Republic of the Philippines
Department of Health
METRO MANILA CENTER FOR HEALTH DEVELOPMENT

SUPPLEMENTAL/ BID BULLETIN NO. 1

IB 2022 – 118E
PROCUREMENT OF 13 UNITS AUTOCLAVE 20L

This Supplemental/Bid Bulletin No. 1 is being issued to revise provisions/specifications in the Bidding Documents for a forecited project:

Revision and clarification to provisions/specifications in the Bidding Documents:	
FROM	**TO**
REQUIREMENTS IF AWARDED THE CONTRACT: 1. Completion Period: The delivery, installation, testing and commissioning of the equipment and its accessories, including the training of end-users and maintenance staff must be completed within 30 calendar days upon receipt of Notice to Proceed.	**REQUIREMENTS IF AWARDED THE CONTRACT:** 1. Completion Period: The delivery, installation, testing and commissioning of the equipment and its accessories, including the training of end-users and maintenance staff must be completed within 30 to 60 calendar days upon receipt of Notice to Proceed.

Bidders are advised to use the following attached forms and submit together with all required documents for the submission of bids on 30th day of August 2022, 9:00 AM:

This Supplemental/Bid Bulletin No. 1 shall form an integral part of the Bidding Documents. All other provisions indicated in the bidding documents which are not affected by this Supplemental/Bid Bulletin No. 1 shall remain in effect.

For guidance and information of all concerned.

Issued this 22nd day of August 2022 in MMCHD

Approved by:

SGD.
ALELI ANNIE GRACE P. SUDIACAL, MD, MPH
Director III / BAC Chairperson

Republic of the Philippines
Department of Health
METRO MANILA CENTER FOR HEALTH DEVELOPMENT

Republic of the Philippines Department of Health Metro Manila Center for Health Development TECHNICAL SPECIFICATIONS			
Item No. **1**	**AUTOCLAVE 20L**	Qty./Unit	**13 Units**
Name of Manufacturer:			Country of Origin
Brand:			Model: (if applicable)
ABC: **780,000.00**			
PURCHASER'S SPECIFICATION			STATEMENT OF COMPLIANCE
Technical Specifications: · Chamber: Stainless Steel, Horizontal · Chamber Capacity: At least 20 Liters · Power Supply: 220V, 60Hz, Single-phase · Heater: At least 1kW · Safety Devices: o Low Water Cutoff Switch o Safety Release Valve o Emergency Exhaust Valve o Door Safety Lock o Pressure Gauge with Pressure Control Switch o Steam Trap o Timer with Alarm · Standard Accessories: Stainless Steel Tray			
DOCUMENTARY REQUIREMENTS: 1. Product brochure or technical data sheet(s) of the equipment showing the technical specifications in English Language. 2. Valid and current Certificate of Compliance of manufacturer of the equipment with the latest version of ISO 13485: Quality Management System – Requirements for regulatory purposes in the name of the manufacturer. The Certificates must be issued by an independent Certifying Body/Agency. 3. Valid Certificate of Distributorship (as first Tier Distributor) issued by			

the Manufacturer of each equipment authorizing the bidder to sell/distribute the offered equipment. 4. Notarized Certificate of the bidder: a. That the brand of the equipment has been in the local and/or international market for at least ten (10) years. b. That the equipment and its accessories are brand new, unused, not discontinued models and were not subjected to any product recall. c. That the supplier has the capability for corrective and preventive maintenance of the unit within the warranty period. 5. Bidder's valid and Current License to Operate (LTO) as a medical device distributor issued by the Philippine Food and Drug Administration. In case of expired LTO, the following must be submitted: i) Copy of expired LTO, ii) Application for renewal, iii) Official Receipt as proof of payment for the renewal of LTO.	
REQUIREMENTS IF AWARDED THE CONTRACT: 1. **Completion Period**: The delivery, installation, testing and commissioning of the equipment and its accessories, including the training of end-users and maintenance staff must be completed within **30 to 60** calendar days upon receipt of Notice to Proceed. 2. **Testing:** Prior to acceptance, the end user shall conduct a physical inspection and functionality test. The equipment must be functioning and must have no physical damage and defect. 3. **Training**: The supplier shall provide a training on the proper use and maintenance of the equipment to the end-users and to the hospital maintenance staff within (3) days upon delivery of the equipment. 4. **Warranty**: Warranty certificate for two (2) years on parts and service. The supplier shall either repair or replace any item or part in the equipment that is found to be defective in material or in workmanship under normal use. The warranty period shall commence from the date of acceptance by the end-user after testing and commissioning. 5. **Notarized undertaking** that the supplier shall conduct the necessary corrective maintenance within five (5) calendar days upon notification of the equipment breakdown from the end-user. The undertaking shall include a statement that the number of days where the equipment is unusable due to defective material or workmanship, shall be added to the warranty period.	

6. **Manuals**: The supplier must provide the end-user one (1) hard and one (1) soft copy of the following: a) Service manual in English language b) Operation manual in English language 7. With "**DOH-MMCHD HFEP**"(Government Property not for sale) sticker in each unit. x-x-x-x-x-x-x-x-x-x-x-x-x-x-x-x-x-x-x-x-x-x-x-x-x-x-x-x-x-x-x-x-x-x-x-x-x-x-x-x-x **Recipient:** Valenzuela City Emergency Hospital – 3 units Muntinlupa – 7 units Health Facilities in Muntinlupa City - 3 units	
Additional Requirement to be submitted by the Single/Lowest Calculated Bidder (SCB/LCB) as part of post qualification: 1. One (1) original sample of manufacturer's product to be submitted and returned after evaluation. The sample submitted and approved during the evaluation shall be the same item to be delivered upon award of contract. Prototype of the labelling instruction must be part of the sample submitted however, the technical specifications of the labelling instruction of the product must be complied upon delivery.	