



Republic of the Philippines
Department of Health

METRO MANILA CENTER FOR HEALTH DEVELOPMENT

SUPPLEMENTAL/ BID BULLETIN NO. 1

IB 2022 – 023E

PROCUREMENT OF LEVOTHYROXINE 50 MCG/25 MCG/ 12.5 MCG (LOT BIDDING)

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:	
ITEM No. 1	
ABC: P 367,500.00	
FROM	TO
IB 2022-04E	IB 2022-023E
ITEM No. 1 Sterilizing Temperature: 40°C, 125°C, -5°C	Sterilizing Temperature: 40°C, 125°C
ITEM No. 2 Voltage: 110V or 220V/50Hz or customized	Voltage: 220V/ 60Hz

Bidders are advised to use the following attached forms and submit together with all required documents for the submission of bids on 4th day of April 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 25th day of March, 2022 in MMCHD

Approved by:


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

Technical Specifications

Item	Specification	Statement of Compliance
		<p><i>[Bidders must state here either “Comply” or “Not Comply” against each of the individual parameters of each Specification stating the corresponding performance parameter of the equipment offered. Statements of “Comply” or “Not Comply” must be supported by evidence in a Bidders Bid and cross-referenced to that evidence. Evidence shall be in the form of manufacturer’s un-amended sales literature, unconditional statements of specification and compliance issued by the manufacturer, samples, independent test data etc., as appropriate. A statement that is not supported by evidence or is subsequently found to be contradicted by the evidence presented will render the Bid under evaluation liable for rejection. A statement either in the Bidder's statement of compliance or the supporting evidence that is found to be false either during Bid evaluation, post-qualification or the execution of the Contract may be regarded as fraudulent and render the Bidder or supplier liable for prosecution subject to the applicable laws and issuances.]</i></p>
<p>IB 2022- 023E</p>	<p>PROCUREMENT OF VARIOUS MEDICAL EQUIPMENTS (LOT BIDDING) (REBID) TOTAL ABC: P 667,500.00</p> <p>ITEM 1 21 UNITS OF STERILIZING UNIT ABC: P 367,500.00</p> <p>TECHNICAL SPECIFICATIONS: 2-in-1 Dry Heat/Ultraviolet Sterilizer Power Supply: 220V, 50-60Hz Rated Power: 500W Loading Limit: at least 5 kg Sterilizing Temperature: 40°C, 125°C Dimensions: Manufacturer's Standard Stainless Steel Body Frame Three-Layer Tray each Side Build Timer: 30-60 mins (Dry Heat) Can be wall-mounted Splash Proof Heater Bulb With "DOH-MMCHD HFEP" (Government Property not for sale) sticker in each unit</p> <p>DOCUMENTARY REQUIREMENTS: 1. Product brochure or technical data sheet(s) of the equipment showing the technical specifications in English Language.</p>	

Muntinlupa - 16 units

ITEM 2

20 UNITS PORTABLE UVC LIGHT STERILIZER

ABC: 300,000.00

TECHNICAL SPECIFICATIONS:

Widely used for sterilization, odor, and mite control in home, clinic, school, kindergarten, hotel, office, vet, spa, and regimen club, etc.

Voltage: 220V/ 60Hz

Power: 40W

Lamp Lifespan: 8,000 to 12,000 hours

Time Setting: 3-mode settings with remote control

Dimensions: Manufacturer's Standard

With "DOH-MMCHD HFEP" (Government Property not for sale) in each unit.

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. Proof (such as sales invoice) that the Brand of the equipment has been sold to other health facilities in the Philippines.
5. 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.
6. 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 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.

