



BID BULLETIN NO. 1
18 JUNE 2026

Public Bidding No. VMC-2026-050

SUPPLY, DELIVERY, INSTALLATION, ACCEPTANCE TESTING AND COMMISSIONING OF VARIOUS MEDICAL EQUIPMENT FOR OB-GYN, OPERATING/DELIVERY ROOM AND DERMA – BATCH 3

Issued pursuant to Sec. 51.5 of the IRR of RA 12009 to clarify and/or amend certain provisions in the Bidding Documents issued for this project, considering the issues raised and clarifications made by the prospective bidders during the Pre-bid Conference held on June 18, 2026, likewise, respond to bidder’s written queries received within the prescriptive period for filing.

Posted in Bidding Documents	Amendment
1. SUPPLY, DELIVERY, ACCEPTANCE, TESTING AND COMMISSIONING OF ELECTROSURGICAL UNIT	
Technical specifications:	Technical specifications:
Remote power control	Remote power control or pen control
- monopolar coagulation modes: spray, standard, and pinpoint	- monopolar coagulation modes: spray, standard, and pinpoint or its equivalent
ELECTRICAL REQUIREMENTS	ELECTRICAL REQUIREMENTS
With compatible Uninterruptible Power Supply (UPS) with voltage regulation function that can provide back-up power at least 30 minutes	With compatible Uninterruptible Power Supply (UPS) with voltage regulation function that can provide back-up power at least 30 minutes, (Mounted on a heavy-duty, compatible mobile cart for transport "OR" designed to be securely mounted within the Electro-surgical Unit (ESU) cart, and equipped with a grounding power cord of at least 5 meters in length.)
GENERAL REQUIREMENTS	GENERAL REQUIREMENTS
Certification that needs to be presented during Post-Qualification:	Certification that needs to be presented during Post-Qualification:
With valid FDA Certificate of Medical Device Notification (CMDN)/Certificate of Medical Device Registration (CMDR) or pending renewal at least one (1) month before expiration	With valid FDA Certificate of Medical Device Notification (CMDN)/Certificate of Medical Device Registration (CMDR) or pending renewal at least one (1) month before expiration Submit the following as proof of renewal application: - Proof of payment - Acknowledgement Receipt (e.g. FDA DocTrack)
The equipment or devices must conform to the IEC 60601 Standards and any of the following Standards: a. USFDA (Food and Drug administration,US Standards b. CE(European conformity) Standards c. ISO Standards (9000, 9001, 9002) d. UL (Underwriters Laboratories) Standards e. Theproposed unit shall also conform to all relevant International, National and local standards and requirements of medical device regulatory agencies. Supplier shall list in the various international standards met by the supplier	REMOVED

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Address: **Padrigal St., Karuhatan, Valenzuela City, 1441**
Telephone Nos: **8294-6711 to 17**
Director's Office Direct Line: **8291-4259**
Email Address: **valgen_hosp@yahoo.com**
Website: **https://vmc.doh.gov.ph/**

NONE	The equipment or devices shall conform to the IEC 60601 (Medical Electrical Safety Standards) or its internationally recognized equivalent.
NONE	The manufacturer of the medical equipment shall possess a valid ISO 13485 Certification (Medical Devices – Quality Management System) or its localized national equivalent.
NONE	The manufacturer of the medical equipment shall comply with ISO 14971 (Risk Management for Medical Devices) or its localized national equivalent.
Procedures to be performed upon delivery:	Procedures to be performed upon delivery:
Conduct Calibration / Verification Test performed by the supplier	Conduct Calibration / Verification Test performed by the supplier, and shall be witnessed by the designated End-User and the VMC Medical Equipment Technician
2. Supply, Delivery, Acceptance, Testing and Commissioning of SMOKE EVACUATOR	
GENERAL REQUIREMENTS	GENERAL REQUIREMENTS
Certification that needs to be presented during Post-Qualification:	Certification that needs to be presented during Post-Qualification:
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3. Supply, Delivery, Installation, Acceptance, Testing and Commissioning of Obstetrical Table	
Technical Specifications:	Technical Specifications:

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Weight Capacity: Minimum 180 kg or more <i>maximum carrying capacity</i>	Weight Capacity: up to 300 kgs.
Accessories:	Accessories:
-One (1) set consisting of three (3) individual mattresses. Material: -high density sponge (foam) with leather coating for easy disinfection -thickness at least 4"	-One (1) set consisting of three (3) individual mattresses. Material: -high density sponge (foam) with leather coating for easy disinfection -thickness at least 3-4"
One (1) Power Cable	One (1) Power Cable (5 meters)
GENERAL REQUIREMENTS	GENERAL REQUIREMENTS
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NONE	The manufacturer of the medical equipment shall possess a valid ISO 13485 Certification (Medical Devices – Quality Management System) or its localized national equivalent.
NONE	The manufacturer of the medical equipment shall comply with ISO 14971 (Risk Management for Medical Devices) or its localized national equivalent.
OTHER TERMS & CONDITION:	OTHER TERMS & CONDITION:
Delivery: Sixty (60) calendar days from the date of receipt of Purchase Order	Delivery: Ninety (90) calendar days from the date of receipt of Purchase Order
4. Supply, Delivery, Acceptance, Testing and Commissioning of OB-GYN ULTRASOUND MACHINE	
Certification that needs to be presented during Post-Qualification:	Certification that needs to be presented during Post-Qualification:
With valid FDA Certificate of Medical Device Notification (CMDN)/Certificate of Medical Device Registration (CMDR) or pending renewal at least one (1) month before expiration	With valid FDA Certificate of Medical Device Notification (CMDN)/Certificate of Medical Device Registration (CMDR) or pending renewal at least one (1) month before expiration

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	Submit the following as proof of renewal application: - Proof of payment - Acknowledgement Receipt (e.g. FDA DocTrack)
The equipment or devices must conform to the IEC 60601 Standards and any of the following Standards: a. USFDA (Food and Drug administration, US Standards) b. CE (European conformity) Standards c. ISO Standards (9000, 9001, 9002) d. UL (Underwriters Laboratories) Standards e. The proposed unit shall also conform to all relevant International, National and local standards and requirements of medical device regulatory agencies. Supplier shall list in the various international standards met by the supplier	REMOVED
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NONE	The manufacturer of the medical equipment shall possess a valid ISO 13485 Certification (Medical Devices – Quality Management System) or its localized national equivalent.
NONE	The manufacturer of the medical equipment shall comply with ISO 14971 (Risk Management for Medical Devices) or its localized national equivalent.
Procedures to be performed upon delivery:	Procedures to be performed upon delivery:
Verification Test performed by the supplier	Verification and Acceptance Testing: The supplier shall perform comprehensive functional and image quality verification testing on-site. This testing must utilize a standardized ultrasound phantom test tool to evaluate axial/lateral resolution, depth of penetration, and distance accuracy. The entire testing procedure must be physically witnessed and formally signed off by both the end-user clinical representative and a VMC Medical Equipment Technician prior to final acceptance.
Other Terms & Condition:	Other Terms & Condition:
Delivery: Sixty (60) calendar days from the date of receipt of Purchase Order (PO)	Delivery: Ninety (90) calendar days from the date of receipt of Purchase Order (PO)

BID OPENING: JULY 2, 2026 @ 9:00AM

All other provisions of the bidding documents not affected by revisions included in this Bid Bulletin shall remain in effect. Amendments/inclusions/clarifications made herein shall be considered an integral part of the Bidding Documents.

Prepared by:

MS. KRISTINE JOY MANUEL
BAC Secretariat

Noted by:

MS. RUBY S. GURREA, RN, MAN
Chairperson, BAC II

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