### **TENDER NOTICE**

No.M/11014/9/21/BB/ Dated: 11.03.2021

Subject: Purchase of the following Blood Bank items for IRCS (NHQ) Blood Bank - reg.

Sealed quotations are invited for purchase of the following items for IRCS (NHQ) Blood Bank at site Indian Red Cross Society Blood Bank, National Headquarters, 1- Red Cross Road, New Delhi -110001, kindly quote your rates for the same.

The sealed quotations, in the name of the Director, Blood Bank, Indian Red Cross Society, National Headquarters, 1-Red Cross Road, New Delhi-110001, should reach before 05.04.2021. Quotation should be marked "Quotation for the purchase of the (Name of the Items) on the envelope. Quotations shall be received only under sealed covers, Quotations received after due date shall not be considered.

Please send the sealed technical bid for the specifications of the items in one envelop and price bid in another envelope. Financial bids of only those vender/firms who are declared qualified for technical evaluation will be opened.

The Competent Authority has the right to reject or accept any or all quotations without assigning any reason whatsoever. Please send the sealed quotation with your lowest rates for the purchase of following items:-

- 1. Haemoglobin Cuvettes (30000 nos.) along-with four machine free of cost (supply in installment)
- 2. Puncher Proof Container = 800 nos. (Small 600, medium 100 & large 100 nos.) (supply in installment)
- 3. New Double distillation water plant (5 to 10 liters)
- 4. Bio Mixer Monitor & Collector = 6 nos.

Rates will valid for two years for items no.1 & 2

The specifications for the above items are given at **Annexure 'A'**.

Security money @10% of total value of the items will be deposited at the time of order letter and Security money will be released without interest after the completion service. No interest will be paid for security money.

# 1

## Specifications - Finger prick, Point of care testing for Hemoglobin

| SI.<br>No. | Name of Specification   | Specifications                                                                                                                                |
|------------|-------------------------|-----------------------------------------------------------------------------------------------------------------------------------------------|
| 1          | System                  | Hemoglobinometer                                                                                                                              |
| 2          | Method                  | Absorbance measurement of whole blood at Hb/HbO2<br>Isobestic Point based on microcuvette technology or any<br>better technology              |
| 3          | Microcuvette            | Should be made of polystyrene plastic either contain<br>active ingredient or non active ingredient                                            |
| 4          | Measuring Range         | 0-25.6g/dL                                                                                                                                    |
| 5          | Measurement             | Dual wavelengths of 500 – 520nm & 850 – 900nm for<br>turbidity compensation.                                                                  |
| 6          | Analyzer                | Analyzer should be USFDA Certified / CE mark                                                                                                  |
| 7          | Measuring Time          | -3seconds - 5 seconds                                                                                                                         |
| 8          | Correlation of Analyzer | Correlation of Analyzer with ICSH reference method with r~0.995                                                                               |
| 9          | Sample Material         | Capillary whole blood / Venous /arterial                                                                                                      |
| 10         | Sample volume           | 3 -10 µL                                                                                                                                      |
| 11         | Calibration             | The system should be factory calibrated and need no further calibration                                                                       |
| 12         | Quality Control         | Built - in self test liquid control should be available                                                                                       |
| 13         | Interface               | RS232 (Printer, PC)                                                                                                                           |
| 14         | Working temperature     | 05 - 40°C                                                                                                                                     |
| 15         | Storage Temperature     | Cuvettes: 10-40°C, Analyzer: 0-50°C                                                                                                           |
| 16         | Power                   | Battery operated                                                                                                                              |
| 17         | Power saver mode        | When operating on battery power, the analyzer will<br>automatically turn off after 5 minutes of no use.                                       |
| 18         | Manufacture             | Manufacture should have ISO 9001:2008 certification/TUV<br>Certificate                                                                        |
| 19         | Analyzer Complies       | Analyzer complies with IVD Medical Device Directive<br>98/79€C                                                                                |
| 20         | Quality Control reagent | All quality control reagents such as low, medium & high<br>control supply free of cost periodically for daily quality control<br>of equipment |
| 21         | Performance certificate | For earlier / recent executed purchase order of reputed,<br>Government / Pvt. Hospital shall be furnished along with<br>the offer.            |

- 22. Technical clearance will be based as precision of the Hemoglobin meter at the Demo site.
- 23. Selection will be based on final technical evaluation.

### Technical specification of Blood Bio Mixer / Monitor and Collector

Input voltage : 100-250 vs C 50 /60 Hz

Electrical Safety Protection against electrical shock class 1 type b internal powered.

Operation : Continuous operation should operate on mains as well as

rechargeable battery on battery it should be operate for a minimum

of 5-8 hours

Power Suitable automatic voltage regulator /stabilizer meeting ISI

specification should be supplied broad specification are automatic. Type input 150-280 volt outup 220 volt +/- 7%, 50 Hz single phase. AC with automatic 2-4 second cut off and 6-9 minutes restart delay.

Working Environment : Temperature : 10"C to 40°C, Relative humidity : 32 - 75%, ATM

pressure 700 to 1060 HPA

Volume & Setting Ranges : In steps of 5ml from 50 to 500 ml automatic storage and recall off set

volume

Alarm Indication LCD, LED indication and audible alarm when the flow rates goes

below 20 ml p/m or above 180 ml p/m. LED, LCD indication and audible alarms when mains power fails. Overload indication LED indication blinking with audible alarm when battery low. Automatic clamping band flow rates sustain in less than 20 ml p/m for more

than 2 minutes.

Manufacturing Standard ISO 9001 2008 EN ISO 13485:2012

Standards and Approval : CE Mark, EURoHS

Medial Devise Calcification : Class I

Continuation Agitation of Blood Bag

during Collector

12-16 p/m

Calibration : Automatic setting / manual setting. Certificate of calibration and

inspection must be provided.

Safety : Design should be meet to all international safety requirement of

EN60601-1 Ensure safety against electrical shock hazards, fire hazards, mechanical hazards, elector maginitic interfreneces etc.

Equipment Carry Case Should be provided by the company of stainless steel for portability

Warranty : Comprehensive on site warranty inclusive of all spares and labour for

five years and five year CMC after warranty.

Performance certificate : For earlier / recent executed purchase order of reputed government

/ private hospital shall be furnished along with the offer

Consumable : All consumable required for installation and standardization of

equipment should be supplied free of cost with equipment.

Americk Slocks

Cornelist 3. 2

### BIO HAZARD

## SPECIFICATIONS FOR SHARP/ PUNCHER PROOF CONTAINERS

- 1. Leak proof, tamper proof, puncture proof containers of virgin plastic quality.
- 2. Capacity 3 liters, 5 liters and 10 liters.
- Non PVC container
- 4. Color Translucent
- 5. Cover of sharp container must have provisions for dropping/removing of blades, needles syringes with non removable needles etc.
- 6. Multiple use Cover must be open able and having a sturdy lock system. Container must be complied with AN ISO 90001:2008 / CE.
- 7. A non washable and non-removable label with Bio-Hazard symbol in red color and word Blo-hazard should be prominently displayed.
- 8. A warning of filling level with a line also be marked on label. Label also carried a warning: DO NOT TILT THE CONTAINER WHEN FULL.
- 9. Performance Certificate for earlier / recent executed purchase order of reputed. Government / Pvt. Hospital shall be furnished along with the offer.



BIO - HAZARD

DO NOT TILT THE CONTAINER WHEN FULL.

### Technical specification of Double Distillation Water Plant



Description of functions

Required for distilled water production for lab

Operational Requirements

- Double distillation plant with stand not wall mounted and approximately 5 to 10 litres p/h output.
- 2. Instant distilled water flow should be there.
- 3. Easy to operate, durable, safe for routine use.

Technical specification

- 1. Quartz distiller, demountable boiler.
- Panel box and stand to accommodate regulator and electrical supply, clamps etc.
- Quality of distillate pyrogen free, PH 6.9 7.0 high purity, low conductivity.
- 4. Distilled water should be heavy metals, salts, pyrogen and iron free.
- Specific conductivity at 25 degree Celsius less than 0.4 multiply 10-2 65 / CM
- 6. Equipment should be thermal shock proof
- Gas vent should be there to remove volatile impurities leaving the condensate free from gaseous impurities
- 8. Automatic low water cut off
- 9. Tubing should be made of good quality rubber (heat resistant)
- 10. Wiring of the equipment should be enclosed in case.
- It should have the concentrator a bleeder device on the evaporation that constantly removes a part of the boiling water from its so that the cumulate concentration of non volatile impurities in the water is prevented.

System Configuration, Accessories, Spares And Consumable Systems as specified.

- The boil chamber and condensing coil should be high grade 304 stainless steal.
- All consumable required for installation and standardization of system to be given free of cost.
- c) Low temperature circulating water bath (Chiller)
- d) Working temperature 10°C to 100°C
- e) Temperature Accuracy +/- 0.1 °C

**Environmental Factors** 

The unit should be capable of being stored continuously in ambient temperature of 0-50 degree Celsius and relative humidity of 15 - 90% the unit shall be capable of operating continuously in ambient temperature of 10 - 40 degree Celsius and relative humidity of 15 - 90%.

Power Supply

: Power input to be 220-240 VAC, 50 Hz fitted with Indian plug.

Voltage corrector / stabilizer of appropriate rating meeting ISI specification input 160-260 volt and output 220-240 volt and 50 htz.

Standard Safety and Training Should be FDA/ CE/ UL BIS approved product.

Should be compliant to ISO:13485:2012, quality system medial device particular requirement for the application for ISO: 9001 applicable to manufacturer and service provider that perform their own design activities.

Consumable

All consumable required for installation and standardization of equipment should be supplied free of cost with equipment.

Safety

Design should be meet to all international safety requirement of IEC 60601 or better general requirement. Ensure safety against electrical shock hazards, fire hazards, mechanical hazards, elector maginitic interferences etc.

Warranty

Comprehensive on site warranty inclusive of all spares and labour for five years and five year CMC after warranty.

Performance certificate

For earlier / recent executed purchase order of reputed government / private hospital shall be furnished along with the offer/

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