



Bid Notice Abstract

Request for Quotation (RFQ)

Reference Number13261479

Procuring EntityVALENZUELA MEDICAL CENTER

TitleSupply, Delivery, Acceptance, Testing and Commissioning of one (1) unit Video Laryngoscope

Area of DeliveryMetro Manila

Solicitation Number:	VMC-2026-200	Status	Active
Trade Agreement:	Implementing Rules and Regulations		
Procurement Mode:	Negotiated Procurement - Small Value Procurement (Sec. 34)	Associated Components	1
Classification:	Goods	Bid Supplements	0
Category:	Medical and Dental Equipment		
Approved Budget for the Contract:	PHP 765,000.00	Document Request List	2
Delivery Period:	30 Day/s		
Client Agency:		Date Published	29/09/2026
Contact Person:	KRISTINE MANUEL BAC Secretariat Padrigal St., Karuhatan, Valenzuela City Metro Manila Philippines 63-2-82944625 Ext.000 vmc_bac2026@yahoo.com	Last Updated / Time	29/09/2026 00:00 AM
		Closing Date / Time	02/10/2026 10:00 AM

Description

PR# 26-09-1149

ITEM NO. QTY. UNIT ITEM DESCRIPTION UNIT COST TOTAL COST

1 1 unit Supply, Delivery, Acceptance, Testing and Commissioning of one (1) unit Video Laryngoscope 765,000.00 765,000.00

Description:

- is a modern medical device used to facilitate endotracheal intubation by providing a real-time, high-definition view of the airway

Features:

- Portable host with at least 2.7" LED touchscreen with resolution of at least 640x480 or 400x400 pixels
- Monitor can be tilted >130° and rotated up to 270°
- Multi-media function: Capable for record video with sound and take photo / play & playback the photos and videos
- Memory: At least 8gb built-in memory
- Comes with case and various specifications of metal blades (Newborn/Pediatric/Adult/difficult intubation)
- Real time anti-fog; turn on and intubate, no need preheating or waiting

Accessories: Manufacturer's Standard including the following:

- Leakage tester-1pc
- Waterproof plug-2pcs
- Vent Cap-1pc
- Aluminum case-1pc
- Power Adapter-1pc
- Micro- USB Cable-1pc
- Metal Blades
- o Newborn
- o Pediatric
- o Adult
- o Difficult Intubation

General Requirements:

Electrical Requirements

- With rechargeable Lithium-Ion battery pack min. 6,400 mAh up to 240 working time or equivalent
- The unit must automatically charge the integrated battery while connected to AC mains power

Certification that needs to be presented during Post-Qualification / Evaluation:

- 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. 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
- Training certificate from manufacturers of field service engineers / technician performing Preventive and Corrective Maintenance and Calibration Services
- The bidder/manufacturer shall submit a valid certification and complete address details confirming the existence of an established and operational service center in Metro Manila with corresponding names, contact number and email address
- Certificate of Comprehensive Warranty which states the following clause:
 - a. The period of warranty shall be deemed to be fully comprehensive
 - b. All inclusive of warranty, labor, spare parts, accessories, service consumables, manufacturer's Preventive Maintenance
- Certificates to be submitted upon delivery:
- Warranty and Preventive Maintenance Services:
 - One (1) year comprehensive warranty on parts and services for equipment
 - Certificate of availability of spare parts within ten (10) years
 - Certification that the 95% uptime of the product is within the warranty period and that any accumulated downtime in excess of 5% shall be added to the warranty period.
 - Submit Semi-Annual Preventive Maintenance Schedule and Annual Calibration Service Schedule during warranty period (include month and year)
 - Calibration Report from the Manufacturer or Verification Report from the Supplier (2 copies, 1 copy for MMS and 1 for EFMS)
 - Certification that the bidder will provide a Service Unit that the end-user can use in case the equipment or any system component will be pulled-out for repair or maintenance within the warranty period.
 - Submit complete parts, accessories and consumables list with identification number or codes
 - Free software upgrades and updates (security, revision updates) for all Clinical and Technical applications during the period of warranty.
- Procedures to be performed upon delivery:
 - Training of End-User (Operation) with Certificate of Training should be provided and should contain the following details:
 - a. Name of Trainee
 - b. Modality, Brand, Model of Equipment
 - c. Type of Training Conducted
 - d. Inclusive Dates of the Training
 - e. Name of Trainer, Date and Venue
 - The unit to be delivered must show proof that its manufacturing date must be CY 2024 onwards
 - Verification Test performed by the supplier
 - Electrical Safety Verification (IEC 62353)
 - Provide two (2) sets of colored Technical Manual, Hardcopy and Softcopy (Flashdrive) in English Manual for all equipment including peripherals, etc.
 - a. User's Operational Manual
 - b. Quality and Maintenance Manual
 - c. Service and Technical Manual
 - The supplier must conduct training for the hospital's biomedical technicians and unit head covering all equipment, accessories, and peripherals. The supplier must perform and guide staff through the actual:
 - a. Standard Operations
 - b. Disassembly and Assembly
 - c. Troubleshooting
 - d. Recommended Maintenance (as per manufacturer guidelines)
- Other Terms & Condition:
 - Preferably with green eco-products specification (made of recyclable and non-toxic materials, energy efficient, sustainability, etc.)
 - Delivery: Thirty (30) calendar days from the date of receipt of Purchase Order (PO)

Total 765,000.00

A copy of your legal requirements is also required to be submitted along with your quotation/proposal:

- Certificate of Registration from BIR
 - Securities & Exchange Commission (SEC)/Department of Trade & Industry (DTI) Registration
 - Business/Mayor's Permit with O.R. renewal
 - Platinum PhilGEPS Certificate
 - Tax Clearance Certificate
 - Original Notarized Hard copy of Omnibus Sworn Statement
 - Brochure/Data Sheet
 - LTO from FDA
 - *Certificates that needs to be presented during Evaluation*
 - 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
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 - a. The period of warranty shall be deemed to be fully comprehensive
 - b. All inclusive of warranty, labor, spare parts, accessories, service consumables, manufacturer’s Preventive Maintenance

Note: The Bidder will submit also a Guarantee letter stating that the certificates to be submitted upon delivery will be provided.

Other Information

MS. RUBY S. GURREA, RN, MAN, MPSA
BAC Chairperson
VALENZUELA MEDICAL CENTER
Padrigal St., Karuhatan,
Valenzuela City

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