VIII Ph Eur. 200IU-350IU vial<br>OR<br>Recombinant Antihemophilic Factor USP200IU-350IU vial<br>OR<br>Dried Recombinant Antihaemophilic Fraction IP200IU-350IU vial<br>Each vial to contain 200IU-350IU of concentrated, purified and viruses inactivated 3rd or generation recombinant dried factor VIII Fraction BP, Ph Eu USP or IP.<br>Note:<br>1. The item should be stable at temperature 2C – 8C.<br>2. The product should have minimum 24 months shelf life at the time of delivery to MSD.<br>3. Tenderer should submit detailed specifications of the product offered.<br>4. The product should ensure, at least two steps on virus inactivation as recommended by WHO/US.FDA<br>5. The purification process should be specified by the manufacturer.<br>6. Each batch should be certified as free from HIV hepatitis and other viruses.<br>7. Anti viral test methods used for screening for HIV and Hepatitis viruses should be declared by the manufacturer. The test methods used should be approved by WHO/US.FDA.<br>8. Each vial to be supplied with suitable diluent<br>9. The product should be protected from light. | Vial | 9,996 | | | | |

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| 31 | Dried Recombinant Factor VIII Fraction 500IU Vial | Dried Recombinant Factor VIII Fraction 500IU vial<br>Dried factor VIII Fraction BP (Dried Recombinant Antihaemophilic Fraction) 500IU vial<br>OR<br>Recombinant Coagulation Factor VIII Ph Eur. 500IU via<br><OR<br>Recombinant Antihemophilic Factor USP 500IU vial OR<br>Dried Recombinant Antihaemophilic Fraction IP 500IU vial<br>Each vial to contain 500IU of concentrated, purified and viruses inactivated 3rd or 4th generation recombinant dried factor VIII Fraction BP, Ph Eur, USP or IP<br>Note:<br>1. The item should be stable at temperature 2C – 8C<br>2. The product should have minimum 24 months shelf life at the time of delivery to MSD<br>3. Tenderer should submit detailed specifications of the product offered<br>4. The product should ensure, at least two steps on virus inactivation as recommended by WHO/US.FDA<br>5. The purification process should be specified by the manufacturer.<br>6. Each batch should be certified as free from HIV hepatitis and other viruses.<br>7. Anti viral test methods used for screening for HIV and Hepatitis viruses should be declared by the manufacturer. The test methods used should be approved by WHO/US.FDA.<br>8. Each vial to be supplied with suitable diluent.<br>9. The product should be protected from light. | Vial | 15,000 | | | | |
| 32 | Papaverine HCl Inj.60mg/2 mL Amp. | Papaverine hydrochloride injection 60mg in 2ml ampoule Each ampoule to contain sterile solution of Papaverine Hydrochloride 60mg/2ml, in water for injection BP/USP for slow intramuscular or intravenous injection. Note: 1. The injection should be stable for a minimum of 2 years when stored at at a temperature range of 28'C-32'C 2. Each ampoule should be labelled accordingly. | AMP | 3,200 | | | | |

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| 33 | Triclofos Oral Solution 500 mg/5 mL in 30 mL Bottle | Triclofos Oral Solution BP/IP 500 mg/5 mL in 30 mL Bottle Each 5 mL of solution contains 500 mg of Triclofos Sodium BP/IP. | Bot | 8,000 | | | | |
| 34 | Paracetamol Infu. 10mg/mL,100mL | Paracetamol 10mg/mL Solution for Infusion OR Paracetamol Infusion IP 10mg/mL in 100mL Glass bottle/Vial or collapsible bag Each 100mL glass bottle/vial or collapsible bag with hanger to contain 10mg/1mL of Paracetamol BP or Acetaminophen USP, solution for infusion. Note: 01. Each bottle or collapsible bag should be labelled accordingly. 02. The shelf life of the product should be minimum of 24 months | VIAL | 76,000 | | | | |
| 35 | Ethosuximide Cap 250mg | Ethosuximide Capsule BP/USP 250mg. Each capsule to contain 250mg of Ethosuximide BP/USP Note: 01.The shelf life of the product should be minimum of 24 months. | CAP | 85,000 | | | | |
| 36 | Flunarizine HClTab. 5mg | Flunarizine hydrochloride Tablet 5mg Each tablet to contain Flunarizine Hydrochloride equivalent to 5mg of Flunarizine Note: 01.The shelf life of the product should be minimum of 24 months. | TAB | 75,00,000 | | | | |
| 37 | Disulfiram Tab. 250mg | Disulfiram Tablet 250mg Each tablet to contain Disufiram BP/USP 250mg Note; 01.Tablets should be protected from light. 02.The shelf life of the product should be minimum of 24 months. | TAB | 1,00,000 | | | | |
| 38 | Levetiracetam Inj. 500mg/5mLVial/Amp. | Levetiracetam Injection 500mg in 5mL Vial/Ampoule Each 5mL ampoule/vial to contain 500mg of Levetiracetam USP in water for injection BP/USP Note: 01. The injection should be stable for a minimum of 24 months when stored under stipulated conditions. 02. Each ampoule/vial to be labelled accordingly. | VIAL | 31,500 | | | | |

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| 39 | Fat emulsion Inj. 20%,100m LBot. | Fat emulsion injection 20% in 100ml bottle Each 100ml bottle to contain in 20% w/v of soya oil for parenteral use BP, 1.2 - 1.5% w/v of Lecithin USNF 2.2 - 2.5% w/v of glycerol BP in a suitable vehicle. Note: Alternative formulate may be acceptable,therefore tenderers should give detailed composition of product offered. 1. The bottle should be graduated to 100ml and be made of medicinal grade Polyethylene or Polypropylene. PVC is not acceptable. 2. The bottle should have a device for hanging. 3. To prevent leakage of fluid with insertion or withdrawal of spike or continuous use the site of the insertion of piercing spike should be covered with a sterile rubber disk or bung and the plastic cover of the disk or bung should be fused to neck of the bottle right round. 4. Each bottle should be packed individually in a protective polyethylene cover. 5. The product should be sterile and stable for a minimum of 24 months within a temperature range of 30'C - 35'C. 6. The bottle should be labelled accordingly. | BOT | 5,100 | | | | |

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| 40 | Amino acid solution for intravenous infusion, 100mL Bottle | Protein Hydrolysate injection USP in 100ml Bottle. Each 100ml solution to contain Amino Acids and short chain peptides which represents the approximate nutritive equivalent to the casein, Lactalbumin, plasma fibrin or other suitable protein which derived by the acid, enzymatic or other methods of hydrolysis. It may contain alcohol, Dextrose or other carbohydrate suitable for intravenous infusion not less than 50% of the total nitrogen present is in the form of L-amino Nitrogen for intravenous infusion for infants and children. Note: 1. The bottle should be graduated to 100ml and be made of medicinal grade Polyethylene or Polypropylene. PVC is not acceptable. 2. The bottle should have a device for hanging. 3. To prevent leakage of fluid with insertion or withdrawal of spike or continuous use the site of the insertion of piercing spike should be covered with a sterile rubber disk or bung and the plastic cover of the disk or bung should be fused to neck of the bottle right round. 4. Each bottle should be packed individually in a protective polyethylene cover. 5. The product should be sterile and stable for a minimum of 24 months within a temperature range of 30 C- 35 C. 6. The bottle should be labelled accordingly. | BOT | 4,800 | | | | |
| 41 | Thiamine Tab.100mg | Thiamine tablet BP/USP 100mg Each tablet to contain 100mg of Thiamine hydrochloride BP/USP. Note: 1. Shelf life of the product should be minimum of 24 months. | TAB | 6,90,000 | | | | |
| 42 | Epoetin Inj. 4000IU PFS | Epoetin Injection 4,000IU Prefilled syringe Each pre-filled syringe to contain 4,000IU of Epoetin (Recombinant Human Erythropoietin) in Water for injection BP/USP for subcutaneous or intravenous injection. Note: 1. The product should be stable for minimum of 02 years when stored under 2'C-8'C. 2. Each Pre-filled syringe should be labeled accordingly. | PFSY | 8,00,000 | | | | |

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| 43 | Desferrioxamine Inj. 500mg | Desferrioxamine Mesylate for Injection BP, 500 mg. Each vial to contain 500mg of Desferrioxamine Mesylate BP as dry powder for reconstitution with water for injection BP/USP before use for subcutaneous, intramuscular and intravenous injection OR Desferrioxamine Mesylate for injection USP 500mg. Each vial to contain 500mg of Desferrioxamine Mesylate USP as dry powder for reconstitution with water for injection BP/USP before use for subcutaneous, intramuscular and intravenous injection Note: 1.The product should be stable for minimum of 02 years when store below 25'C 2.Each vial should be labelled accordingly. | VIAL | 2,70,000 | | | | |
| 44 | Human Albumin Sol. 20%, 50mLBot. | Human Albumin Solution BP/USP/Ph Eur 20% in 50ml bottle Each 50ml vial to contain at least 95% of the Albumin Protein BP / Ph Eur, derived from human plasma serum. Note: 1.The product should fulfill the requirement of virus elimination to include both enveloped and non-enveloped viruses (including HIV,HBV,HCV,HAV, Parvo virus B19)using virus reduction inactivation methods with pasteurization of the final product as per the current WHO approved techniques(WHO,TRS,No941,2007 & WHO,TRS,No 924,2004) 2.The product should be stable for minimum of 02 years when stored under the storage condition stipulated by the manufacturer. 3.The product should be protected from light. 4.Each vial should be labeled accordingly. 5.The manufacturer should submit a certificate for each batch of the product that it has been tested free of Hepatitis B, Hepatitis C, and HIV infections | BOT | 2,00,000 | | | | |
| 45 | Phosphate Tab. 500 mg | Phosphate tablet 500 mg Each tablet to contain 500mg of Phosphate Note: The shelf life of the product should be minimum of 24 months | TAB | 1,56,000 | | | | |

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| 46 | Total Parenteral Nutrition in500mL-1,500mL Collap. Bag | Total Parenteral Nutrition in multiple component in collapsible bag (central or peripheral)<br>Per pack should contain;<br>-Glucose/Dextrose calories 300kCal-1000kCal<br>- Nitrogen 4g-6g<br>- Calories from Fat 300kCal-800kCal<br>- Lipid source - SMOF 20% (Soyabean 30%, MCT 30%, Olive Oil 25%, Fish oil 15%) or 50% MCT+40% Soya bean +10% omega 3 or<br>Medium & long chain triglyceride (20% w/v)<br>- pH 5 to 6<br>Note: 1. Each collapsible bag should be labeled accordingly | BAG | 15,000 | | | | |
| 47 | Salbutamol Respiratory Sol.0.5%,15mL Vial | Salbutamol Respiratory Solution 5mg/mL Vial OR Salbutamol Nelbulizer Solution BP 5mg/mL Vial Each 10 mL to 30 mL vial is to contain Salbutamol Sulphate BP equivalent to 0.5% w/v of Salbutamol in a suitable base for use with a nebulizer Note: Each vial should be labelled accordingly. | VIAL | 2,75,000 | | | | |
| 48 | Pirfenidone Tablets 200mg | Pirfenidone Tablets 200mg Each coated tablet to contains 200mg of Pirfenidone BP. Note: 01.The shelf life of the product should be minimum of 24 months. | TAB | 5,50,000 | | | | |
| 49 | Rabies Vaccine (Human use)(0.5ml/1ml). Each dose 0.1ml to contain at least 7 IU/ml of rabies Antigen as an inactivated freeze-dried powder form.Each vial should be either 0.5mL or 1mL. The Vaccine should be suitable for human use, produced in cell culture.(Eg. Primary chickembryo cell culture Rabies Vaccine, Vero cell Rabies Vaccine). The vaccine should comply with the Rabies Vaccine as the Pharmacopoeial requirement of Vaccine and biological products. Note: 1.Cold chain monitors or WHO recommended other cold chain monitoring device should be included for each cartoon and the cold chain should be maintained according to the manufacturer's instructions during storage, transport & delivery of vaccine. 2.The following documents should be submitted pre- shipment for | DOSE | 15,00,000 | | | | |

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| | | each lot of vaccine dispatched. (a).Invoice (b).Certificate of origin (c).Certificate of analysis.(d).Lot release certificate from National control Laboratory(NCL) from country of origin.(e).Summary lot protocols of production procedure & quality control testing. (f).Packaging list. (g). Copy of product information leaflet(PIL). These documents should be submitted pre-shipment to the National Control Laboratory for Vaccines (NCL-MRI) Sri Lanka. 3.Airway bill & temperature monitoring data should be submitted with the sample submission for lot release to NCL. 4.The product should be only from fresh stocks and should have a minimum of 2/3 of remaining shelf life at the time of delivery at MSD. 5.The vaccine should be stored at a temperature range of 2'C - 8'C & should not be frozen. 6.Each vial should be labelled accordingly indicating both date of manufacture and expiry. 7.The vaccine should be recommended by WHO for intradermal use / the vaccine should be recommended and registered by the National Regulatory Authority of the manufacturing country or a reference country for intradermal use 8.Each vial should be provided with a suitable sterile diluent (5 vials of vaccine + 5 vial of diluent to be packed in the same box). 9. Immunization record cards should be provided for each vial. Patient information should be provided in Sinhala language for 75% of the consignment and Tamil language for the remain 25%(Template to be obtain from MRI). 10. Director/MSD should be informed two weeks prior to the arrival of vaccine. | | | | | | |

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| 50 | Pneumococcal Vaccine SingleDose Vial (PPSV-23) | Pneumococcal Polysaccharide Vaccine BP (PPSV-23) Each 0.5mL single dose vaccine vial/prefilled syringe to contain a mixture of 23 equal parts of highly purified immunochemically different capsular Polysaccharide antigens, prepared from suitable pathogenic strains of Streptococcus pneumoniae for subcutaneous or intramuscular use. Note: 1. Vaccines recommended for bulk purchase by the WHO for UN agencies will be considered as an added advantage (WHO pre- qualified). 2. The following documents should be submitted pre-shipment for each lot of vaccine dispatched. (a).Invoice (b).Certificate of origin (c).Certificate of analysis.(d).Lot release certificate from National control Laboratory (NCL) from country of origin.(e).Summary lot protocols of production procedure & quality control testing. These documents should be submitted pre-shipment to the National Control Laboratory for Vaccines (MRI) in Sri Lanka. 3. The product should be only from fresh stocks and should be stable for a minimum 2/3 of remaining shelf life at the time of delivery at MSD when stored under storage condition specified by the manufacturer. 4. The vaccine should also comply with the general requirements for vaccines in the BP or USP or other pharmacopeial standards approved by NMRA, Sri Lanka. 5. Cold Chain monitors or WHO recommended other cold chain monitoring device should be included for each carton during storage transport and delivery of vaccine. 6. The vaccine should be protected from light and should be stored at a temperature range of 2'C-8'C. Vaccine should not be frozen. 7. Each vaccine vial/prefilled syringe should be labelled accordingly indicating both date of manufacture and expiry. 8. The vaccine should meet the most recent requirements of WHO when tested by the methods outlined by WHO. 9. Director/MSD should be informed two weeks prior to the arrival of vaccine. | VIAL | 4,500 | | | | |

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| 51 | Varicella Vaccine 0.5mL Vial | Varicella Vaccine Live BP single dose vials Each single dose vial of freeze - dried vaccine (live ) to contain attenuated OKA strain of Varicella virus BP Packing: 0.5ml (single dose vial) Note i. Vaccines recommended for bulk purchase by the WHO for UN agencies will beconsidered as an added advantage (WHO pre qualified). ii. The following documents should be submitted pre-shipment for each lot of vaccine despatched. (a).Invoice (b).Certificate of origin (c).Certificate of analysis.(d).Lot release certificate from National control Laboratory(NCL) fromcountry of origin.(e).Summary lot protocols of production procedure & quality control testing. These documents should be submitted pre-shipment to the National Control Laboratory for Vaccines (NCL-MRI) Sri Lanka. iii.. Each consignment should have a minimum remaining shelf life of 75% at the time of receiving the consignment at MSD. iv.The vaccine should also comply with the general requirements for vaccine in the BP, USP or other pharmacopeial standards approved by NMRA, Sri Lanka v. The vaccine should meet the most recent requirements of W.H.O. when tested by the methods outlined by WHO. vi. Cold chain Monitors should be included in each carton and the Cold Chain should be maintained according to the manufacturer's instructions during storage,transport and delivery of vaccine. vii.The vaccine should be stored at temperature between +2'C and + 8'C. viii.Each vial should be provided with a suitable sterile diluent. ix. Each vial should be labled accordingly indicating batch No., date of manufacture and expiry. x. Director/MSD should be informed two weeks prior to the arrival of vaccine. | VIAL | 450 | | | | |

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| 52 | Anti Rabies Serum Inj. 1,000I.U./5 mL Amp. | Rabies Antiserum BP 1000 IU/5mL OR<br>Anti Rabies Serum USP 1000 IU/5mL<br>Each 5mL ampoule to contain 1000 IU of Rabies Antiserum BP or Anti Rabies Serum USP Note 1:<br>1. This injection should be stable for 24 months when stored within a temperature range of 2'C-8'C. Do not freeze.<br>2.Each consignment should have a minimum remaining shelflife of 75% at the time of receiving the consignment at MSD.<br>3. Cold chain monitors should be included for each carton and cold chain should be maintained according to the manufacturer's instructions during storage, transport and delivery of item.<br>4. Each vial should be labelled accordingly.<br>Note<br>The following documents should be submitted pre-shipment for each lot of serum dispatched.<br>1. Summary protocols for production and quality testing (Including protocols for immunization, initial pool, final bulk and finished product ) based on WHO model summary protocols.<br>2. Lot release certificate from National Control Laboratory of manufacturing country for that particular batch.<br>3. Certificate of analysis of finished product.<br>4. Certificate of analysis of diluents, if applicable.<br>5. The package information leaflet. | VIAL | 24,000 | | | | |

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| 53 | Anti Rabies HumanImmunoglobulin 300IU Vial | Human Rabies Immunoglobulin BP OR Human Rabies Immune Globulin USP Each vial to contain 300IU of Rabies Immunoglobulin/ Rabies Immune Globulin obtained from Human Plasma containing specific anti bodies against Rabies virus suitable for intramuscular use. Note: 1.This injection should be stable for 20 months when stored within a temperature range of 2'C - 8'C. Do not freeze. 2.Each consignment should have a minimum remaining shelf life of 75% at the time of receiving the consignment at MSD. 3. The manufacturer should submit a certificate for each batch of the product that it has been tested free of Hepatitis B, Hepatitis C and HIV infections. 4. Each vial should be labelled accordingly indicating batch No., date of manufacture and expiry. 5. Cold Chain monitors or WHO recommended other cold chain monitoring device should be included in each pack during storage, transport and delivery of vaccine. | VIAL | 1,200 | | | | |
| 54 | Biphasic Isophane Insulin(Human) Inj. 30/70 Vial | Biphasic Isophane Insulin injection BP,1000 IU / 10ml. Each 10ml rubber capped multidose vial to contain 100 IU/ml (Human 30% as soluble 70% as Isophane ) suspention of Human insulin complexed with Protamin sulphate in a solution of Human Insulin. Note: 1.This injection should be stable for a minimum of 02 years when stored within the temperature range of 2'C - 8'C . 2.Manufacturer should confirm the product is free of HIV and Hepatitis virus. 3.This injection to be presented in airtight tamper proof vial. 4.Each vial should be labelled accordingly. | VIAL | 22,00,000 | | | | |

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| 55 | InsulinIsophane (human) 1,000IU/10 mLVial | Isophane Insulin injection BP (NPH Insulin) derived from human 1000IU in 10ml. OR Isophane Insulin suspension USP (NPH Insulin) derived from huaman 1000 IU/10ml. Each rubber capped multidose Vial to contain 1000 IU/10ml of sterile purified suspension of Isophane Insulin BP/USP derived from human. Note: 1. This injection should be stable for a minimum of 24 months when stored within 2'C - 8'C. 2. Do not allow to freeze. 3. Each vial should be labelled accordingly. | VIAL | 36,000 | | | | |
| 56 | Insulin Soluble (Human) Inj.1,000IU/ 10mL Vial | Insulin Injection BP (Soluble Insulin) 1000 I.U. in 10ml derived from human. Each rubber capped multidose vial to contain sterile neutral solution of 1000 I.U./10ml of human insulin BP. OR Insulin Injection USP (Soluble Insulin)1000IU in 10ml derived from human. Each rubber capped multidose vial to contain sterile neutral solution of 1000 IU/10ml of human Insulin USP. Note: 1. This injection should be stable for a minimum of 24 months when stored within 2'C -8'C. 2. Do not allow to freeze. 3. Each vial should be labelled accordingly. | VIAL | 72,000 | | | | |
| 57 | Glucagon HCl Inj. 1 IU Vial | Glucagon Hydrochloride Injection BP/USP 1IU vial. Each vial to contain 1IU (1mg) of Glucagon Hydrochloride BP/USP as dry powder for reconstitution with water for injection BP/USP for subcutaneous, intramuscular or intravenous injection. Note; 1.The product should be stable for minimum of 02 years when stored under 2'C - 8'C. 2.Each vial should be labeled accordingly. | VIAL | 1,200 | | | | |
| 58 | Dexamethasone Tab. 4 mg | Dexamethazone Tablets BP 4mg Each tablet to contain 4mg of Dexamethazone BP OR Dexamethazone Tablets USP 4mg Each tablet to contain 4mg of Dexamethazone USP Note: 01.Tablets should be scored 02.The shelf life of the product should be minimum of 24 months | TAB | 2,70,000 | | | | |

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| 59 | Hydroxyprogesterone Inj.250mg/1mL Amp. | Hydroxyprogesterone injection BP, 250mg/ml Each 1ml ampoule to contain 250mg of Hydroxyprogesterone Hexanoate BP in a suitable ester in a suitable fixed oil, or in a mixture of these for intramuscular injection. OR Hydroxyprogesterone Caproate injection USP, 250mg/ml Each 1ml ampoule to contain 250mg of Hydroxyprogesterone Caproate in a suitable ester in a suitable fixed oil, or in a mixture of these for intramuscular injection. Note: 1.This injection should be stable for a minimum of 24 months when stored at a temperature range of 28'C-32'C 2.Each ampoule should be labelled accordingly. | AMP | 9,000 | | | | |
| 60 | Tetracosactrin Inj.250mcg/1mL Amp. | Tetracosactrin (Tetracosactide) Injection 250microgram/1ml Each 1ml ampoule to contain 250microgram of Tetracosactide BP in water for injection BP/USP, for intramuscular or intravenous injection. Note: 1. This injection should be stable for a minimum of 24 months when stored within a temperature range or 2'C - 8'C. Protect from light. 2. Each ampoule should be labelled accordingly | AMP | 1,300 | | | | |
| 61 | Somatropin for Inj. 2IU-30IU | Human Growth Hormone or Somatropin for Injection BP/USP/IP<br>Each container to contain of Somatropin for injection BP/USP/IP for subcutaneous or intramuscular injection.<br>OR<br>Somatropin Powder and solvent for solution for inj.<br>Note:<br>1.This Injection should be stable for a minimum of 24 months when stored at a temperature which is mentioned by the manufacturer<br>2. The shelf life of the reconstituted solution should be minimum of 28 days when stored at a temperature which is mentioned by the manufacturer<br>3.Each container should be labelled accordingly.<br>4.The product should be approved by the stringent regulatory authorities or WHO Pre-qualified | IU | 2,24,000 | | | | |

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| 62 | Desmopressin Tab. 100mcg | Desmopressin Tablet 100microgram Each tablet to contain 100microgram of Desmopressin Acetate BP. Note: 01.Tablets should be protected from light. 02.The shelf life of the product should be minimum of 24 months. | TAB | 60,000 | | | | |
| 63 | Desmopressin acetate NasalSpray 10mcg/metered spray( 50 MS) | Desmopressin acetate Nasal spray 10 microgram/metered spray ( 50 metered spray) Each vial to contain 50 metered sprays (50 doses) and each metered spray to contain 10 microgram of Desmopressin Acetate. Note: 01.The shelf life of the product should be minimum of 24 months. | VIAL | 4,200 | | | | |
| 64 | Triamcinolone Acetonide Inj (preser. free) 40mg/mLVial | Triamcinolone acetonide (preservative free) injection 40mg in 1ml vial. Each vial to contain 40mg of Triamcinolone acetonide BP as sterile suspension in very fine particles, in water for injections containing suitable dispersing agents for intra- articular or intramuscular use. OR Each vial to contain 40mg of Triamcinolone acetonide USP,as sterile suspension in a suitable aqueous medium for intra-articular or intramuscular use. Note: 1. Product should be protected from light. 2. Each vial should be labeled accordingly. | VIAL | 1,500 | | | | |
| 65 | Diazoxide tab. 50mg | Diazoxide tablets 50mg Each tablet to contain 50mg of Diazoxide BP/USP Note: 01.The shelf life of the product should be minimum of 24 months. 02.The product should be protected from light. | TAB | 12,000 | | | | |
| 66 | Dutasteride Cap. 0.5 mg | Dutasteride capsule 0.5 mg Each tablet to contain 0.5mg of Dutasteride Note: 01.The shelf life of the product should be minimum of 24 months | CAP | 30,000 | | | | |
| 67 | Estradiol Valerate Tab. 1mg | Estradiole valerate tablet 1mg Each coated tablet to contain 1mg of Estradiol valerate USP Note: 01.The shelf life of the product should be minimum of 24 months. 02.The product should store in the original package in order to protect from light and moisture. 03.Each calendar pack should contains 28 tablets | TAB | 18,000 | | | | |

| | | | | | | | | |
|---|---|---|---|---|---|---|---|---|
| 68 | Metoclopramide Tab. 10mg | Metoclopramide tablets BP 10mg Each tablet to contain Metoclopramide Hydrochloride BP equivalent to 10mg of anhydrous Metoclopramide Hydrochloride BP OR Metoclopramide tablets USP 10mg Each tablet to contain Metoclopramide Hydrochloride USP equivalent to 10mg of anhydrous Metoclopramide Hydrochloride | TAB | 11,00,000 | | | | |
| 69 | Metoclopramide Inj.10mg/2 mLAmp. | Metoclopramide Injection BP 10mg/2ml Each 2ml amber coloured ampoule to contain Metoclopramide Hydrochloride BP/USP equivalent to 10mg of anhydrous Metoclopramide Hydrochloride Note: 1.This injection should be stable for a minimum of 24 months when stored at a temperature range of 28'C-32'C. 2.Each ampoule should be labelled accordingly. | AMP | 11,00,000 | | | | |
| 70 | Ranitidine Inj. 50mg/2mL Amp. | Ranitidine Injection BP/USP, 50mg/2ml Each 2ml ampoule to contain Ranitidine Hydrochloride BP/USP equivalent to 50mg of Ranitidine in Water for Injection BP/USP Note: 01.Each ampoule should be labelled accordingly. 02.The shelf life of the product should be minimum of 24 months | AMP | 6,50,000 | | | | |
| 71 | Mesalazine tab. 400mg | Mesalazine tablet 400mg Each tablet to contain 400mg of Mezalazine BP OR Mesalamine Tablet USP 400mg Each tablet to contain 400mg of Mesalamine USP Note: 01.Tablet should be stored in an airtight container and protected from light. 02.The shelf life of the product should be minimum of 24 months | TAB | 1,50,000 | | | | |
| 72 | Macrogol 3350/4000 (10g-13.125g of PEG) Oral powder sachets (with or without electrolytes) | Macrogol 3350/4000 (10-13.125g of PEG) oral powder sachets with or without electrolytes Each sachet to contain 10-13.125g of Macrogol 3350/4000. Note: The shelf life of the product should be minimum of 24 months. Each sachet should be labelled accordingly. | Sach | 9,498 | | | | |

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|---|---|---|---|---|---|---|---|---|
| 73 | Terlipressin Acetate Inj.1mg Vial/Amp. | Terlipressin Acetate Injection 1mg vial Each vial to contain 1mg of Terlipressin BP as sterile, dry powder for reconstitution and solvent for solution for injection. Note: 1. This product should be stable for a minimum of 24 months when stored at a temperature below 25'C and protected from light. 2. Each vial should be labeled accordingly. OR Terlipressin Acetate Solution for Injection 1mg Ampoule Each ampoule to contain 1mg of Terlipressin BP as sterile and solution for injection. Note: 1. This product should be stable for a minimum of 24 months when stored at a temperature of 2 驴 C-8 驴 C and protected from light. 2. Each ampoule should be labeled accordingly. | VIAL | 21,000 | | | | |
| 74 | Fluorescein Sodium Inj.10%, 2mL-5mL Vial | Fluorescein Injection BP/USP,10%w/v Each 2mL-5mL vial to contain Fluorescein sodium BP/USP 10%, in water for injection BP/USP for intravenous use as diagnostic fluorescein angiography or angioscopy of fundus and iris. Note: 1.This injection should stable for a minimum of 24 months when stored at a temperature range of 28'C-32'C. 2.Each vial should be labelled accordingly | VIAL | 2,900 | | | | |
| 75 | Olopatadine HCl Eyedrops 1mg/mL, 5mL Dropper Bot. | Olopatadine hydrochloride eye drops 1mg/mL, 5mL dropper bottle Each 5mL dropper bottle to contain 1mg per 1mL Olopatadine hydrochloride for ophthalmic use Note: 01.The shelf life of the product should be minimum of 24 months 02. Each dropper bottle should be labelled accordingly. 03. Product should be individually packed in light resistant boxes/cartons with a leaflet . | BOT | 18,000 | | | | |
| 76 | Chlorambucil Tab. 2mg | Chlorambucil Tablet BP 2mg Each coated tablet to contain 2mg of Chlorambucil BP Note: 1.The tablet should be stable for a minimum of 24 months when stored within temperature range of 2'C-8'C | TAB | 15,000 | | | | |

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|---|---|---|---|---|---|---|---|---|
| 77 | Cyclophosphamide Tab. 50mg | Cyclophosphamide Tablet BP/USP 50mg Each coated tablet to contain Cyclophosphamide BP equivalent of 50mg of anhydrous Cyclophosphamide. Note: 1.The tablet should be stable for a minimum of 24 months when stored at a temperature range of 28'C - 32'C. | TAB | 1,05,000 | | | | |
| 78 | Epirubicin HCl Inj.50mg/25 mL Vial | Epirubicin Hydrochloride for injection 50mg vials. Each vial to contain 50mg of Epirubicin Hydrochloride (CAS-56390-09-01) as sterile dry powder for reconstitution with water for injection BP/USP before use for intravenous infusion OR Epirubicin Hydrochloride injection 50mg/25ml Each 25ml vial to contain 50mg of Epirubicin Hydrochloride (CAS-56390-09-1) in a suitable vehicle for intravenous infusion. Note: 1. This injection should be stable for a minimum of 24 months when stored within a temperature range of 2'C - 8'C 2. Each vial should be packed in light resistant box. 3. Each vial should be labelled accordingly | VIAL | 4,500 | | | | |
| 79 | Mitomycin for Inj. 2mg Vial | Mitomycin for injection USP/JP 2mg vial Each vial to contain a mixture of 2mg of Mytomycin USP/JP and Mannitol USP as sterile dry powder for reconstitution before intravenous use Note: 1.This injection should be stable for a minimum of 24 months when stored at a temperature range of 28'C - 32'C 2.Each vial should be labelled accordingly | VIAL | 6,000 | | | | |
| 80 | Cytarabine Inj. 100mg/5m L Vial(Not for Intrathecal use) | Cytarabine Injection BP 100mg/5ml,Not for Intrathecal use Each 5ml vial to contain 100mg of Cytarabine BP in water for injection BP/USP for intravenous and subcutaneous injection OR Cytarabine for Injection BP/USP 100mg,Not for Intrathecal use Each vial to contain 100mg of Cytarabine BP/USP as sterile dry powder for reconstitution with water for injection BP/USP before use for intravenous and subcutaneous injection Note: 1.This injection should be stable for 24 months when stored at a temperature range of 28'C - 32'C 2.Each vial should be labelled accordingly | VIAL | 1,020 | | | | |

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|---|---|---|---|---|---|---|---|---|
| 81 | Cytarabine Inj. 100mg/mL Vial ,Preservative free | Cytarabine for injection BP 100mg in 1ml,Preservative free,for intrathecal use Note: 1.This injection should be stable for 24 months when stored at a temperature within 28'C-32'C. 2.Each vial should be labelled accordingly | VIAL | 1,100 | | | | |
| 82 | Fluorouraci Inj. 250mg/10 mLVial | Fluorouracil injection BP/USP 250mg/10ml Each 10ml amber coloured vial to contain Fluorouracil Sodium equivalent to 250mg of Fluorouracil BP/USP for intravenous injection, intravenous infusion and intra-arterial infusion. Note: 1.This injection should be stable for a minimum of 24 months when stored within temperature of 15'C - 25'C 2.Each vial should be labelled accordingly | VIAL | 600 | | | | |
| 83 | Gemcitabine HClInj. 200mg Vial | Gemcitabine Hydrochloride Injection 200mg Vial Each vial to contain Gemcitabine Hydrochloride as sterile dry powder for reconstitution with water for injection for intravenous injection or intravenous infusion. 1. The product should be stable for minimum of 24 months when stored under storage condition specified by the manufacturer. 2. Each vial to be labelled accordingly. | VIAL | 3,000 | | | | |
| 84 | Methotrexate Inj. 1g/10mL Vial | Methotrexate injection BP/USP 1g/10ml Each 10ml vial to contain Methotrexate Sodium BP/USP equivalent to 100mg/ml Methotrexate in water for injection BP/USP for intramuscular, intravenous use. OR Methotrexate for injection USP 1g Each vial to contain Methotrexate Sodium USP/BP equivalent to 1g of Methotrexate as sterile freeze dried powder for reconstitution with water for injection BP/USP for intramuscular, intravenous use. Note: 1. This injection should be stable for minimum of 24 months when stored at a temperature range of 28'C-32'C. 2. Each vial should be labelled accordingly. | VIAL | 13,200 | | | | |
| 85 | Etoposide Cap.100mg | Etoposide Capsule USP/BP 100mg Each capsule to contain 100mg of Etoposide USP/BP. Note: 01.The shelf life of the product should be minimum of 24 months | CAP | 2,400 | | | | |

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|---|---|---|---|---|---|---|---|---|
| 86 | Etoposide Cap. 50mg | Etoposide Capsule USP 50mg Each capsule to contain 50mg of Etoposide USP. Note: 01. The shelf life of the product should be minimum of 24 months | CAP | 900 | | | | |
| 87 | Temozolomide Cap. 100mg | Temozolomide capsule 100mg Each capsule to contain Temozolomide 100mg Note: 01.The shelf life of the product should be minimum of 24 months | CAP | 9,600 | | | | |
| 88 | Carboplatin inj.150mg/15mLVial with Diluent | Carboplatin Injection 150mg in 15ml Vial with suitable diluent Each 15ml vial to contain 150mg of Carboplatin BP with suitable diluent for intravenous use Note: 01.The shelf life of the product should be minimum of 24 months | VIAL | 2,250 | | | | |
| 89 | Carboplatin Inj. 450mg/45 mLVial | Carboplatin Injection BP,450mg Each 45ml vial to contain 450mg of Carboplatin BP in water for injection BP/USP for intravenous use. Note: 1.This injection should be stable for a minimum of 24 months when stored under manufacturer's conditions. 2.Each vial should be labeled accordingly | VIAL | 12,600 | | | | |
| 90 | Procarbazine Cap. 50mg | Procarbazine Hydrochloride Capsule USP 50mg Each capsule to contain Procarbazine Hydrochloride USP equivalent to 50mg of Procarbazine Note: 01.The shelf life of the product should be minimum of 24 months | CAP | 2,400 | | | | |
| 91 | Imatinib Mesilate Tab/Cap100mg | Imatinib Mesilate Tablet/ Capsule 100mg Each Tablet/ capsule to contain 100mg of Imatinib Mesilate. Note: 01.The shelf life of the product should be minimum of 24 months | CAP | 3,60,000 | | | | |
| 92 | Filgrastim Inj 300mcg in0.5mL-1mL, PFS/Vial | Filgrastim Injection 300microgram in 0.5ml/1ml. Each Pre-filled syringe / vial to contain 300microgram of Filgrastim for subcutaneous injection or intravenous infusion. Note: 1.Shelf life of the product should be minimum of 24 months 2.Each pre-filled syringe/vial should be labelled accordingly. | VIAL | 1,35,000 | | | | |

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|---|---|---|---|---|---|---|---|---|
| 93 | Tamoxifen Tab. 20mg | Tamoxifen tablets BP 20mg Each tablet to contain Tamoxifen Citrate BP equivalent to 20mg of Tamoxifen. OR Tamoxifen Citrate tablets USP 20mg Each tablet to contain Tamoxifen Citrate USP equivalent to 20mg of Tamoxifen Packing: Pack suitably to ensure to protect from light until the time of administration of the drug Pack size: 100 tablets in a pack. Note: 01.The shelf life of the product should be minimum of 24 months | TAB | 9,00,000 | | | | |
| 94 | Goserelin Acetate Implant3.6 mg | Goserelin Acetate 3.6mg implant in syringe applicator. Each implant to contain Goseraline Acetate Ph Eur 3.6mg for subcutaneous use. Note; 1.The product should be stable for a minimum of 24 months when stored below 25'C . 2.The product should be protected from light. 3.Each implant should be labelled accordingly. | IMPL | 11,400 | | | | |
| 95 | Abiraterone Acetate Tab.250mg | Abiraterone acetate tablet 250mg Each tablet to contain 250mg of Abiraterone acetate Note: 01.The shelf life of the product should be minimum of 24 months | TAB | 3,00,000 | | | | |
| 96 | Mesna Inj. 200mgin 2mL Vial/Ampoule | Mesna injection 200mg in 2ml Each vial/ampoule to contain aqueous solution of mesna (sodium 2-mercapto-ethanesulphonate) 200 mg in 2 ml intravenous injection Note: 1.The injection should be stable for a minimum of 24 months when stored below 30'C 2.This injection should be protected from light. 3. Each vial/ampoule should be labelled accordingly. | VIAL | 18,000 | | | | |
| 97 | Clotrimazole Pessaries 100mg | Clotrimazole Pessaries BP 100mg Each vaginal tablet to contain 100mg of Clotrimazole BP. OR Clotrimazole vaginal insert 100mg USP Each vaginal insert to contain 100mg of Clotrimazole USP. Note: 1. The shelf life of the product should be minimum of 24 months. 2. Pack size: One pack should contain 3 or 6 pessaries. | PESS | 1,02,000 | | | | |
| 98 | Tolterodine SR Cap. 2mg | Tolterodine Sustained Release Capsule BP 2mg Each sustained release capsule to contain 2mg of Tolterodine BP/USP. Note: 01.The shelf life of the product should be minimum of 24 months | CAP | 60,000 | | | | |

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|---|---|---|---|---|---|---|---|---|
| 99 | Solifenacin Tab. 5mg | Solifenacin tablet 5mg Each tablet to contain 5mg of Solifenacin. Note: 01.The shelf life of the product should be minimum of 24 months. | TAB | 24,000 | | | | |
| 100 | Oxybutynine Tab. 2.5mg | Oxybutynine Tablets BP 2.5mg Each tablet to contain Oxybutynine Hydrocloride BP 2.5mg OR Oxybutynine Chloride Tablet USP 2.5mg Each tablet to contain Oxybutynine Hydrochloride USP 2.5mg Note: 01.The shelf life of the product should be minimum of 24 months. | TAB | 1,80,000 | | | | |
| 101 | Etomidate Inj. 20mg/10mLVial/Amp. | Etomidate Injection USP 20mg in 10mL Vial/Ampoule Each 10mL vial/ampoule to contain 20mg of Etomidate BP/USP/Ph Eur for slow intravenous injection. Note: 1. The injection should be stable for a minimum of 24 months when stored at a temperature range of 28'C-32'C. 2. Each vial should be labelled accordingly. | VIAL | 1,800 | | | | |
| 102 | GlycopyrrolateInj. 200mcg/mL Vial | Glycopyrrolate Injection USP 200 microgram in 1ml vial Each 1ml vial to contain 200microgram of glycopyrrolate USP in water for injection BP/USP for intramuscular or intravenous use. Note: 1. The injection should be stable for a minimum of 24 months when stored at at a temperature range of 28'C-32'C 2. Each vial should be labelled accordingly. | AMP | 24,000 | | | | |
| 103 | Flumazenil Inj. 500mcg/5mLVial/Amp. | Flumazenil Injection 500microgram in 5ml Vial or ampoule Each 5ml vial/ampoule to contain 500microgram of Flumazenil BP/Ph Eur for intravenous injection or intravenous infusion. Note: 1. The injection should be stable for a minimum of 24 months when stored at a temperature range of 28'C-32'C 2. Each ampoule should be labelled accordingly. | VIAL | 660 | | | | |

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|---|---|---|---|---|---|---|---|---|
| 104 | Naloxone Inj. 400mcg/mL Amp. | Naloxone Injection BP,400mcg/1ml OR Naloxone Hydrochloride Injection USP, 400mcg/1ml Each 1ml amber coloured ampoule to contain Naloxone Hydrochloride BP/USP equivalent to 400mcg of anhydrous Naloxone Hydrochloride for subcutaneous, intramuscular and intravenous use. 1.The Injection should be stable for minimum of 24 months when stored at a temperature range of 28'C-32'C 2.Each ampoule should be labelled accordingly. | AMP | 6,000 | | | | |
| 105 | Dantroline Sodium Inj. 20mg Vial | Dantroline Sodium Injection USP 20mg vial Each vial to contain 20mg of sterile reconstitution powder of Dantroline Sodium BP/USP for rapid intravenous injection. Note: 1. The injection should be stable for a minimum of 24 months when stored at a temperature range of 28'C-32'C 2. Each vial should be labelled accordingly. | VIAL | 350 | | | | |
| 106 | Lidocaine Topical aerosol10 %, 50mL Bot. | Lidocaine Topical aerosol USP 10% w/v Bottle Each 50mL bottle to contain Sterile solution of Lidocaine in suitable vehicle with suitable propellants in a pressurized container equipped with a metering valve and 10 plastic nozzles with 10cm-12cm arm to deliver 5mg to 10mg of Lidocaine USP. Note: 1.Minimum of 500 doses should be filled in non-reactive aerosol container equipped with metered dose valve. | BOT | 2,700 | | | | |
| 107 | Lidocaine HCl Gel 2%,30g Tube | Lidocaine Gel BP, 2% w/v (Sterile) OR Lidocaine Hydrochloride Gelly USP 2% w/v (sterile) Each air tight 30g tube to contain Lidocaine hydrochloride BP/USP equivalent to 2% w/v of anhydrous Lidocaine hydrochloride in a suitable water soluble sterile viscous base. Note: 1.The product should be stable for a minimum of 24 months when stored at a temperature range of 28'C-32'C 2.Each tube should be labelled accordingly. | TUBE | 2,40,000 | | | | |

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|---|---|---|---|---|---|---|---|---|
| 108 | Lidocaine 2% + Adrenalin1: 80,000 Inj. 30mL Vial | Lidocaine and Adrenaline Injection BP/Lidocaine and Epinephrine Injection BP, 2%, 30ml Vial Each 30ml amber coloured glass ampoule to contain 2% w/v of Lidocaine Hydrochloride BP and Adrenaline Acid Tartrate BP equivalent to 1 in 80,000 of Adrenaline/Epinephrine base OR Lidocaine Hydrochloride and Epinephrine injection USP 2% w/v, 30ml Each 2ml amber coloured glass ampoule to contain 2% w/v of Lidocaine Hydrochloride and Epinephrine USP equivalent to 1 in 80,000 of Epinephrine base. Note: 1.This injection should be stable for a minimum of 02 years when stored at a temperature range of 28'C-32'C, protect from light. 2.Each ampoule should be labelled accordingly. 3.The label should state both Lidocaine with Adrenaline Injection and Lidocaine and Epinephrine injection. | VIAL | 2,10,000 | | | | |
| 109 | Dexmedetomidine HCl Inj200mcg/ 2mL Vial | Dexmedetomidine Hydrochloride Injection 200microgram/2ml vial Each vial to contain Dexmedetomidine Hydrochloride 200microgram. Note: 1.The injection should be stable for minimum of two years when stored under Tropical climate conditions. 2.Each vial should be labelled accordingly. | VIAL | 2,400 | | | | |
| 110 | Morphine Tab. 30mg | Morphine Tablet BP 30mg. Each Tablet to contain 30mg of Morphine sulphate BP. Note: 01.The shelf life of the product should be minimum of 24 months. | TAB | 1,50,000 | | | | |
| 111 | Morphine Sulphate Oral solu.2mg/1 mL, 100mL Bott. | Morphine Sulfate Oral solution (BP) 2mg/1ml 100ml bottle, Each 100ml bottle to contain 2mg/1ml of Morphine Sulfate BP/USP in a suitable flavored and sweetened syrup/suspension base. Note: 01.Product should be free from alcohol 02.The product should be stable for 24 months when stored at 28'C-32'C 03.The product should be protected from light | BOT | 750 | | | | |

- ***In BOQ, Bidders are requested to quote for per piece (per unit) rate only***
- ***GST amount for per piece to be mentioned (percentage of GST cannot be entered).***

**ANNEXURE – 5**

| Packing Material Specification | |
|---|---|
| BABY CARTON | 350Gsm foreign art card with 4 Colour printing, single side printing,Tuck in flap system,finishing with outer gloss lamination with dye punching and pasting |
| DISPLAY CARTON | 350Gsm ITC saffaire graphic 4 Colour printing, single side printing,Tuck in flap system with locked bottom,finishing with outer gloss lamination with dye punching and pasting |
| MASTER CARTON (CORRUGATED BOX) | Narrow Flute 7 Ply Corrugated Card Board Box Total Gsm = >1147 inner & outer ply virgin kraft paper of which outer ply to be alkali resistant with bitumen. The box shall be single piece with double stapling using flat wire of MS or GI material as per ISI 10066, 1981. Gsm: - (outer Line bituminised) 160, Inner lining 120x3 flute= 150x3 (@35% extra for 3 ply corrugating). Direction of flute: Vertical, nature of flute: Narrow. Punch Resistance - Not less than 45deg. C 0Zs per tear inch. Bursting strength: 18 kg/cm2 (min.) (bursting factor not less than 20, Gum -Nature: Starch Based.). |
| ALUMINIUM FOIL | Thickness- 0.021mm to 0.022 mm (21 to 22 micron), Gsm - 59 (54 to 56 aluminium + HSL 3 min.) |
| PVC/PVDC | Food Grade Thermo formable transparent blister foil. Thickness= 0.35 mm max. Gsm= 320 to 330, Sealing= Proper sealing, PVC= Non Toxic - PVC food grade, Yield= 3.125 to 3.03 mt2/ kg |
| LEAFLET | Maplitho Deluxe Paper Of 70 Gsm Min. Two Folds Printed In Single Colour (Black) |

**Note: Bidders may adopt appropriate packing mode, however shall ensure that the packing must be suitable for cargo handling/export by air. There may also be a branding requirement in the tertiary packing.**

**Product literature must be printed in English**

**SPECIMEN LABEL FOR OUTER CARTON**

**Product Name: (like Paracetamol IP - 500mg)**
**Batch No. :**
**Mfg. Date:**
**Exp. Date:**
**Total Quantity:**
**Net Weight of the Carton:**
**Manufactured By:**

**Annexure-06**

# MANUFACTURER'S AUTHORIZATION FORM

**No. _____ Dated _____**

**To**
_____
_____
_____
_____

**Dear Sir,**

**Bid Ref. No. _____**

We _____ who are established and reputable manufacturers of _____ having factories at _____ Registered office at_____ possessing Manufacturing Licence No. _____, dated_____, valid upto _____ (copy enclosed) do hereby authorize M/s_____ (Name and Address of Representative) to submit a bid, and subsequently negotiate and sign the contract with you against the above mentioned tender.

No company or Firm or individual other than M/s _____ are authorized to bid, negotiate and conclude the contract in regard to this business against this specific tender.

We hereby extend our full guarantee and warranty as per the tender conditions for the goods offered for supply against this invitation for bid by the above firm.

Your faithfully,

(Name )

For and on behalf of M/s _____

(Name of Manufacturers)

**Note :** This letter of authority should be on the letterhead of the manufacturing concern and should be signed by a person competent and having the power of attorney to bind the manufacturer.

**For and behalf of the firm**
**(Firm Name & Address)**

**Annexure 07**

## Category details of organization

| SL No. | Description | Yes/No |
|---|---|---|
| 1. | *Whether the organization belongs to the MSE category | |
| 2. | *If yes whether the organization belongs to MSE category | |
| 3. | *Whether the MSE organization belongs to SC/ST entrepreneur. | |
| 4. | *Whether the MSE organization belongs to woman entrepreneur. | |
| 5 | Whether the MSE organization is registered under MSE Type of Enterprise **'Trading'** | |

***Kindly furnish the copies of documents supporting your above claim along with this Annexure duly filled.**

***The Udyog Aadhar no of the bidder ……..**

**(Self-attested copy of Udyog Aadhar registration certificate should be submitted along with the technical bid)**

**Date:**

**Place:**

**Signature of the Bidder:**

**Name with seal:**
**Designation:**
**Address:**

**Annexure 08**

To,

Vice President (PS) and GH (HCS)
HLL Lifecare Limited,
HLL Bhavan, Poojappura,
Thiruvananthapuram -695012 Kerala, India
Tel: 0471 2775500, 0471 2350959 (EXTN - 606 /531)
Website – www.lifecarehll.com

## INDEMNITY CERTIFICATE

Dear Sir,

As a supplier to HLL, the indemnifier assumes liability for and irrevocably agrees to indemnify, defend and hold harmless Government of India and HLL Lifecare Limited, its Affiliates, shareholders, officers, directors, employees, agents, and their respective successors and assigns from and against any and all losses, damages, claims, actions, liabilities, proceedings, injury, cost or expenses (including counsel's fees of whatsoever kind of nature arising out of or in any way connected with the licenses granted or the manufacture of the products or out of any defect (whether obvious or hidden) in the products or arising from the indemnifier's failure to comply with applicable laws.

Dated this [insert: number] day of [insert: month], [insert: year].

Signature.....

Name.....

Full Address with contact person Name, Phone number and Email

Designation and Common Seal...

**Annexure 09**

## Performance Bank Guarantee Format*

To: _____ (Name of Purchaser) **WHEREAS** _____ (Name of Supplier) (hereinafter called "the Supplier") has undertaken, in pursuance of Contract No._____ dated _____20___ to supply _____ (Description of Goods and Services) (hereinafter called "the Contract").

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

**AND WHEREAS** we have agreed to give the Supplier a Guarantee:

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

This guarantee is valid until the _____ day of_____ 20___.

Signature and Seal of Guarantors

_____

_____

Date: _____ 20 _____

Address: _____

***"The successful Bidder on whom the purchase order / Letter of Intent is placed shall be required to furnish a Performance Guarantee mandatorily only in the form of e-Bank Guarantee (e-BGs) from the Banks which are integrated with NeSL any other mode of Performance Bank Guarantees from such Banks will not be accepted. The details of Banks which are integrated with NeSL is listed at Annexure 16 and Indicative Challan for obtaining e-BG from Bank is at Annexure 17"**

**Annexure 10**

## CHECK LIST

| Sl No | PARTICULAR OF DOCUMENT | ATTACHED / NOT ATTACHED | PAGE NO | Remarks |
|---|---|---|---|---|
| 1 | Forwarding letter indicating the submission of Technical documents along with check list of document | | | |
| 2. | EMD/ Tender Fee in the form of BG/DD (copy of the NEFT/RTGS details) | | | |
| 3 | Tender document duly signed and stamped in all pages along with corrigendum if Any) | | | |
| 4 | Duly attested copies of manufacturing license along with product list highlighting the quoted product | | | |
| 5 | All Pharmaceutical products should have relevant & valid WHO GMP certificate | | | |
| 6 | Valid Drug license for quoted Products in case for authorized agents | | | |
| 7 | Copy of Udyog Aadhaar, in case of MSE bidders | | | |
| 8 | Authenticated copy of the Memorandum of Association/Articles of Association / Partnership deed etc and certificates of incorporation/ registration of the organization with details of Name, Address, Tel. No., Fax No., E-mail Address of firm and the M. Director / Partner / Proprietor | | | |
| 9 | Documentary proof attested by Charted Accountant for establishing the average annual turnover of Original Manufacturers having a minimum average annual turnover of Rs.1 Crore (Rupees One Crore only) during the last three years i.e. 2021-22 , 2022-2023 and 2023-2024 (Original/ provisional). In case of Authorized agents they must submit the documentary proof attested by Charted Accountant for minimum average turnover in the last three years i.e., 2021-22 , 2022-2023 and 2023-2024 (Original/ provisional) is Rs.50 lakh (Rupees Fifty lakhs only) And documentary proof attested by Charted Accountant for establishing their Principal manufacturers meets the eligibility criteria for original manufacturer as specified above. In case of bid by authorized agents, manufacturers authorization form must be attached with the bid submitted | | | |
| 10 | Copy of Recent Non conviction certificate issued by State drug controller | | | |
| 11 | Power of Attorney in stamp paper (RS.200/-) duly notarized authorizing the signatory to sign the bids and transact business. | | | |
| 12 | Authorization letter from manufacturer (Self–attested Copy). | | | |
| 13 | Annexure 1 - Self Declaration | | | |
| 14 | Annexure 2 - Bid Form | | | |
| 15 | Annexure 3 - Under taking letter for replacement of complaint/defective goods | | | |
| 16 | Annexure 4 – List of quoted products with specification Compliance | | | |
| 17 | Annexure 5 – Packing Specifications | | | |
| 18 | Annexure 6 - Manufacture Authorization Form (if applicable) | | | |
| 19 | Annexure 7 - Category details of Organization | | | |
| 20 | Annexure 8 - Indemnity Certificate | | | |
| 21 | Annexure 10 - Check List | | | |
| 22 | Annexure 11 – Compliance To Rule 144 (XI) of GFR 2017 (Self Declaration) | | | |
| 23 | Annexure 12 - Make In India Preference (Self Declaration) | | | |
| 24 | Annexure 13 – Pre Contract Integrity Pact | | | |
| 25 | Annexure 14- Fall Clause Declaration | | | |
| 26 | Annexure 15- Bidder Info | | | |
| 27 | Annexure 16- List of Banks integrated with NeSL | | | |
| 28 | Annexure 17- Indicative Challan | | | |
| 29 | Copy of PAN Card & GSTN details | | | |

**Annexure 11**

## SELF DECLARATION – COMPLIANCE TO RULE 144 (XI) OF GFR 2017

We,

……

……

…….

**(Include name and address of the bidder)**

Hereby declare that we are eligible to bid for the tender: ……

**(Include tender number and date)**

As per the eligibility stipulated by Government Order no F.No.6/18/2019-PPD dated 23-July-2020 inclusive of the latest amendments regarding insertion of rule 144(Xi) in the General Financial Rules (GFR) 2017, issued by Ministry of Finance, Government of India.

We are aware that any bidder indenting to participate in this tender who is from a country which shares a land border with India will be eligible to bid in this tender only if the bidder is registered with Competent Authority as per the GO.

Date:
Place:

Signature of the Bidder:

Name with seal:

Designation:

Address:

**Annexure 12**

## SELF DECLARATION – MAKE IN INDIA PREFERENCE

In line with Government Public Procurement Order No. P-45021/2/2017-BE-II dt. 15.06.2017, as amended from time to time and as applicable on the date of submission of tender, we hereby certify that we M/s_____ (supplier name) are local supplier meeting the requirement of minimum Local content (50%) as defined in above orders for the material against Tender No_____ Details of location at which local value addition will be made is as follows: ---------- ---------- ---------- We also understand, false declarations will be in breach of the Code of Integrity under Rule 175(1)(i)(h) of the General Financial Rule for which for which a bidder or its successors can be debarred for up two years as per Rule 151 (iii) of the General Financial Rules along with such other actions as may be permissible under law.

Seal and Signature of Authorized Signatory

**Annexure 13**

## PRE-CONTRACT INTEGRITY PACT

This Pre-Contract Integrity Pact (herein after called the Integrity Pact) is made on ------t day of the month of ----------,

**Between**

HLL Life Care Limited, a Government of India Enterprise with registered office at HLL Bhavan, Poojappura, Thiruvananthapuram 695 012, Kerala, India. (Hereinafter called "HLL", which expression shall mean and include, unless the context otherwise requires, his successors in office and assigns) of the First Party.

**And**

---------- India represented by Shri ----------
(hereinafter called the "BIDDER / Seller" / Contractor which expression shall mean and include, unless the context otherwise requires, his successors and permitted assigns) of the Second Party.

**Preamble**

[Both HLL and BIDDER referred above are jointly referred to as the Parties]

HLL intends to award, under laid down organizational procedures, Purchase orders / contract/s against Tender /Work Order. HLL desires full compliance with all relevant laws and regulations, and the principles of economic use of resources, and of fairness and transparency in its relations with its Bidder/s and Contractor/s.

NOW, THEREFORE,

To avoid all forms of corruption by following a system that is fair, transparent and free from any influence /prejudiced dealings prior to, during and subsequent to the currency of the contract to be entered into with a view to:-

1. Enable HLL to obtain the desired materials/ stores/equipment/ work/ project done at a competitive price in conformity with the defined specifications by avoiding the high cost and the distortionary impact of corruption on public procurement; and
2. Enable the BIDDER to abstain from bribing or indulging in any corrupt practice in order to secure the contract by providing assurance to them that their competitors will also abstain from bribing and other corrupt practices and HLL will commit to prevent corruption, in any form, by its officials by following transparent procedures.

The parties hereto hereby agree to enter into this Integrity Pact and agree as follows:

**Clause.1. Commitments of HLL**

1.1 HLL undertakes that HLL and /or its Associates (i.e. employees, agents, consultants, advisors, etc.) will not demand, take a promise for or accept, directly or through intermediaries, any bribe, consideration, gift, reward, favour or any material or immaterial benefit or any other advantage from the BIDDER, either for themselves or for any person, organization or third party related to the contract in exchange for an advantage in the bidding process, bid evaluation, contracting or implementation process related to the contract.

1.2 HLL will, during the tender process / pre-contract stage, treat all BIDDERs with equity and reason, and will provide to all BIDDERs the same information and will not provide any such information or additional information, which is confidential in any manner, to any particular BIDDER which could afford an advantage to that particular BIDDER in comparison to other BIDDERs in relation to tendering process or during the contract execution.

1.3 All the officials of HLL will report to Chief Vigilance Officer of HLL (CVO), any attempted or completed breaches of the above commitments as well as any substantial suspicion of such a breach.

1.4 HLL will exclude from the process all known prejudiced persons and persons who would be known to have a connection or nexus with the prospective bidder.

1.5 If the BIDDER reports to HLL with full and verifiable facts any misconduct on the part of HLL's Associates (i.e. employees, agents, consultants, advisors, etc.) and the same is prima facie found to be correct by HLL, necessary disciplinary proceedings, or any other action as deemed fit, including criminal proceedings may be initiated by HLL. Further, such an Associate may be debarred from further dealings related to the contract process. In such a case, while an enquiry is being conducted by HLL the proceedings under the contract would not be stalled.

**Clause 2. Commitments of BIDDERs/ CONTRACTORs**

2. The BIDDER commits itself to take all measures necessary to prevent corrupt practices, unfair means and illegal activities during any stage of its bid or during any pre-contract or post-contract stage in order to secure the contract or in furtherance to secure it and in particular commit itself to the following:-

2.1 The BIDDER will not offer, directly or indirectly (i.e. employees, agents, consultants, advisors, etc.) any bribe, gift, consideration, reward, favour, any material or immaterial benefit or other advantage, commission, fees, brokerage or inducement to any official of HLL, connected directly or indirectly with the bidding process, or to any person, organization or third party related to the contract in exchange for any advantage in the bidding, evaluation, contracting and implementation of the contract.

2.2 The BIDDER further undertakes that it has not given, offered or promised to give, directly or indirectly any bribe, gift, consideration, reward, favour, any material or immaterial benefit or other advantage, commission, fees, brokerage or inducement to any official of HLL or otherwise in procuring the contract or forbearing to do or having done any act in relation to obtaining or execution of the contract or any other contract with the Government for showing or forbearing to show favour or disfavor to any person in relation to the contract or any other contract with the Government.

2.3 The BIDDER will not engage in collusion, price fixing, cartelization, etc. with other counterparty(s).

2.4 The counterparty will not pass to any third party any confidential information entrusted to it, unless duly authorized by HLL.

2.5 The counterparty will promote and observe ethical practices within its Organization and its affiliates.

2.6 BIDDER shall disclose the name and address of agents and representatives and Indian BIDDERS shall disclose their foreign principals or associates.

2.7 The counterparty will not make any false or misleading allegations against HLL or its Associates.

2.8 BIDDERs shall disclose the payments to be made by them to agents / brokers or any other intermediary, in connection with this bid/contract.

2.9 The BIDDER further confirms and declares to HLL that the BIDDER is the original integrator / manufacture /authorized government sponsored export entity of the defense stores and has not engaged any individual or firm or company whether Indian or foreign to intercede, facilitate or in any way to recommend to HLL or any of its functionaries, whether officially or unofficially to award the contract to the BIDDER, nor has any amount been paid, promised or intended to be paid to any such individual, firm or company in respect of any such intercession, facilitation or recommendation.

2.10 The BIDDER while presenting the bid or during pre-contract negotiations or before signing the contract, shall disclose any payments he has made, is committed to or intends to make to officials of HLL or their family members, agents, brokers or any other intermediaries in connection with the contract and the details of services agreed upon for such payments.

2.11 The BIDDER will not accept any advantage in exchange for any corrupt practice, unfair means and illegal activities.

2.12 The BIDDER commits to refrain from giving any complaint directly or through any other manner without supporting it with full and verifiable facts.

2.13 If the BIDDER or any employee of the BIDDER or any person acting on behalf of the BIDDER, either directly or indirectly, is a relative of any of the officers of HLL, or alternatively, if any relative of an officer of HLL has financial interest /stake in the BIDDER's firm, the same shall be disclosed by the BIDDER at the time of filing of tender.

The term 'relative' for this purpose would be as defined in Section 6 of the Companies Act 1956.

2.14 The BIDDER shall not lend to or borrow any money from or enter into any monetary dealings or transactions, directly or indirectly, with any employee of HLL.

2.15 The BIDDER will not collude with other parties interested in the contract to impair the transparency, fairness and progress of the bidding process, bid evaluation, contracting and implementation of the contract, and will not enter into any undisclosed agreement or understanding with other Bidders, 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.

2.16 The BIDDER will not commit any offence under the relevant Indian Penal Code, 1860 or Prevention of Corruption Act, 1988; further the Bidder(s)/ Contractor(s) will not use improperly, for purposes of competition or personal gain, or pass on to others, any information or document provided by the HLL as part of the business relationship, regarding plans, technical proposals and business details, including information contained or transmitted electronically. The BIDDER also undertakes to exercise due and adequate care lest any such information is divulged.

2.17 The BIDDER will not instigate third persons to commit offences outlined above or be an accessory to such offences.

2.18 The Bidder(s)/Contractors(s) of foreign origin shall disclose the name and address of the Agents /representatives in India, if any. Similarly the Bidder(s) /Contractors(s) of Indian Nationality shall furnish the name and address of the foreign Principal(s), if any.

**Clause.3. Previous contravention and Disqualification from tender process and exclusion from future contracts**

3.1 The BIDDER declares that no previous contravention occurred in the last three years immediately before signing of this Integrity Pact, with any other company in any country in respect of any corrupt practices envisaged hereunder or with any Public Sector Enterprise in India or any Government Department in India that could justify BIDDER's exclusion from the tender process

3.2 The BIDDER agrees that if it makes incorrect statement on this subject, BIDDER can be disqualified from the tender process or the contract, if already awarded, can be terminated for such reason.

If BIDDER before award or during execution has committed a contravention through a violation of Clause 2, above or in any other form such as to put his reliability or credibility in question, HLL is entitled to disqualify the BIDDER from the tender process.

**Clause .4. Equal treatment of all Bidders / Contractors / Subcontractors**

4.1 The Bidder(s) / Contractor(s) undertake(s) to demand from his Subcontractors a commitment in conformity with this Integrity Pact.

4.2 HLL will enter into agreements with identical conditions as this one with all Bidders and Contractors.

4.3 HLL will disqualify from the tender process all bidders who do not sign this Pact or violate its provisions.

**Clause .5. Consequences of Violation / Breach**

5.1 Any breach of the aforesaid provision by the BIDDER or any one employed by it or acting on its behalf (whether with or without the knowledge of the BIDDER) shall entitle HLL to take all or any one of the following action, wherever required:-

i. To immediately call off the pre-contract negotiations without assigning any reason or giving any compensation to the BIDDER. However, the proceedings with the other

ii. If BIDDER commits violation of Integrity Pact Policy during bidding process, he shall be liable to compensate HLL by way of liquidated damages amounting to a sum equivalent to 5% to the value of the offer or the amount equivalent to Earnest Money Deposit /Bid Security, whichever is higher.

iii. In case of violation of the Integrity Pact after award of the contract, HLL will be entitled to terminate the contract. HLL shall also be entitled to recover from the contractor liquidated damages equivalent to 10% of the contract value or the amount equivalent to security deposit/ performance guarantee, whichever is higher.

iv. To immediately cancel the contract, if already signed, without giving any compensation to the BIDDER.

v. To recover all sums already paid by HLL, and in case of an Indian BIDDER with interest thereon at 2% higher than the prevailing Prime Lending Rate of State Bank of India, while in case of a BIDDER from a country other than India with interest thereon at 2% higher than the LIBOR. If any outstanding payment is due to the BIDDER from HLL in connection with any other contract for any other stores, such outstanding payment could also be utilized to recover the aforesaid amount.

vi. To encash the advance bank guarantee and performance guarantee / warranty bond, if furnished by the BIDDER, in order to recover the payments already made by HLL, along with interest.

vii. To cancel all or any other contract with the BIDDER. The BIDDER shall be liable to pay compensation for any loss or damage to HLL resulting from such cancellation/recession and HLL shall be entitled to deduct the amount so payable from the money(s) due to the BIDDER.

viii. To debar the BIDDER from participating in future bidding processes of HLL for a minimum period of five (5) years, which may be further extended at the discretion of HLL or until Independent External Monitors is satisfied that the Counterparty will not commit any future violation.

ix. To recover all sums paid in violation of this Pact by BIDDER(s) to any middleman or agent or broker with a view to securing the contract.

x. In cases where irrevocable Letters of credit have been received in respect of any contract signed by HLL with the BIDDER, the same shall not be opened.

xi. Forfeiture of performance guarantee in case of a decision by HLL to forfeit the same without assigning any reason for imposing sanction for violation of the pact.

5.2 HLL will be entitled to all or any of the actions mentioned in para 5.1(i) to (x) of this pact also on the commission by the BIDDER or any one employed by it or acting on its behalf (whether with or without the knowledge of the BIDDER), of an offence as defined in Chapter IX of the Indian Penal Code, 1860 or Prevention of Corruption Act, 1988 or any other statute enacted for prevention of corruption.

5.3 The decision of HLL to the effect that a breach of the provisions of this Pact has been committed by the BIDDER shall be final and conclusive on the BIDDER. However, the BIDDER can approach the Independent External Monitor(s) appointed for the purposes of this Pact.

**Clause.6. Fall Clause**

The BIDDER undertakes that it has not supplied/is not supplying similar product/systems or subsystems OR providing similar services at a price / charge lower than that offered in the present bid in respect of any other Ministry/Department of the Government of India or PSU and if it is found any stage that similar product/systems or sub systems was supplied by the BIDDER to any to the Ministry/Department of the Government of India or a PSU at a lower price, then that very price, with due allowance for elapsed time will be applicable to the present case and the difference in the cost would be refunded by the BIDDER to HLL, if the contract has already been concluded.

**Clause .7. Independent External Monitor(s)**

7.1 HLL has appointed Independent External Monitor(s) (hereinafter referred to as Monitor(s)) for this Pact in consultation with the Central Vigilance Commission.

7.2 The responsibility of the Monitor(s) shall be to review independently and objectively, whether and to what extent the parties comply with the obligations under this Pact.

7.3 The Monitor(s) shall not be subject to instructions by the representatives of the parties and perform their functions neutrally and independently.

7.4 Both the parties accept that the Monitor(s) have the right to access all the documents relating to the project/ procurement, including minutes of meetings.

7.5 As soon as the Monitor(s) notices, or has reason to believe, a violation of this pact, he will so inform the CVO.

7.6 The BIDDER(S) accepts that the Monitor(s) have the right to access without restriction to all project documentation of HLL including that provided by the BIDDER. The BIDDER will also grant the Monitor(s), upon his request and demonstration of a valid interest, unrestricted and unconditional access to his project documentation. The same is applicable to subcontractors engaged by the BIDDER. The Monitor(s) shall be under contractual obligation to treat the information and documents of the BIDDER/ Subcontractor(s) with confidentiality.

7.7 HLL will provide to the Monitor(s) sufficient information about all meetings among the parties related to the Project provided such meeting could have an impact on the contractual relation between the parties. The parties will offer to the Monitor(s) option to participate in such meetings.

7.8 The Monitor(s) will submit a written report to the CVO of HLL within 8 to 10 weeks from the date of reference or intimation to him by HLL/BIDDER and, should consent arise, submit proposals for correcting problematic situations.

**Clause.8.Criminal charges against violating Bidder(s)/**

**Contractor(s)/ Subcontractor(s)**

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

**Clause.9. Facilitation of Investigation**

In case of any allegation of violation of any provisions of this Pact or payment of commission, HLL or its agencies shall be entitled to examine all the documents, including the Books of Accounts of the BIDDER and the BIDDER shall provide necessary information and documents in English and shall extend all possible help for the purpose of such examination.

**Clause.10. Law and Place of Jurisdiction**

Both the Parties agree that this Pact is subject to Indian Law. The place of performance and hence this Pact shall be subject to Thiruvananthapuram Jurisdiction.

**Clause.11. Other legal Actions**

The actions stipulated in the Integrity Pact are without prejudice to any other legal action that may follow in accordance with the provisions of the extant law in force relating to any civil or criminal proceedings.

**Clause.12. Validity and Duration of the Agreement**

This Pact begins when both parties have legally signed it. It expires for the Contractor/Successful bidder 12 months after the last payment under the contract or the complete execution of the contract to the satisfaction of the both HLL and the BIDDER /Seller, including warranty period, whichever is later, and for all other Bidders/unsuccessful bidders 6 months after the contract has been awarded.

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 as specified above, unless it is discharged / determined by Chairman and Managing Director of HLL.

**Clause. 13. Other provisions**

13.1 Changes and supplements as well as termination notices need to be made in writing. Both the Parties declare that no side agreements have been made to this Integrity Pact.

13.1 If the Contractor is a partnership or a consortium, this agreement must be signed by all partners or consortium members.

13.1 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

INWITNESS THEREOF the parties have signed and executed this pact at the place and date first above mentioned in the presents of following witnesses:

| **HLL** | **BIDDER** |
|---|---|
| | (Name & Designation) |
| Chairman and Managing Director<br>HLL Lifecare Limited,<br>Thiruvananthapuram. | |
| Witness | Witness |
| 1…….. | 1…… |
| 2…… | 2……..…….. |

* Provisions of these clauses would be amended /deleted in line with the policy of HLL in regard to involvement of Indian agents of foreign suppliers.

**Annexure 14**

## FALL CLAUSE DECLARATION

The BIDDER undertakes that it has not supplied/is not supplying similar product/systems or subsystems OR providing similar services at a price/ charge lower than that offered in the present bid in respect of any other Ministry/Department of the Government of India or PSU and if it is found any stage that similar product/systems or sub systems was supplied by the BIDDER to any to the Ministry/Department of the Government of India or a PSU at a lower price, then that very price, with due allowance for elapsed time will be applicable to the present case and the difference in the cost would be refunded by the BIDDER to HLL, if the contract has already been concluded.

Seal and Signature of Authorized Signatory

**Annexure-15**

## Bidder Info

The bidder should provide the following details in company letter head

- Bidder Name
- Address
- Contact Person
- Contact Number
- Email ID for communication
- GSTIN & PAN details

**Annexure-16**

**The details of Banks which are integrated with NeSL for obtaining e-BG is listed as below:**

**Sr. No. Banks / NBFCs**

1. Axis Bank Ltd
2. Bank of Baroda
3. Bank of India
4. Bank of Maharashtra
5. Canara Bank
6. Central Bank of India
7. HDFC Bank Ltd
8. ICICI Bank Ltd
9. IDBI Bank Ltd
10. IFIN Securities Finance Ltd
11. Indian Bank
12. Indian Overseas Bank
13. Karnataka Bank
14. Indus Ind Bank Ltd
15. Kotak Mahindra Bank Ltd
16. Punjab & Sind Bank
17. Punjab National Bank
18. SIDBI
19. State Bank of India
20. Sundaram Finance Limited
21. Tata Capital Financial Services
22. The Federal Bank Ltd
23. The Karur Vysya Bank Ltd
24. The South Indian Bank
25. UCO Bank
26. Union Bank of India

**Annexure- 17**

**For issuance of Electronic Bank Guarantees (e-BG) through National e-Governance Services Limited (NeSL) platform, the details of Challan form for obtaining e-BG from Bank is as under:-**

| Sr.No. | Field | Description of Field |
|---|---|---|
| 1 | PAN / UIN* | ANFPA6883M |
| 2 | Name | HLL Lifecare Limited |
| 3 | Email ID | sdrbdsouth@lifecarehll.com |
| 4 | Name of the Representative | Vice president (PS) & GH (HCS) |
| 5 | Mobile Number | 9400027975 |
| 6 | Relation to tender | Options are :<br>Beneficiary |
| 7 | Tender Reference Number | |
| 8 | BG Amount | xxxxxxxxxx |
| 9 | BG CCY (currency by default this could be Rs. | Indian Rupees |
| 10 | Expiry Date | xxxxxxxxxx |
| 11 | Claim Expiry Date | xxxxxxxxxx |