Republic of the Philippines
DEPARTMENT OF HEALTH
Metro Manila Center for Health Development

BAGONG PILIPINAS

SUPPLEMENTAL/ BID BULLETIN NO. 1

IB 2024 – 095E
PROCUREMENT OF 5 UNITS RADIANT WARMER (REBID)

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

1. Query during Pre-bidding Conference:

Technical Specifications	Query	Response of the End User Unit
Ceramic heater elements and golden reflectors	Ceramic heater elements and silver reflectors.	Ceramic heater elements and silver reflectors. GRAMTED
With at least 2 heater elements located at the Radiant warmer.	With at least 1 heater element located at the Radiant warmer	With at least 1 heater element located at the Radiant warmer GRANTED
Lamps (Work Light): 30 Watts • Lamps (Night Light): 9 Watts	Lamps (Work Light): 6 watts to 30 Watts Lamps (Night Light): 6 watts to 9 Watts	Lamps (Work Light): 6 watts to 30 Watts GRANTED Lamps (Night Light): 6 watts to 9 Watts GRANTED
Accuracy (Sensor): ±0.1 degrees Celsius - Setting Range: 30 degrees Celsius to 38.5 degrees Celsius	Accuracy (Sensor): ±0.1 degrees Celsius - Setting Range: 30 degrees Celsius to 38.5 degrees Celsius	Accuracy (Sensor): ±0.2 degrees Celsius Setting Range: 34.5 degrees Celsius to 37.5 degrees Celsius GRANTED

Bidders are advised to use the **following attached forms and submit them together with all required documents for the submission of bids on the 20th day of November 2024, 9:00 AM:**

This Supplemental/Bid Bulletin No. 1 shall be integral to the Bidding Documents. All other provisions indicated in the bidding documents not affected by this Supplemental/Bid Bulletin No. 1 shall remain in effect.

For guidance and information of all concerned.

Issued this 12th day of November 2024 in MMCHD

Approved by:

JEREMIAS FRANCIS Y. CHAN, MD
Licensing Officer V / BAC Chairperson

Republic of the Philippines

Department of Health

Metro Manila Center for Health Development

TECHNICAL SPECIFICATIONS

Item No. **1**	**RADIANT WARMER**	Qty./Unit	**5 Units**
Name of Manufacturer:			Country of Origin
Brand:			Model: (if applicable)
ABC: **1,250,000.00**			

PURCHASER'S SPECIFICATION	STATEMENT OF COMPLIANCE
TECHNICAL SPECIFICATIONS: • Movable Lamp mechanism that automatically focuses radiant warmth on the baby. • Lamp can be positioned on the overhead left or right side of the equipment. • *Ceramic heater elements and silver reflectors.* • Radiant warmth with Two (2) brightness level Examination lamp features. • With at least 10 heater level. • Radiant Warmer can be positioned from left to right direction for up to 90 degrees. • *With at least 1 heater element located at the Radiant warmer* . • *Lamps (Work Light): 6 watts to 30 Watts* *Lamps (Night Light): 6 watts to 9 Watts* . • Radiant Power at Heat Level 3 between the bed and radiant warmer: 10mW/cm2 • Radiant Power at Heat Level 10 between the bed and radiant warmer: 30mW/cm2 • Lamps (Work Light): 30 Watts	

• Lamps (Night Light): 9 Watts • Minimum clearance between the top edge of the Radiant warmer and Ceiling: >50cm • Temperature - Capable of measuring baby's central or peripheral temperature. - Manual mode - Skin servo mode - With a temperature measuring range of at least 15 degrees to 42 degrees - *Accuracy (Sensor): ±0.2 degrees Celsius* - *Setting Range: 34.5 degrees Celsius to 37.5 degrees Celsius* • Variable Height Adjustment (VHA): At least 290mm • Standard Features - Swivel cupboards located at the right or left side of the machine. - Manual bed-tilt feature Head-up position is 20 degrees. - Manual bed-tilt feature Head down position is 15 degrees. - The inner side walls must be 7cm. - Equipped with a Standard X-ray tray. • Accessories - One (1) unit Skin temperature probe - One (1) unit Power cable - One (1) unit Power adapter	
Requirements if awarded the Contract 1 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 with 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. 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 the delivery of the equipment. 4**. Warranty** a) 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 workmanship under normal use. The warranty period shall commence from the date of acceptance by the end-user after testing and commissioning. b) Preventive maintenance at least every six (6) months or according to the manufacturer's recommendations; c) Corrective maintenance within five (5) calendar days upon notification from the end-user regarding equipment breakdown/defects. d) The number of days where the equipment is unusable due to equipment defects/faults shall be added to the warranty period. e) The supplier shall specify post-warranty comprehensive preventive maintenance costs including list and prices of major spare parts of the equipment for three (3) years after the warranty period. 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 Source of Fund: SAA 2023-02-000687 (HFEP 2023 ConAp) Recipient: Caloocan City Medical Center	

Republic of the Philippines Department of Health Metro Manila Center for Health Development TECHNICAL SPECIFICATIONS			
Item No. **1**	**INFANT INCUBATOR**	Qty./Unit	**2 Units**
Name of Manufacturer:			Country of Origin
Brand:			Model: (if applicable)
ABC: **4,200,000.00**			
PURCHASER'S SPECIFICATION			STATEMENT OF COMPLIANCE
TECHNICAL SPECIFICATIONS: • Intensive care incubator for treatment of premature and newborn babies • Can accommodate premature and full-term babies up to a maximum of 10kg (22 lbs.) • Has manual and automatic humidity adjustment • Double-wall with automatic air curtain to support thermoregulation • *Storage volume (cabinet (30 L): not less than 80 L* • Has ventral skin-to-skin contact (SSC) mode: - maintains the ideal microclimate inside the incubator while the neone is undergoing skin-to-skin care - with display in elapsed time and central peripheral temperature for			

continuous monitoring of staff • Has its condensate from the incubator compartment isolated from the clean water supply of the humidity system - The entire humidity system can be easily removed for quick, convenient yet effective hygienic reprocessing after every patient • Has dual-skin temperature monitoring • With variable height adjustment • With keypad lock function • Has 7-day trend for weight gain and loss • With integrated xray • 24-hour trend • Air temperature and skin temperature • Relative humidity, Oxygen Concentration, Heater Power • Has central temperature probe • Peripheral Temperature Probe • Temperature control modes Skin and Air Temperature Control Mode • Air temperature mode set point range 20.0˚C (68.0˚F) to 39.0˚C (102.2˚F) • Air temperature mode set point override temperature range 37.0˚C (98.6˚F) to 39.0˚C (102.2˚F) • Skin temperature mode set point range 34.0˚C (93.2˚F) to 37.0˚C (98.6˚F) • Skin temperature mode set point override temperature range 37.0˚C (98.6˚F) to 38˚C (100.4˚F) • Temperature rise time at 22˚C (72˚F) ambient <35 min • Operating noise level in hood <47 dBA (with servo Oxygen Control) • Humidity control ranges from 30 to 95% in 1% increments • Humidity control capacity 1,500 ml • Accessories: - Storage - Utility shelf - IV pole • Device Classification - Protection class: Class I, Type BF, Continuous Operation	

- Ingress of liquids IPX0	
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 with 60-90 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 the delivery of the equipment. 4**. Warranty** a) 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 workmanship under normal use. The warranty period shall commence from the date of acceptance by the end-user after testing and commissioning. b) Preventive maintenance at least every six (6) months or according to the manufacturer's recommendations; c) Corrective maintenance within five (5) calendar days upon notification from the end-user regarding equipment breakdown/defects. d) The number of days where the equipment is unusable due to equipment defects/faults shall be added to the warranty period. e) The supplier shall specify post-warranty comprehensive preventive maintenance costs including list and prices of major spare parts of the equipment for three (3) years after the warranty period. 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-	1.

Source of Fund: SAA 2023-02-000687 (HFEP 2023 ConAp) Recipient: CALOOCAN CITY MEDICAL CENTER	