ANNEX – 1

BID NO. : DHS/L/WW/30/25
DATE OF ISSUE : 06TH AUGUST, 2024
CLOSING DATE & TIME : 19TH SEPTEMBER, 2024 AT 09.00 HOURS SRI LANKA TIME

Special Conditions for tendering :

1. Offers should be accompanied with the original of valid registration certificate/any **Subsequent renewal certificates where applicable** or **a copy certified by an Attorney at law**, **of aforesaid document** issued by the National Medicine Regulatory Authority in Sri Lanka.
2. Offered item should bear both our SR number and the Item number. However at the bid opening only the item numbers will be read out. Therefore price quoted should be shown against each item number.
3. **A break-up of** FOB and Freight charges should be quoted separately against each item in addition to quoted C&F price.
4. The volume of the total quantity of each item should be given in cubic meters (m^3)
5. Foreign offers should be on C&F {CPT/CFR (into FOB and freight]} Colombo basis. **Only** FOB offers are not acceptable. If offers are received on Import & Supply basis from local suppliers, those offers should be in LKR. All local suppliers/manufacturers should in LKR for the total delivery price to MSD stores.
6. Fax/E-mail offers directly sent to State Pharmaceuticals Corporation are not acceptable. Tenderers are requested to draw their attention to the clause "Submission of Bids" of the bid document in this regard**.**
7. If awarded supplier is unable to adhere the delivery schedule due to no fault of the SPC/Ministry would result in the supplier being surcharge 0.5% of total bid amount per day from the due delivery date.
8. The original payment receipt has to be annexed to the offer. Offers without same will be rejected.
9. We reserve the right to reject offers which do not comply above.
10. **The offer should be valid up to 18.03.2025**

CONDITIONS FOR SUPPLY FOR SPC ORDERS – APPLICABLE FOR SURGICAL & LAB ITEMS

(a) Part A-General Order Conditions (GOC) of Supply

1. The consignments supplied in respect **of** an order concerned, shall exactly match with the reference sample submitted and the product information (item descriptions, shelf life/warranty where applicable, manufacturer's name, country of manufacture, country of origin, etc.) provided in the bid document by the supplier, which has been accepted by the procurement committee, and included in the Indent / Purchase Order (PO), issued by SPC.

2. All items shall be supplied, sourcing from the manufacturer and country of manufacturer, stated in the Purchase Order (PO)/Indent of SPC and wherever applicable shall have a valid product registration or waiver of registration from NMRA.

3. Maintaining the validity of the product registration during the period of supply(delivery schedule), obtaining waiver of registration &/ import license / manufacture licensing at NMRA, is a pre-requisite for the supply of surgical, pharmaceutical & relevant laboratory items. Hence all suppliers shall produce relevant valid registration certificates/licenses, when requested by MSD/SPC.

When the validity of the product/manufacturing licenses and registrations of NMRA (eg; manufacturing license, product registration and GMP certificates), of local manufacturers / local suppliers, lapses during the year or during the period of supply (delivery schedule), it shall be extended / renewed by the supplier. A certified copies of afore mentioned valid certificates shall be submitted to MSD by the supplier when deliveries are made.

4. The number of batches per consignment shall be minimal. Batch quantity shall be an equal multiple of the quantity of the consignment and the proportionate size of the batch quantity shall be not less than 15% of the quantity in the consignment.

5. If MSD decides to accept a part or full consignment, with deviations from certain tender conditions (eg: with regard to labeling/packaging etc. due to an urgency, that shall be done subject to, either rectifying the defect within 05 working days by the supplier, or recovering the total cost [a] of rectifying the defect by MSD (via a duly contracted third party providing such services), from the supplier with a 25% surcharge on the labeling cost. (total charge = [a]+[a]x0.25) or 2% of the invoiced value, whichever is the highest.

All possible tender deviations such as Packing, labeling, delivery schedule, storage status, payment mode & conditions, etc., shall be communicated and agreed upon before accepting the tender award by the supplier. Noncompliance of same shall be considered as tender violations, to apply surcharge (as clause No. 37).

6. The specifications of the product offered in the bid by the supplier shall match with the tender specifications for the item and **any form of alternate offers will not be entertained.**

Shelf life & Warrantees

7. ~~In the supply of all Non consumables; Manufacturer or supplier or local agent shall provide a minimum of 02 year warranty period or as specified in the specification, for each such item or it's sub components supplied (through the local agent), unless otherwise agreed upon with MSD, prior to awarding the tender. Foreign suppliers of all such items shall have their own local agent in Sri Lanka, capable of providing technical support, repairs & spares, when necessary.~~

8. Freshly manufactured stocks of the product shall be supplied; thereby the residual Shelf Life (shelf life remaining at the time of delivery of goods in Sri Lanka/MSD stores in case of local supplies) of the product, shall be 85% of the shelf life requested (specified in order/Indent/PO).
In respect of the items with requested shelf life equal or more than 24 months, any deficit between the residual shelf life and requested shelf, shall not be more than 04 months.

In the violation of the above tender condition, SPC/MSD reserves the right to accept a reduced quantity, that is usable (as per the consumption rate) up to three months before the expiry of same and will subject to application of a penalty (as clause No. 37).
When the shelf life is not specified in the indent/PO/item spec; the requested shelf life shall be considered as, 24 months.

Standards & Quality

9. ~~Standards; In addition to Pharmacopoeial Standards that are indicated in the item specifications, other Pharmacopoeial Standards that are registered at National Medicines Regulatory Authority in Sri Lanka are also acceptable when no bidders have quoted for the standard specified in the item specification.~~

10. ~~Any product deficient of its sub components/ accessories, not at the specified quality standards or all its components not unitized appropriately in packaging (as a set), shall be rejected.~~

11. Withdrawal from use of items due to quality failure found as manufacturer's fault:

(a). In case of batch withdrawal, **value of entire batch quantity supplied** shall be recovered from the supplier.

(b).In case of product withdrawal, **value of entire product quantity** supplied shall be recovered from the supplier.

(c). In the event of either a) or b) above, supplier shall be surcharged the total **cost involved for MSD, of the quality failed supplies** with 25% administrative surcharge of the same**.**

12. The storage conditions and the packing requirements of the product shall conform to the information given by the manufacturer and accepted by NMRA for the product registration or shall conform to the information submitted for waiver of registration granted by NMRA in exceptional circumstances.(refer clause No.24)

If the offered product, deviate from NMRA registered product features, supplier must provide with the bid, a declaration to certify the NMRA accepted product details such as; storage conditions, pack details/contents/sizes and standard batch quantity/size of the product.

13. Immediately after delivery at MRI, the consignments shall be subjected to testing appropriately drawn, one random batch sample (Post-delivery sample) of the consignment at a government/semi-government/accredited laboratory. (to be selectively applied for Surgical & Lab items, depending on availability of testing methodology & facilities) If the sample is found to be substandard, random batch samples will be tested from all the batches/lots in the consignment, and entire expenses on such tests, like value of samples, transport, sampling & testing charges, etc, will be recovered from the supplier.

14. Consignments supplied to MRI violating the storage conditions indicated on product labels and/or product information leaflet (as accepted for product registration at NMRA), shall be considered as quality affected consignments and quality assurance of such consignments shall be carried out by post-delivery testing at government / semi government laboratory in Sri Lanka or at an accredited laboratory (foreign/local). All the expenses on such an event, including storage cost shall be borne by the supplier. If found to be quality affected the consignment will be treated as quality failed (as clause No.11).

Pack size, Labeling & Packaging

15. Offers for pack sizes at a lower level(smaller quantity per pack) than the pack size specified in the item description/specification and MSD order List, are also acceptable, but higher level (larger quantity per pack) pack sizes will not be entertained unless otherwise offered with the original bid and accepted by the procurement committee, with the concurrence of MSD.

16. ~~In respect of bulk packs (not applicable for blister/strip packs), "DHS" mark shall be ;~~

~~(a). embossed or printed in case of tablets~~

~~(b). printed in case of capsules~~

~~Above condition can be waved off, if the quantity in the purchase order is less than 100,000 tablets/capsules, (any exemptions to this condition, is notified in the relevant MSD order list)~~

17. ~~Each; innermost pack, vial/ampoule, pre-filled syringe or bottle, shall bear the item Description, SR No, Batch No/Lot no., Reference/Catalogue no.(not for pharmaceuticals), Date of Manufacture, Date of Expiry and "STATE LOGO" of Government of Sri Lanka.~~
~~It is essential to include and exactly match the dates of Expiry & date of Manufacture (in any form as "Year & Month" or "No Exp."), in the innermost pack and supplier's invoice.~~

18. Description of the Item, SR No, Date of Manufacture, Date of Expiry, Batch No, Name and address of manufacturer and "STATE LOGO" of Sri Lanka Government shall be clearly marked on the outer covering of the individual/innermost pack containing the minimum unit of measure, including blister & strip cards and on the outer cover of the carton/box. Any deviations of the Date of Manufacture (DOM)/ Date of Expiry(DOE)declared in the offer shall be approved by MSD and DOM & DOE shall consist of at least the year & month.

19. All outer most cartons (shipping packages) shall bear the MSD Purchase Order No, SPC Indent No., SR No, Batch No, and Date of Expiry in size 1.5cm letters / figures in prominently visible manner. This may be printed, stenciled or label properly affixed.

20. Batch Number of the product shall be separately Barcoded (in Code 128 or 2D formats) and Barcode shall be printed on the labels at all levels of packing as described below, conforming to the industry standards in Barcode printing and pasting.
Format shall be according to Code 128 or 2D standards.
Maximum barcode size shall be 5.0cm (length) x 2.5cm (width).

21. In case of receiving goods under inappropriate packaging conditions(not in good order), was to be sorted out by MSD to select the items in good order by 100% checking/handling of the consignment, all expenses incurred to MSD in such an event (including demurrage charges, cold stores charges, labor charges etc. or any other charges incurred until goods are ready for acceptance), have to be paid to MSD by the local supplier, before attending to checking the consignment 100%, by MSD.
In respect of SPC imported supplies, if the local agent does not follow suit as above, such extra expenses incurred to MSD shall be recovered from the supplier by SPC and refund to MSD.

Storage Conditions & Temperature

22. If the storage temperature & conditions are not specified in the item specification, NMRA accepted product storage conditions, shall conform to Sri Lankan ambient storage conditions in the ranges of 30^0c +/- 2^0c temperature and 75% +/-5% relative humidity. The product storage conditions shall be clearly indicated at all levels of labels/packages/boxes.

23. Maintenance of Cold Chain;

a. In case of cold storage items, cold chain monitors (temperature recording devices) shall be included for each carton and the cold chain shall be maintained according to the manufacturer's instructions during storage, transport and delivery.

b. Supplier shall use suitable prominently visible identification marks of international standard, with appropriate colours and sizes for easy identification of cold cargo. Supplier shall use standardized **USB Devices** for temperature data logging inside the packages and shall provide free of charge, data logger readers **&/ software (reading apps compatible with Windows-07/latest)** to wharf department of SPC in advance, to enable examining the maintenance of cold chain in transit, and before taking over the consignment by MSD.

c. If the cold chain break is observed at the time of taking over the consignments by MSD, such consignments shall be rejected, indicating the reason on the relevant **WDNor copy of the delivery documents.** In such an event, the SPC shall arrange necessary cold storage for the consignment until

'observed cold chain break' is investigated leading to acceptance / total rejection of consignment and the expenses born by MSD / SPC in arranging the cold storage shall be recovered from the supplier.

d. The vehicles transporting cold cargo to MSD shall be equipped with temperature monitoring devices and the vehicle shall have NMRA approval for transport of pharmaceuticals.

e. The suppliers shall dispatch consignments of the items, which require coldchain maintenance, to arrive in Sri Lanka during Monday to Thursday to avoid additional demurrage & storage charges during weekends, during which MSD stores is closed. In case of non-compliance of this condition, any additional expenses incurred to MSD and SPC, to Custom clear/store/receive such consignments shall be recovered from the supplier.

24. In respect of the products requiring controlled temperature storage (Eg. < 25^0c, 2-25^0c, 15-20^0c/30^0c, 2-8^0c etc.), supplier shall provide MSD with latest product stability study reports with the invoice of the consignment.(report shall include studies; at 30^0c +/- 2^0c & 75% +/- 5% RH for **AC stored** items and at 25^0c +/- 2^0c & 60% +/- 5% RH for **Cold stored** items. It shall be a true copy of the latest report submitted to NMRA or a report issued within last 05 years). (refer clause No.12)

Delivery Requirements

25. All items shall be supplied as per the latest/final delivery schedule, communicated to the supplier, as an amended Indent/PO delivery schedule (if not amended, original schedule in the Indent/PO will apply) mutually agreed between MSD& SPC, at the time of establishing the payment terms (L/C, DP, TT, etc). Any deviation from this agreed delivery schedule shall be treated as a defaulted delivery.

Contravening the above directions, if the delivery schedule is violated by the supplier for no fault of MSD/SPC/MOH and in the event MSD decides to accept any such consignment in full or part thereof, that is delivered after the due delivery date, Condition No. 27 on delayed deliveries, shall be applied.

26. All consignments shall be delivered at Medical Supplies Division or an alternate receiving point as directed. However sending consignments **to reach Sri Lanka from 15th December to 10th January** shall be avoided, unless otherwise prior approval has been granted by MSD for such deliveries.

27. Defaulted consignments with respect to delivery schedule shall only be considered for acceptance, subject to a penalty imposed for the delay due to suppliers fault, allowing a grace period up to two weeks. Consignments delivered after that grace period shall be considered for acceptance subject to a penalty to the supplier as described below ;

(a). A penalty of 0.5% per day of the consignment value, calculated commencing from the 15th day up to 60th day delay from the due delivery date, as per the indent/PO or its' latest amended delivery schedules.

(b). When the delay exceeds 60days purchase order will be considered as automatically cancelled, on defaulted performance. In such a situation, MSD reserve the right to recover liquidated damages or to revoke the cancellation (eg. if payments have been released prior to such a cancellation), and accept the consignment subject to a 25% admin surcharge.

28. (i). If any local purchases were to be made by MSD/SPC to ensure continuity of supply (due to noncompliance of Indent/PO/its' amended; delivery schedule); in the ensuing period inclusive of the grace period for delivery from due delivery date, extra expenditure incurred on such local purchases, over the landed cost of relevant SPC main order, shall be recovered from the supplier.

(ii). If a delivery defaulted (violating delivery schedule in the indent/PO) SPC supplier/his local agent, who participate in an urgent local purchase tender of SPC or MSD for the same item, quoting the same product or any similar product, is bound to supply the local purchase order at the landed cost of the defaulted SPC

main order. In violations of the same, the cost difference will be set off from the payments to the supplier of the corresponding SPC main order.

29. In respect of local manufacturers/ local suppliers, all deliveries shall be made only on week days excluding public holidays, also allowing adequate time to enable the completion of the receiving process at MSD stores before 3.30 p.m. In the event of failure to meet this deadline due to supplier's fault (eg. In delivery; time, product, document, etc.) goods shall be accepted on the following working day, such date shall be counted for working out penalties as per No. 27 (regarding defaulted consignment) of the conditions of supply.

As an alternative, supplier can request MSD to take over the consignment on the same day, subject to settling all adl. expenses (i.e. staff OT, forklift charge, etc.) of MSD, by the supplier.

30. The extension of L/C's overstepping delivery schedules in the Indent/PO/its' amendments, shall not in any way affect the recovery of late delivery charges, as per Condition No. 27 (regarding defaulted consignments) and any other direct or indirect additional costs/liquidated damages, relating/consequent to extension of L/C.

31. When adequate storage space is not available at MSD, to accept a delivery defaulted consignment (deviating from the delivery schedule in the Indent/PO/its' amendments) under the condition No. 27, any additional expenses caused to MSD or SPC in arranging temporary external storage and other expenses (eg. demurrage, detention, container storage, re-handling cum transport, etc.) shall be borne by the supplier.

Documents & Information

32. MSD Order No, Item Description, SR No, Batch No., Date of Manufacture, Date of Expiry and product Storage Condition, shall be indicated in all Supply Invoices and detailed Packing Lists.

33. One of the tender samples of the selected bid shall be forwarded to MSD, for using as a reference sample (can make it; a part of the last consignment or a returnable to supplier) for checking the conformity of the consignments received under the indent/PO.(applicable for all surgical items and regular category of laboratory items, when specified in respective order lists).
The images of the; specimen labels, minimum pack and outer most box/shipper carton, that satisfies the above mentioned labeling conditions, shall also be provided within 14 days of releasing the indent by SPC.

34. The supplier shall submit all shipping documents to (Including Bills of Lading / Draft Air Way Bills etc.) SPC Imports department and MSD by e-mail(**follow instructions in website www.msd.gov.lk**), at least 03 days before the Expected Time of Arrival (ETA) of sea freighted consignments & 02 days before the ETA of Air freighted consignments.

35. After releasing the Indent/PO or establishing L/C, the latest logistical position of manufacturing & supply on the Indent/PO, shall be updated biweekly through e-mails to SPC with a copy to MSD[6] by the supplier.(follow instructions in the website www.msd.gov.lk)
If it is not complied or the information so provided are found to be incomplete/false, the grace period (for supply delays) mentioned in the clause 27 will not be applicable.

Common conditions

36. In addition to the general conditions of supply given herein, item/order-list specific amendments, exclusions or additions to the same, stated in the covering letter of the order list and any other relevant conditions as per the tender document issued by SPC, are also applicable. The order/item specific; new conditions or amendments to General Order Conditions, will be included in the order list itself and as a remark entry in the MSMIS order records.

37. Administrative surcharge of 25%(on the value of goods), will be applied for tender condition violations that cause deficiencies in supply with respect to; quality, standards & specifications, short packing & short supply or delayed delivery as per the cabinet decision. **(eg. As in conditions No. 08,05,10,13)**

Abbreviations :*NMRA ; National Medicines Regulatory Authority/Sri Lanka*, SPC ; State *Pharmaceuticals Corporation, MSD; Medical Supplies Division/Ministry of Health-Sri Lanka.*

Special order conditions sent by NBTS are attached herewith.
Items to be delivered to National Blood Transfusion Service, Colombo 05.

Special Conditions

(I) Suppliers should submit all shipping documents including the Bill of lading or Air Way Bill to SPC at least 2-3 days prior to arrival of the consignments to prevent any delay in clearance.

Demurrage / additional charges if any which become payable due to supplier's failure to comply with this requirement will be claimed from the supplier.

(II) In the event of an award made to you on this tender, SPC reserve the right to cancel/suspend the procuring of said order in any stage, if you would be placed the defaulted supplier's list due to quality failure found in your previous supplies made to SPC or non compliance of contractual agreement.

(III) This bid is administered by the provisions of the "Public Contract Act No. 3 of 1987" and therefore, in the event bidder is to retain an agent, representative, nominee for and on behalf of Bid or shall register himself and such public contact act in accordance with the section 10 of the Public Contract Act and produce such valid original certificate of registration with the bid.

(IV) Where a purchase for a particular item is being made for the first time from a supplier, or where there are previous quality failures on goods supplied by a Particular supplier payments will only be made upon testing the quality and standards of the goods and comparing the bulk supply with the samples provided along with the offer.

(V) Destination Terminal Handling charges (THC) should be borne by the supplier at the Port of Loading. Hence when the C&F prices are quoted this should be inclusive of THC.

(VI) The bid submitted should be duly signed and endorsed by the Bidder/ Tenderer himself (with the name and designation of the signatory) or by the representative. Representatives submit offers on behalf of their principals should submit a letter of authorization and power of attorney (if signing on behalf of the principals) and also should submit documentary proof on their registration as per the Act. No. 03 of 1987 with the Department of the Registrar of Companies – Sri Lanka.

(VII) In the event of delivery of consignments deviating from given delivery schedule by MSD due to default of supplier and same is rejected due lack of storage space available at MSD warehouses, any resulting demurrage charges incurred shall be borne by the suppliers concerned.

(VIII) All Shipment should be made exclusively on vessels belonging to the Ceylon Shipping Corporation or those chartered by CSC. Shipments on other vessels will be permitted in instances where vessels of the Ceylon Shipping Corporation do not call at the Port of Shipment or if they are not available for timely shipment of cargo. In which event the supplier should attach a waiver certificate issued by Ceylon Shipping Corporation on their Authorized Agent in the supplier's country**.**

(IX) Procurement Committee has the authority to decide whether pre-shipment/pre delivery / post delivery samples to be tested. In such cases the supplier will have to bear the cost of testing samples.

(X) The recommended storage mentioned on the product label should be maintained at transit also and storage condition should be clearly showed on Bill of Lading/Airway Bill and Invoice**.**

(XI) a) for products which are imported to Sri Lanka the registration should also be valid until at least six (06) months after the last consignment of the products to be procured are due to be received in Sri Lanka.

b) For products which are manufactured in Sri Lanka the registration should be valid for at least six (06) months after the last consignment of the products to be procured are received by the Procurement Entity.

(XII) Local Agent shall attend to renew the product registration with NMRA six months prior to expiration of the existing product registration.

(XIII) Supplier shall submit the signed contract within 14 days of receiving of the contract agreement from SPC.

(XIV) The below mentioned documents (the original or copies certified by the attorney at law) should be submitted along with copy documents to SPC.

1. **A certified copy of the Certificate of Analysis/warranty**
2. **A certified copy of the Customs declaration**
3. **Original of the Import License**
4. **Original of the Customs Assessment Notice**
5. **Certificate copy of NMRA or WOR.**

Note : A certified copy of Business registration certificate (by an Attorney at Law) should be submitted with the offer.

Annexure

Special –Order Conditions –HLA Histocompatibility Reagents

1. All items to be air freighted and residual shelf lie of the item should be at least 12 months ahead from the date of receipt of reagents in Sri Lanka. In circumstances here the residual shelf life is limited to <12 months (such as manufacturer recommended maximum shelf life is limited as the nature of reagents or production run), the rationale should be provided as reasoned by the manufacturer.

2. Reagents are preferred to contain the certificate of use in the country of origin with one or more of the following criteria.

    1) FDA and/or CE
    2) WHO certification
    3) End user evaluation certificate for the product and /or evidence of registration on one or more of the following countries; USA, UK, France, Germany, Canada, New Zealand, Australia or countries following equitable quality standards.
    4) Reagents are with RUO or IUO licensing considered when no equivalent products with IVD and /or CE certification is not offered and/or accompanying a documentation proving evidence (E.g. Widely acclaimed, peer reviewed publications or user reference amounting to wide usage) of specific benefit for the tests intended (to be evaluated in the local context by the TEC) or on the basis of cost/benefit assessment in the context of user reference)

3. Item should be suitable packed while maintaining the "Cold Chain" throughout transit up until the acceptance from the end user (such as NBC and /or peripheral lab), exactly as recommended/ intended by the manufacturer for each item. Temperatures in particular should be maintained exactly as the manufacturer recommended for each item separately (Proof of manufacturer recommendations on transit conditions should be provided)

4. All reagents intended purpose and limitations declarations, IFUs, worksheets, safety data sheets, quality certificates, brochures, user references and all academic literature supportive of the recommended use and /or comparative advantages, accredited guidelines featuring (E.g., ASHI, BSHI, EFI etc) should be provided, should be in or translated to English language. Academic literature will be evaluated in the context of the quality of publications (Backing of claims by 'larger-scale' met analyses, strength of individual trials, grade of evidence, statistical proofs, appearance in accredited guidance), publication media, evidence of acceptance in wider academia, publications disputing claims by competitive offers etc. by the TEC.

5. If evaluation samples are needed, should be provided within a reasonable time frame (such as 1-2 months of request, as determined by the evaluator), to the address of Director, National Blood Transfusion Services, Colombo -5, for evaluation /re-evaluation and selection. * Statement agreeing to provide evaluation samples in rationalized quantities required by the evaluator if required by the evaluator with terms and conditions should be submitted with the original offer.

6. All items should be eventually delivered to the National Blood Transfusion Service.

7. All assay offers should accompany evidence supportive of (as detailed and extensive as possible) their **comparative merits** (including **comparative advantages in technical characteristics, analysis software characteristics, academic evidence** for the claims) & comprehensive support structure offered (see below). Academic evidence shall be evaluated for standard parameters evaluating such evidence as backing of claims by 'larger-scale' met analysis, strength of individual trials, grade of evidence, statistical proofs, appearance in accredited guidance & industry-wide acceptance by the TEC. Offers will be appraised for the primary assay qualifying for the majority portion procurement & secondary assay qualifying for the minority & overlap portion (for their perceived unique individual merits) of procurement, the ratio of which will be determined by the end-user. (Clinical, Laboratory & administration framework originating the tender)

8. If the need arises, the end-user would reserve the right to individually enhance (independent of each other) either the primary and/or the secondary assay portion procurements, as per the unpredictable nature of reagent consumption, complexities of clinically implementing such comparable, yet uniquely individual assay systems for a single general purpose & cost –benefit dynamics.

9. All offers should accompany, separately be quote & competitively evaluated by the TEC for comprehensive assay application support, specific hardware AND software support & comprehensive continuous capacity building support from technical to clinical application levels in line with global industry standards, **for the total period of usage of the procured assay system quantities by the NBTS.** All offers should have declaration confirming the ability of the end-user to accesses 24hrs & 7 days of local in-person, online global & in-person/in-institution global (within clinically reasonable time frames) application, hardware, software & capacity building support. Technical & capacity building support will be evaluated **at the same level of importance as the assay merits**, while being competitively evaluated for the number, the quantity of events spanning and number of opportunities offered per event. All offers should accompany a list of events & opportunities offered & parameters defining their quality (content delivered and/or experience granted, credentials of presenters, demonstrators and hosts, accreditations, credentials of, global participation & general credentials of participants & historic track-record of delivering such opportunities) & will be critically & competitively evaluated as for **the extremely technical, complex & uniquely proprietary nature of the assays systems, local unavailability of expertise & training resources** (as in molecular histocompatibility & immunogenetics, transplant immunology) **& ultimate impact such capacity building had over the years in clinical application of such platforms.**

10. General SOC conditions will also be further subjected & scrutinized according to workflow specific evaluation criteria (criteria for molecular HLA typing stream vs. screening workflow) when required, and will be published with necessary authorization at the time of tendering.

BID NO: DHS/L/WW/30/25 CLOSING ON : 19.09.2024 at 9.00 a.m.

ORDER LIST NUMBER: 2025/SPC/N/C/D/00026

(A) ITEM NO	(B) SR NO	(C) ITEM	(D) QTY	(E) DELIVERY	(F) Bid Bond value (LKR)
1	52120201	Short Sample Probe for One Lambda Luminex3D & HLA Fusion Software for National Blood Transfusion Service	03 Nos	Mar/2025	
2	52100101	Long Sample Probe for One Lambda LABScan 100 (Model: Luminex 100/200) & XPONENT Software	02 Nos	Mar/2025	
3	43606201	Luminex 200 (performance) Verification Kit {Machine-specific General Purpose Reagent (GPR)} for Luminex corporation xMAP microbead technology-based Luminex 200 (LX-200) compact flow cytometry platform	03 Kits	1 - Mar/2025 1 – Jul/2025 1 – Nov/2025	36,844.98
4	43606101	Luminex 200 (performance) Calibration Kit {Machine-specific General Purpose Reagent (GPR)} for Luminex corporation xMAP microbead technology-based Luminex 200 (LX-200) compact flow cytometry platform	03 Kits	1 - Mar/2025 1 – Jul/2025 1 – Nov/2025	31,818.82
5	43602001	FLEXMAP 3D Performance Verification Kit for One Lambda Luminex3D (Model: FLEXMAP 3D) & XPONENT Software	04 kits	2 - Mar/2025 1 – Jul/2025 1 – Nov/2025	20,297.77
6	43601901	FLEXMAP 3DCalibration Kit for One Lambda Luminex3D (Model: FLEXMAP 3D) & XPONENT Software	04 kits	2 - Mar/2025 1 – Jul/2025 1 – Nov/2025	
7	43601801	XMAP Sheath Fluid for One Lambda Luminex3D (Model: FLEXMAP 3D) & HLA Fusion Software	400L	200L – Mar/2025 200L – Sep/2025	38,101.52

All tenderers should furnish an unconditional Bid Bond for each SR No. encashable on demand to the value mentioned in the Column F.

Amount of Bid Bond should be 2% of the bid value of each item to be submitted along with the bid**, when the tendered value of each item exceeds LKR 01 million. (when not indicated in the Column F).**

Bid Bond should be submitted with valid up to 17.04.2025 together with the tender

Bidding Document Fee- As per the guideline 6.1.1 (a) of the Government Procurement Guidelines 2006.

A non refundable fee of Rs. 3000/= + taxes should be paid in cash to SPC for each set of Tender Documents and attached it to the Bid.

ANNEX – 1

SPECIMEN OF ANNEX – 1

BID NO/ BID REFERENCE :
(TENDER NO)

DATE OF ISSUE :

CLOSING DATE & TIME :
(SRI LANKAN TIME)

ORDER LIST NO :

SR No	Item Description/ Specifications	Quantity	Delivery Schedule

Amount of Bid Bond : LKR …….. or USD …….. to be submitted along with the Bid Bond valid till …… (date)

Bid validity period : Bid should be valid till …….. (date)

Bid Document Fee : ……
(should be paid in cash to SPC for each set of Bid Documents)

MSD CONDITIONS OF SUPPLY

1.
2.

Abbreviations : SPC ; State Pharmaceuticals Corporation, MSD; Medical Supplies Division

Annex VI

SPECIMEN OF CONTRACT FORM (IB)

STATE PHARMACEUTICALS CORPORATION OF SRI LANKA

(Established under the State Industrial Corporation Act, No. 49 of *1957)*
"Mehewara Piyasa", 16th Floor, No. 41, Kirula Road, Colombo 05,Sri Lanka.
Telephone (00)94-1-2335008 Fax: (00)94-11-2582495
E-mail: dgmsurgical@spc.lk or mgrsurgical@spc.lk

DEMOCRATIC SOCIALISTREPUBLIC OF SRI LANKA
AGREEMENT

SPC Ref. No Date :
Bid Ref.

This **AGREEMENT** made and entered into between the State Pharmaceuticals Corporation of Sri Lanka, a Corporation established under the State Industrial Corporation Act. No. 49 of 1957 and having its Head Office at "Mehewara Piyasa", 16th Floor, No. 41, Kirula Road, Colombo 05, Sri Lanka (hereinafter called the "SPC" which term or expression shall mean and include the said State Pharmaceuticals Corporation and its successors and permitted assigns) of the **FIRST PART**

AND

M/s ……..
…business under the time, style and firm of a company duly registered and carrying business (hereinafter called "the supplier" and which term or expression shall mean and include the said and its/their/its heirs executors administrator and permitted assign/successors in business or permitted assigns) of the **SECOND PART.**

AND

M/s ……..
…business under the time, style and firm of a company duly registered and carrying business (hereinafter called "the Local Agent" and which term or expression shall mean and include the said and its/their/its heirs executors administrator and permitted assign/successors in business or permitted assigns) of the **THIRD PART.**

Whereas the State Pharmaceuticals Corporation has accepted the bid of M/s …… …… for the supply and delivery of…… as per the attached indent for …… marked SPC ……. Dated ……and M/s …… will act as local agent of the supplier for all matters arising out of supplies here of.

NOW IT IS HEREBY AGREED AS FOLLOWS:

1. The following documents:

   (a) Conditions of Contract marked – Annex 1
   (b) Bid Documents marked – Annex 2
   (c) Copy of Indent marked – Annex 3

(hereinafter called "the Contract Documents") showing and describing the nature and scope of the agreement duly signed by three parties shall be deemed to form and be read and construed as part and partial of this agreement.

2. In consideration of the payment to be made by SPC to the supplier the contract sum hereinafter mentioned the supplier hereby covenants with SPC to supply and deliver the goods in conformity in all respects with the provisions of this contract, and the local agent will be responsible for all the matters regarding the supplies which do not confirm to required standard..

   The supplier shall be paid for such supply and delivery of the goods according to the Indent marked and in the manner and at the times hereinafter specified.

   This contract as herein before defined constitutes the entire agreement between SPC, the supplier and the local agent may only be modified or repealed by formal agreement in writing duly executed by the parties or their authorized representatives.

   In witness whereof the State Pharmaceuticals Corporation has caused its Common Seal to be affixed and …… and ……. of State Pharmaceuticals Corporation have set their hands and Supplier and the Local Agent has placed its hand/caused its Common Seal to be affixed hereunto and to two other of the same tenor on this ……20…

The Common Seal of M/s (supplier)…….... herein.

1. ……..

President/Managing Director/C.E.O.

2. ……..

Director

Witnesses

Signature	Name, Address and ID No./Passport No.
1. ……	……..
2. ……	……

The Common Seal of M/s. ……(Local Agent) herein.

…….
President/Managing Director/C.E.O.

…….
Director

Witnesses

Signature	Name, Address and ID No./Passport No.
1. ……	……..
2. ……	……

CONDITIONS OF CONTRACT

3.FREE REPLACEMENT /REIMBURSEMENT

3.1 SPC reserves the right to call for Free Replacement/Reimbursement in the event of

3.1.1 Short packing
3.1.2 Loss/damage or deterioration of goods supplied (within shelf-life if applicable)
3.1.3 Packs which cannot be identified due to labels falling off.
3.1.4 Goods supplied fails to perform or meet requirements of the specification/or quality standards to the satisfaction of Medical Supplies Division of Sri Lanka/ State Pharmaceuticals Corporation of Sri Lanka.

3.2 In the event of a quality problem, Batch samples would be tested by SPC its authorized personnel at the NMQAL or SPC Quality Assurance Laboratory or any other Quality Assurance Laboratory nominated by SPC or it's fitness for use will be determined by and expert Committee appointed by the relevant authority.

3.3 **Withdrawal from use of Item due to quality failure.**

a) In case of batch withdrawal due to quality failure, the supplier/ manufacturer shall reimburse the value of entire batch quantity supplied.
b) In case of product withdrawal due to quality failure, the supplier/ manufacturer shall reimburse the value of entire product quantity supplied.
c) In the event of either a) or b) above the supplier/ manufacturer shall be surcharged additional 25% of the total value concerned as administrative cost.

Samples from the available batches will be retained by SPC and the balance will be destroyed by SPC in the presence of the Local Agent and a certificate of destruction issued by SPC.

The supplier and the Local Agent agreed to reimburse the SPC the landed cost and an additional 25% surcharge of the total quantity supplied.

20. FORCE MAJEURE

20.1 The supplier shall not be liable for any delay or failure in making delivery of the supplies if it was due to any event which interfered with performance and was beyond the control of the supplier. However, at every time the supplier faces a situation disturbing the due performance of the obligations under this contract due to conditions beyond his/ its control he/it should write to SPC and get its approval. Approval/disapproval will be notified within 7 working days of receipt of same in writing. Parties however shall endeavours to remove any obstacles to performance (when possible) and co-operate to remove the harmful effects as far as practicable.

21 . NOTICE

21.1 All notices given in respect of this contract shall be deemed to be sufficiently given if sent by

registered post addressed to the parties at the respective addresses at the beginning hereof written.

The common seal of State Pharmaceuticals Corporation of Sri Lanka was affixed)

hereto……)
……)
……)
……)

Authorized signatory

Authorized signatory

Witnesses

Signature Name, Address and ID No

1…….. ……

2…… ……