



Republic of the Philippines
Department of Health
Metro Manila Center for Health Development
VALENZUELA MEDICAL CENTER



MINUTES OF THE PRE-BID CONFERENCE

02 May 2025

Public Bidding VMC No. 2025-055

Supply and Delivery of Ophtha Equipment – Batch 4

Present during the meeting were as follows:

BIDS & AWARDS COMMITTEE:

Ms. Ruby S. Gurrea - Chairperson
Engr. Zoraida S. Cuadra - Vice Chairperson
Dr. Maria Concepcion Isberto - BAC Member
Dr. Manuel B. Pocsidio Jr. - Provisional Member
Dr. Melquiades Figueiroa - Provisional Member

BAC SECRETARIAT:

Ms. Ligaya Ubalde - Head
Ms. Kristine Joy Manuel
Ms. Angelita Dayego
Ms. Aileen C. Pacheco
Mr. Lester John Jake R. Divino
Ms. Diana Pulido
Ms. Christallyne Castro
Ms. Kezia-Therese Medina

TWG, END-USERS & OBSERVERS:

Engr. Reynato Pascual - TWG
Ms. Esperanza Chiong - TWG
Mr. Jose Liberato Dueñas - TWG
Ms. Liza Demition - Observer, CNO
Engr. Gerardo E. Lingat - Observer, Engineer III

PROSPECTIVE BIDDER/S:

1. Mr. Romell Lingad – MTC Opto Medic Inc.
2. Ms. Mary Grace Anicete – Medilight Inc.
3. Ms. Cheska Florido – MOC OptoMedic Inc.
4. Ms. Maricel G. Pascua – MDRX Enterprise

The conference started at 10:00am and was presided by **Ms. Ruby S. Gurrea**, Chairperson of Bids & Awards Committee (BAC), held at the BAC Office, 2nd Floor, Admin Building, Valenzuela Medical Center, Padrigal St., Karuhatan, Valenzuela City. She acknowledged the presence of all representatives of each prospective bidder, the members of the BAC, BAC Secretariat, TWG as well as the invited observers. She reminded everyone that the Committee strictly adheres to Republic Act No. 12009 and its Implementing Rules and Regulations or the New Government Procurement Act.

BUSINESS MATTERS:

- In accomplishing the Technical Specifications and Schedule of Requirements, state only the item that will be bid. **Kindly include your OFFER (Technical Specs) in the "Statement of Compliance" column and state "Comply" or "Not Comply".**
- Bid Security will be forfeited if withdrawn during the validity period.
- Notice of Award will be emailed to winning bidders. The following day will be counted as 1st day of receipt.
- CTC of documents by the bidder itself are acceptable provided that the bidder will submit the Omnibus Sworn Statement. (Note: State CTC based on original, photocopy, etc.)

"PHIC Accredited Healthcare Provider"

"Valenzuela Medical Center...Where your health matters most"

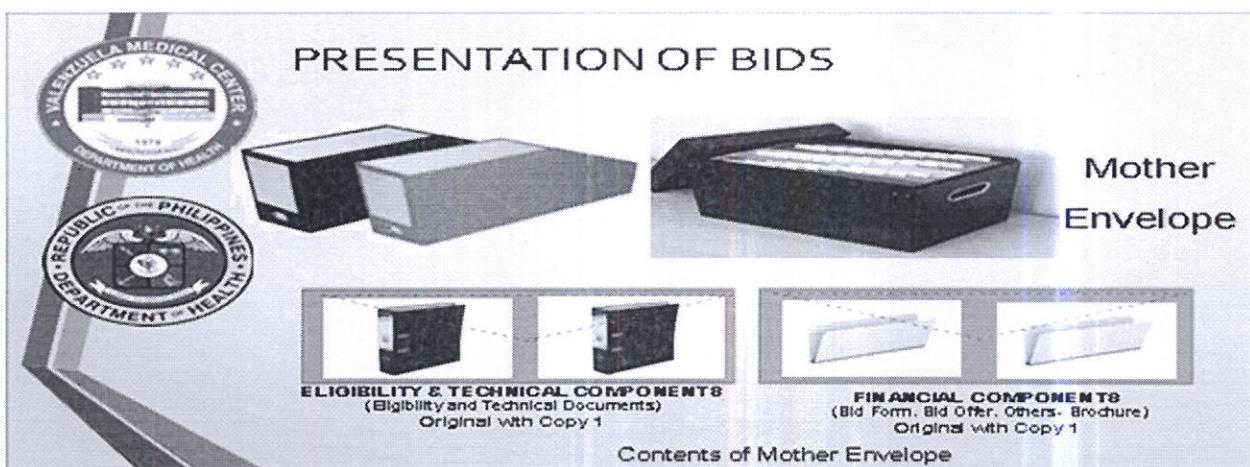


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Director's Office Direct Line: 8291-4259
Email Address: valgen_hosp@yahoo.com
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- Any document or certification issued outside Philippines should be accompanied by the official red ribbon (authentication) by the Philippine Consular Office/Embassy where the subject document or certification is issued.
- Modification of Bid is strictly prohibited. The description stated in the bid offer will be followed and cannot be amended

PRESENTATION OF BIDS:

- Bidders shall submit their bids through their duly authorized representatives using the forms specified in the Bidding Documents in two (2) separate sealed envelopes, which shall be submitted simultaneously.
- Bidders shall enclose the "Original" and "Copy 1" of their Eligibility and Technical Documents in a separate envelope marked ELIGIBILITY and TECHNICAL COMPONENTS. The "Original" and "Copy 1" of their Financial Documents (Bid Form, Bid Offer & others) shall be enclosed in a separate envelope marked FINANCIAL COMPONENTS.
- These 2 envelopes shall be enclosed in any sealed box (preferably Data File Box) with a cover.
- No color preference for the Folders and Boxes.
- **All documents to be submitted as part of the Bid should be arranged in chronological order based in the Checklist provided by the BAC. Further, all bid proposals should be ring bound and tabulated in words. Failure to follow instructions will mean disqualification.**



➤ Documents Comprising the Bid: Eligibility and Technical Components – 1st Envelope

(A) Eligibility Documents

Class "A" Documents:

(i)

- a. Valid PhilGEPS Registration Certificate (Platinum Membership) (all pages).
- b. Statement of the prospective bidder of **ALL** its ongoing Government and Private Contracts including contracts awarded but not yet started, if any, whether similar or not similar in nature and complexity to the contract to be bid;
- c. Statement of the bidder's Single Largest Completed Contract (SLCC) similar to the contract to be bid, except under conditions provided for in Section 52.4.1.3 of RA No. 12009 and its IRR, within the last ten (10) years as provided in the Bidding Documents;
 - Amount of the completed contract should be fifty (50%) of the ABC to be bid
- d. Original copy of Bid Security. If in the form of a surety Bond, submit also a certification issued by the Insurance Commission or an Original copy of the Notarized Bid Securing Declaration
- e. Conformity with the Technical Specifications, which may include production/delivery schedule, manpower requirements, and/or after-sales/parts, if applicable; and

- f. Original duly signed Omnibus Sworn Statement (OSS); and if applicable, Original Notarized Secretary's Certificate in case of a corporation, partnership, or cooperative; or Original Special Power of Attorney of all members of the joint venture giving full power and authority to its officer to sign the OSS and do acts to represent the Bidder.

Financial Documents

- g. The prospective bidder's computation of its Net Financial Contracting Capacity (NFCC); or A committed Line of Credit from a Universal or Commercial Bank in lieu of its NFCC computation.

Class "B" Documents

- h. If applicable, a duly signed joint venture agreement (JVA) in case the joint venture is already in existence or duly notarized statements from all the potential joint venture partners stating that they will enter into and abide by the provisions of the JVA in the instance that the bid is successful.

➤ FINANCIAL COMPONENT ENVELOPE – 2nd Envelope

The second envelope shall contain the financial information/documents as specified in the PBDs

- i. Original of duly signed and accomplished Financial Bid Form;
- j. Original of duly signed and accomplished Price Schedule(s).
- k. Brochure

Other documentary requirements under RA No. 12009 (as applicable)

- (l) [For foreign bidders claiming by reason of their country's extension of reciprocal rights to Filipinos] Certification from the relevant government office of their country stating that Filipinos are allowed to participate in government procurement activities for the same item or product.
- (m) Certification from the DTI if the Bidder claims preference as a Domestic Bidder or Domestic Entity.

ADDITIONAL REQUIRED DOCUMENTS (to be submitted during Post-Qualification)

1. CTC copy of Official Receipt as proof of payment of bidding documents.
2. Registration certificate from the Securities and Exchange Commission (SEC), Department of Trade and Industry (DTI) for a sole proprietorship, or Cooperative Development Authority (CDA) for cooperatives or its equivalent document.
3. Mayors or Business permits issued by the city or municipality where the principal place of business of the prospective bidder is located, or the equivalent document for Exclusive Economic Zones or Areas.
4. Tax clearance per E.O. No. 398, s. 2005, as finally reviewed and approved by the Bureau of Internal Revenue (BIR).
5. The Supplier's audited financial statements, showing, among others, the Supplier's total and current assets and liabilities, stamped "received" by the BIR with 2023 ITR or its duly accredited and authorized institutions, for the preceding calendar year which should not be earlier than two (2) years from the date of bid submission
6. Certificate of **Good Performance** from at least two (2) Government or Private Hospital/Agency except from VMC (CY 2024 to present)
7. Special Power of Attorney (SPA) for Authorized Representative if OSS is Sole proprietorship
8. Proof of evidence for Single Largest Completed Contract (SLCC) should be fifty percent (50%) of the ABC to be bid – Purchase Order or Notice of Award or Contract Agreement.
9. Certificate from the manufacturer to distribute their products or Exclusive Distributorship or any equivalent documents.
10. License to Operate (LTO)
11. Certificate of Stocks Availability from Bidder (Notarized).
12. Other requirements stated in the Bidding Documents/Technical Specifications.

REVIEW OF TECHNICAL SPECIFICATIONS:

ITEM NO.	ITEM DESCRIPTION	QTY.	UOM	UNIT PRICE	TOTAL AMOUNT	AMENDMENT
1	AUTOMATED PERIMETRY	1	unit	2,073,333.00	2,073,333.00	
	Location: Ophthalmology Clinic					
	Personnel to use the equipment: Medical Specialist					
	I. Equipment Specification:					
	Ability to add/re-test points during test					
	Duration: 200ms/ 500 ms					
	Visual field test range : 60° (monocular) / 160° (binocular) _ 10,000 ASB 31.5 ASB					
	Visual field testing distance: 25 cm					
	Stimulus intensity (maximum) 10,000 ASB					
	Background illumination 31.5 ASB					
	Manual Kinetic : Standard					
	Custom Kinetic Patterns: Standard					
	Custom Static Patterns : Standard					
	Stimulus size : I-V					
	Stimulus size : 50 um to 500 um					
	Test methods : Standard Automated Perimetry (SAP), white-on-white					
	Screening tests/patterns:					
	Single stimulus 26, 54 and 86 point tests (incorporating both 24-2 and 10-2 test pattern points)					
	Multiple stimulus 26, 54 and 86 point tests (incorporating both 24-2 and 10-2 test pattern points)					
	Groups 1 (120 point) and 2 (124 point) (EU standard)					
	Test locations can be manually added to all Smart Supra screening tests					
	Standard :Threshold central 10-2: / 30/24-2 (extendable in-test)					
	Fast :Threshold central 10-2: / 30/24-2 (extendable in-test)					
	Average test times:					
	Single stimulus ~1 min (26 points); ~3.5 minutes for fully extended 24-2 plus 10-2 test (86 points)					
	Multiple stimulus Under 30 seconds (26 points)					
	ZATA ~2.5 minutes per eye					
	ZATA Fast ~2 minutes per eye					
	Both eyes camera monitoring for binocular tests					
	Fixation target: Single or 4-point LED diamond pattern					
	Test Mode: Age corrected					
	Threshold Related					
	Stand Alone					
	II. Electrical Specification:					
	Equipment shall be compatible with hospital power supply of 220 to 240VAC, single phase, 60Hz with grounding					
	With compatible Uninterruptible Power Supply (UPS) with voltage regulation function that can provide back-up power at least 30 minutes					

	III. Technical Specification:				
	a. General Requirements				
	FDA Certificate of Medical Device Notification (CMDN)				
	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				
	Manufacturer's Certificate Brand must be in the local market for at least 5 years				
	Certification that there is established Service Center in Metro Manila or Philippines				
	Certificate for the Cost of Preventive Maintenance and Calibration Program for equipment after warranty period up to 7 years. Price validity of 7 years upon delivery of equipment				
	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				
	b. Warranty and Preventive Maintenance Services:				
	Five (5) years comprehensive warranty on parts and services for equipment				
	Three (3) years warranty on parts and services for UPS				
	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.				
	Semi-Annual Preventive Maintenance Service during warranty period				
	Annual Calibration Service during warranty period and should be conducted by manufacturers qualified Service Engineers/Technicians during warranty period				

	Calibration Report from the Manufacturer or Verification Report from the Supplier				
	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.				
	Any corrective action requiring replacement of part/component(s) shall be conducted and completed within at most 3 days, during the warranty period.				
	Service support shall cover 24 hours/day, 7 days/week during the warranty period. Remote service should be provided within 12 hours and on-site service within 24 hours must be provided during the warranty period.				
	Pricelist for major spare parts, accessories and consumables that is valid for 5 years.				
	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.				
	IV. Installation, Acceptance, Testing and Commissioning				
	Installation, Acceptance, Testing & Commissioning of Equipment				
	Certification that the bidder shall be responsible for notification, transportation, delivery, installation and commissioning at no cost to the government.				
	Comprehensive on-site operations and applications training for at least three (3) days for the end-users, to be conducted by the supplier's clinical applications specialist on the installed unit after passing the performance testing. A follow-up/refresher session after at least 3 months shall be conducted. Training should cover operation and use of hardware and software (all applications) of all equipment and their accessories and peripherals.				
	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 is not later than CY 2023 onwards				
	Verification Test during Delivery				
	Electrical Safety Test				

	Provide two (2) sets of colored Technical Manual, Hardcopy and Softcopy (Flashdrive) in English Manual for all equipment including peripherals, UPS, etc.				
	a. User's Operational Manual b. Quality and Maintenance Manual c. Service and Technical Manual				
	Comprehensive technical training for hospital biomedical engineers and unit head on maintenance and troubleshooting of all equipment and their accessories and peripherals				
	Hands-on Training for Biomedical Unit, suppliers must perform an actual:				
	a. Operations b. Disassembly and Assembly c. Troubleshooting d. Recommended Maintenance as per manufacturer				
	V. Other Terms & Condition:				
	Bidder must undertake a site inspection. A Site inspection certificate shall be issued as proof of compliance. This certificate shall be submitted with the bid.				
	Preferably packaging is made of recyclable materials				
	Preferably with green eco-products specification (non-toxic materials, energy efficient, sustainability, etc.)				
	Certification that the Bid offer is in Philippine Peso to include taxes and duties, transportation to site, delivery, installation, and testing expenses on site by the bidder				
2	<u>SLIT LAMP WITH IMAGE CAPTURE</u>	1	unit	1,705,000.00	1,705,000.00
	Location: Ophthalmology Clinic				
	Personnel to use the equipment: Medical Specialist				
	I. Equipment Specification:				
	Capability to capture Color and B/W images				
	Auto Exposure Function				
	Capability to capture rapid series of images with one shot				
	Real time live image will be seen on the monitor				
	Video Capability with LAN connection interface included				
	Slit Length (mm) 0.3~12 continuous				
	Slit Width 0.12~ continuous				
	Slit Projection 1x				
	Aperture Diaphragms 12.5x				
	Filter: Cobalt Blue, Red Free, Grey, Yellow and Heat Absorption , UV Cut Filter, ND Filter				
	Slit Rotation 0° ~ 180° continuous				
	Angle Incidence 0°, 5°, 10°, 15°, 20°				
	Patient's Eye/Prism 80mm				
	Surface Working Distance 66mm				
	Type 5 Position Rotating Drum				
	Eyepieces 12.5x				

	Total Magnifications 5x, 10x, 16x, 25x, 50x					Total Magnifications 5x-6x, 10x, 16x, 25x, 40x-50x
	Vertical Movement 28mm					
	Longitudinal Movement 78mm					
	Instrument Voltage 12V DC					
	Halogen Bulb 12V 30W or LED Lamp (10V 4.4W)					
	Fixation Point Bulb 3.4V 20mA					
	Digital Camera					
	Image Sensor 1/2" Interline CMOS					
	Image Size Up to 3,840 x 2,748 pixels					
	Cell size 1.67 µm X 1.67 µm					
	Resolution Depth 8bit or 2bit Raw RGB, yuv 4 : 2 : 2					
	With USB 2.0					
	Power Consumption less than 2.5W (5V DC, from USB Cable)					
	Accessories:					
	Motorized Stand, Heavy Duty					
	AVR					REMOVED
	Chairs (2 pieces)					
	With Applanation Tonometer, Hang Type					
	Stand Alone					
	II. Electrical Specification:					
	Equipment shall compatible with hospital power supply of 220 to 240VAC, single phase, 60Hz with grounding					
	With compatible Uninterruptible Power Supply (UPS) with voltage regulation function that can provide back-up power at least 30 minutes					
	III. Technical Specification:					
	a. General Requirements					
	FDA Certificate of Medical Device Notification (CMDN)					
	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					
	Manufacturer's Certificate Brand must be in the local market for at least 5 years					
	Certification that there is established Service Center in Metro Manila or Philippines					
	Certificate for the Cost of Preventive Maintenance and Calibration Program for equipment after warranty period up					

	to 7 years. Price validity of 7 years upon delivery of equipment				
	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				
	b. Warranty and Preventive Maintenance Services:				
	Five (5) years comprehensive warranty on parts and services for equipment				
	Three (3) years warranty on parts and services for UPS				
	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.				
	Preventive Maintenance and Calibration Services Schedule				
	Semi-Annual Preventive Maintenance Service during warranty period				
	Annual Calibration Service during warranty period and should conducted by manufacturers qualified Service Engineers/Technicians during warranty period				
	Calibration Report from the Manufacturer or Verification Report from the Supplier				
	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.				
	Any corrective action requiring replacement of part/component(s) shall be conducted and completed within at most 3 days, during the warranty period.				
	Service support shall cover 24 hours/day, 7 days/week during the warranty period. Remote service should be provided within 12 hours and on-site service within 24 hours must be provided during the warranty period.				
	Pricelist for major spare parts, accessories and consumables that is valid for 5 years.				
	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.				
	IV. Installation, Acceptance, Testing and Commissioning				
	Installation, Acceptance, Testing & Commissioning of Equipment				

	Certification that the bidder shall be responsible for notification, transportation, delivery, installation and commissioning at no cost to the government.				
	Comprehensive on-site operations and applications training for at least three (3) days for the end-users, to be conducted by the supplier's clinical applications specialist on the installed unit after passing the performance testing. A follow-up/refresher session after at least 3 months shall be conducted. Training should cover operation and use of hardware and software (all applications) of all equipment and their accessories and peripherals.				
	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 is not later than CY 2023 onwards				
	Electrical Safety Test				
	Provide two (2) sets of colored Technical Manual, Hardcopy and Softcopy (Flashdrive) in English Manual for all equipment including peripherals, UPS, etc.				
	a. User's Operational Manual b. Quality and Maintenance Manual c. Service and Technical Manual				
	Comprehensive technical training for hospital biomedical engineers and unit head on maintenance and troubleshooting of all equipment and their accessories and peripherals				
	Hands-on Training for Biomedical Unit, suppliers must perform an actual:				
	a. Operations b. Disassembly and Assembly c. Troubleshooting d. Recommended Maintenance as per manufacturer				
	V. Other Terms & Condition:				
	Bidder must undertake a site inspection. A Site inspection certificate shall be issued as proof of compliance. This certificate shall be submitted with the bid.				
	Preferably packaging is made of recyclable materials				
	Preferably with green eco-products specification (non-toxic materials, energy efficient, sustainability, etc.)				
	Certification that the Bid offer is in Philippine Peso to include taxes and duties, transportation to site, delivery, installation, and testing expenses on site by the bidder				

3	FUNDUS CAMERA WITH FLUORESCEIN ANGIOGRAPHY	1	unit	6,778,333.33	6,778,333.33	
	Location: Ophthalmology Clinic					
	Personnel to use the equipment: Medical Specialist					
	I. Equipment Specification:					
	OCT Scanning:					
	Principle : Spectral domain Optical Coherence Tomography, Confocal Scanning Laser					
	OCT Resolution : Axial resolution (in tissue) 3.9 μm (digital)					
	Lateral resolution 5.7 micron per pixel					
	30°x15° and 11x11 microns per pixel full field					
	15°x15° with 11x11 microns per pixel high speed					
	10°x10° with 5.7x5.7 microns per pixel high resolution					
	Depth Resolution 1.9 mm					
	OCT light source 820nm and 870nm					
	85,000 A scans/sec					
	Auto Alignment Dual beam active eye tracking					
	Minimum pupil diameter ≥ 2.00 mm					
	With Optical Cpherence Tonography with Angiography Image Fusion					
	With Adaptive Segmentation and OCTA Slider					
	Glaucoma Module					
	Anatomic Positioning System : creates an anatomic map of each patients eye using two fixed structural land mark (Fovea and Bruchs membrane opening)					
	Optic Nerve Head :Measurement of the Neuro-retinal rim width					
	Retinal Nerve Fiber Layer : overview of Axonal distribution along the Macular area					
	Posterior Pole: overview of the Ganglion Cell layer ang other layers of the Retina					
	Fundus Surface Imaging					
	Principle: Spectral domain OCT, Confocal Scannng Laser					
	Angle of View 30° x 30°					
	Iris Imaging: Confocal Scanning Laser Ophthalmoscopy					
	Capable of continuous capturing of images					
	Has an Auto mosaic function					
	Accessories:					
	1 unit Motorized Stand, Heavy Duty					
	Movable and adjustable motorized table, Heavy Duty					
	Stand Alone					
	II. Electrical Specification:					
	Equipment shall compatible with hospital power supply of 220 to 240VAC, single phase, 60Hz with grounding					
	With a dedicated medical equipment compatible Uninterruptible Power Supply (UPS) with voltage regulation function that can provide back-up power at least 30 minutes.					

	III. Technical Specification:				
	a. General Requirements				
	FDA Certificate of Medical Device Notification (CMDN)				
	The equipment or devices must conform to the IEC 60601 Standards and any of the following Standards:				
	<p>a. USFDA (Food and Drug administration, US Standards</p> <p>b. CE(European conformity) Standards</p> <p>c. ISO Standards (9000, 9001, 9002)</p> <p>d. UL (Underwriters Laboratories) Standards</p> <p>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</p>				
	Training certificate from manufacturers of field service engineers / technician performing Preventive and Corrective Maintenance and Calibration Services				
	Manufacturer's Certificate Brand must be in the local market for at least 5 years				
	Certification that there is established Service Center in Metro Manila or Philippines				
	Certificate for the Cost of Preventive Maintenance and Calibration Program for equipment after warranty period up to 7 years. Price validity of 7 years upon delivery of equipment				
	Certificate of Comprehensive Warranty which states the following clause				
	<p>a. The period of warranty shall be deemed to be fully comprehensive</p> <p>b. All inclusive of warranty, labor, spare parts, accessories, service consumables, manufacturer's Preventive Maintenance</p>				
	b. Warranty and Preventive Maintenance Services:				
	Five (5) years comprehensive warranty on parts and services for equipment				
	Three (3) years warranty on parts and services for UPS				
	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.				
	Preventive Maintenance and Calibration Services Schedule				
	Semi-Annual Preventive Maintenance Service during warranty period				
	Annual Calibration Service during warranty period and should be conducted by manufacturers qualified Service Engineers/Technicians during warranty period				

	Calibration Report from the Manufacturer or Verification Report from the Supplier				
	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.				
	Any corrective action requiring replacement of part/component(s) shall be conducted and completed within at most 3 days, during the warranty period.				
	Service support shall cover 24 hours/day, 7 days/week during the warranty period. Remote service should be provided within 12 hours and on-site service within 24 hours must be provided during the warranty period.				
	Pricelist for major spare parts, accessories and consumables that is valid for 5 years.				
	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.				
	IV. Installation, Acceptance, Testing and Commissioning				
	Installation, Acceptance, Testing & Commissioning of Equipment				
	Certification that the bidder shall be responsible for notification, transportation, delivery, installation and commissioning at no cost to the government.				
	Comprehensive on-site operations and applications training for at least three (3) days for the end-users, to be conducted by the supplier's clinical applications specialist on the installed unit after passing the performance testing. A follow-up/refresher session after at least 3 months shall be conducted. Training should cover operation and use of hardware and software (all applications) of all equipment and their accessories and peripherals.				
	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 is not later than CY 2023 onwards				
	Verification Test during Delivery				
	Electrical Safety Test				

	Provide two (2) sets of colored Technical Manual, Hardcopy and Softcopy (Flashdrive) in English Manual for all equipment including peripherals, UPS, etc.				
	a. User's Operational Manual b. Quality and Maintenance Manual c. Service and Technical Manual				
	Comprehensive technical training for hospital biomedical engineers and unit head on maintenance and troubleshooting of all equipment and their accessories and peripherals				
	Hands-on Training for Biomedical Unit, suppliers must perform an actual:				
	a. Operations b. Disassembly and Assembly c. Troubleshooting d. Recommended Maintenance as per manufacturer				
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	Preferably packaging is made of recyclable materials				
	Preferably with green eco-products specification (non-toxic materials, energy efficient, sustainability, etc.)				
	Certification that the Bid offer is in Philippine Peso to include taxes and duties, transportation to site, delivery, installation, and testing expenses on site by the bidder				
4	OPERATING MICROSCOPE WITH ASSIST SCOPE	1	unit	15,250,000.00	15,250,000.00
	Location: Ophthalmology Clinic				
	Personnel to use the equipment: Medical Specialist				
	I. Equipment Specification:				
	1 Unit Power cable, 5m				
	Optics with focus/tilt/assistant's scope				
	Stand F42				
	Stereo base 25 mm				Stereo base 24-25 mm
	Dust cover M600				
	Wired and Wireless Footswitch 14-functions Type B				
	6 pieces Clip-on handle				
	9 pieces Drive knob cover, binoc, tube T				9 pieces Drive knob cover, binoc, tube II
	Working distance 175 mm				
	Continuous zooming				
	Adjustable optical head: 45 to 90 degrees				
	1 unit Tube 10-50°, type II, UltraLow III				
	2 units Eyepiece f. spect.w's 8.33x/22B, type II				2 units Eyepiece f. spect.w's 8.33x/21-22B, type II
	1 unit Laser filter for IVC				
	1 unit Binocular tube inclinable 5.25° with PD				
	Filter: UV, softlight, daylight, blue				Filter: UV
	1 unit Documentation set for IVC				

	With HDR Recorder				
	27" medical HD monitor				
	With Monitor arm				
	Attachment for vitreo-retinal surgery (no inverter needed)				
	Balancing: Mechanical				
	Stand Alone				
	II. Electrical Specification:				
	Equipment shall compatible with hospital power supply of 220 to 240VAC, single phase, 60Hz with grounding				
	With compatible Uninterruptible Power Supply (UPS) with voltage regulation function that can provide back-up power at least 30 minutes				
	III. Technical Specification:				
	a. General Requirements				
	FDA Certificate of Medical Device Notification (CMDN)				
	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				
	Manufacturer's Certificate Brand must be in the local market for at least 5 years				
	Certification that there is established Service Center in Metro Manila or Philippines				
	Certificate for the Cost of Preventive Maintenance and Calibration Program for equipment after warranty period up to 7 years. Price validity of 7 years upon delivery of equipment				
	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				
	Warranty and Preventive Maintenance Services:				
	Five (5) years comprehensive warranty on parts and services for equipment				
	Three (3) years warranty on parts and services for UPS				
	Certificate of availability of spare parts within ten (10) years				Certificate of availability of spare parts within five (5) to 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.				
	Preventive Maintenance and Calibration Services Schedule				
	Semi-Annual Preventive Maintenance Service during warranty period				
	Annual Calibration Service during warranty period and should conducted by manufacturers qualified Service Engineers/Technicians during warranty period				
	Calibration Report from the Manufacturer or Verification Report from the Supplier				
	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.				
	Any corrective action requiring replacement of part/component(s) shall be conducted and completed within at most 3 days, during the warranty period.				
	Service support shall cover 24 hours/day, 7 days/week during the warranty period. Remote service should be provided within 12 hours and on-site service within 24 hours must be provided during the warranty period.				
	Pricelist for major spare parts, accessories and consumables that is valid for 5 years.				
	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.				
	IV. Installation, Acceptance, Testing and Commissioning				
	Installation, Acceptance, Testing & Commissioning of Equipment				
	Certification that the bidder shall be responsible for notification, transportation, delivery, installation and commissioning at no cost to the government.				
	Comprehensive on-site operations and applications training for at least three (3) days for the end-users, to be conducted by the supplier's clinical applications specialist on the installed unit after passing the performance testing. A follow-up/refresher session after at least 3 months shall be conducted. Training should cover operation and use of hardware and software (all applications) of all equipment and their accessories and peripherals.				
	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 is not later than CY 2023 onwards				
	Verification Test during Delivery				
	Electrical Safety Test				
	Provide two (2) sets of colored Technical Manual, Hardcopy and Softcopy (Flashdrive) in English Manual for all equipment including peripherals, UPS, etc.				
	a. User's Operational Manual b. Quality and Maintenance Manual c. Service and Technical Manual				
	Comprehensive technical training for hospital biomedical engineers and unit head on maintenance and troubleshooting of all equipment and their accessories and peripherals				
	Hands-on Training for Biomedical Unit, suppliers must perform an actual:				
	a. Operations b. Disassembly and Assembly c. Troubleshooting d. Recommended Maintenance as per manufacturer				
	V. Other Terms & Condition:				
	Bidder must undertake a site inspection. A Site inspection certificate shall be issued as proof of compliance. This certificate shall be submitted with the bid.				
	Preferably packaging is made of recyclable materials				
	Preferably with green eco-products specification (non-toxic materials, energy efficient, sustainability, etc.)				
	Certification that the Bid offer is in Philippine Peso to include taxes and duties, transportation to site, delivery, installation, and testing expenses on site by the bidder				
5	PHOROPTER HEAD WITH REFRACTIVE CHAIR	1	unit	1,367,666.67	1,367,666.67
	Location: Ophthalmology Clinic				
	Personnel to use the equipment: Medical Specialist				
	I. Equipment Specification:				
	Phoropter head				
	Sphere power range: +16.75D~ -19.00D				
	Step: 0.25D				
	Cylinder power range : OD ~ -6.00D				
	Step: 0.25D (When applied with auxiliary cylinder lens +0.12D, Step: 0.12D)				
	Cylinder axis range : 0°~180° Step: 5°				
	Cross cylinder : ± 0.25D				
	Rotary prism range : 0Δ ~20 Δ Step : 1 Δ				
	P.D. Adjustment: 50-80mm, Step:1mm				

	Convergence adjustment: 40cm (When P.D. 64mm)				
	Forehead rest adjustment: 16mm Vertex distance : 13.75mm (From corneal point to standard lens surface)				
	Dimensions : 323 (L) x 85 (W) x 315 (H) mm				REMOVED
	Weight: 5kg				REMOVED
	Refracting Unit and Chair				
	Up/Down angle of swing arm: ±30°				
	Table size: 670mm x 405mm				Table size: 670-900mm x 405-450mm
	Table rotational angle: 90°				
	Chair height adjustment range : 410~550mm				Chair height adjustment range: 410~750mm
	Loading for chair: 160kgs				Loading for chair: 160-200kgs
	Stand Alone				
	II. Electrical Specification:				
	Equipment shall compatible with hospital power supply of 220 to 240VAC, single phase, 60Hz with grounding				
	With compatible Uninterruptible Power Supply (UPS) with voltage regulation function that can provide back-up power at least 30 minutes				
	III. Technical Specification:				
	a. General Requirements				
	FDA Certificate of Medical Device Notification (CMDN)				
	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				
	Manufacturer's Certificate Brand must be in the local market for at least 5 years				
	Certification that there is established Service Center in Metro Manila or Philippines				
	Certificate for the Cost of Preventive Maintenance and Calibration Program for equipment after warranty period up to 7 years. Price validity of 7 years upon delivery of equipment				
	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				
	Warranty and Preventive Maintenance Services:				
	Five (5) years comprehensive warranty on parts and services for equipment				
	Three (3) years warranty on parts and services for UPS				
	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.				
	Preventive Maintenance and Calibration Services Schedule				
	Semi-Annual Preventive Maintenance Service during warranty period				
	Annual Calibration Service during warranty period and should conducted by manufacturers qualified Service Engineers/Technicians during warranty period				
	Calibration Report from the Manufacturer or Verification Report from the Supplier				
	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.				
	Any corrective action requiring replacement of part/component(s) shall be conducted and completed within at most 3 days, during the warranty period.				
	Service support shall cover 24 hours/day, 7 days/week during the warranty period. Remote service should be provided within 12 hours and on-site service within 24 hours must be provided during the warranty period.				
	Pricelist for major spare parts, accessories and consumables that is valid for 5 years.				
	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.				
	IV. Installation, Acceptance, Testing and Commissioning				
	Installation, Acceptance, Testing & Commissioning of Equipment				
	Certification that the bidder shall be responsible for notification, transportation, delivery, installation and commissioning at no cost to the government.				
	Comprehensive on-site operations and applications training for at least three (3) days for the end-users, to be conducted				

	by the supplier's clinical applications specialist on the installed unit after passing the performance testing. A follow-up/refresher session after at least 3 months shall be conducted. Training should cover operation and use of hardware and software (all applications) of all equipment and their accessories and peripherals.				
	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 is not later than CY 2023 onwards				
	Verification Test during Delivery				
	Electrical Safety Test				
	Provide two (2) sets of colored Technical Manual, Hardcopy and Softcopy (Flashdrive) in English Manual for all equipment including peripherals, UPS, etc.				
	a. User's Operational Manual b. Quality and Maintenance Manual c. Service and Technical Manual				
	Comprehensive technical training for hospital biomedical engineers and unit head on maintenance and troubleshooting of all equipment and their accessories and peripherals				
	Hands-on Training for Biomedical Unit, suppliers must perform an actual:				
	a. Operations b. Dissassembly and Assembly c. Troubleshooting d. Recommended Maintenance as per manufacturer				
	V. Other Terms & Condition:				
	Bidder must undertake a site inspection. A Site inspection certificate shall be issued as proof of compliance. This certificate shall be submitted with the bid.				
	Preferably packaging is made of recyclable materials				
	Preferably with green eco-products specification (non-toxic materials, energy efficient, sustainability, etc.)				
	Certification that the Bid offer is in Philippine Peso to include taxes and duties, transportation to site, delivery, installation, and testing expenses on site by the bidder				
6	POSTERIOR VITRECTOMY MACHINE	1	unit	12,750,000.00	12,750,000.00
	Location: Ophthalmology Clinic				
	Personnel to use the equipment: Medical Specialist				
	I. Equipment Specification:				

	Phacoemulsification by ultrasound with modes: Continuous, burst, pulsed				
	Dual pump aspiration system (venturi and peristaltic)				
	IOP controlled infusion for better IOP control.				
	5000 Anterior cuts/min				
	17" Articulating touch panel screen with advanced GUI				
	(Graphic User Interface) and video DFUs (Directions for Use) with remote control and high vacuum occlusion technology				
	Bipolar diathermy compatible with gauge 23, 25, 27				REMOVED
	Bipolar diathermy compatible with gauge 23, 25, 27				
	High speed vitrectomy with a cutting rate of up to 20,000 cuts/minute				
	Equipped with pneumatic and electric vitrectomy mode.				
	Auxiliary Illuminator				
	With air fluid exchange function				
	Wired single linear foot pedal				
	(Additional Features)				
	Laser Embedded 532mm Laser offers advanced laser technology controlled from the Vision System screen				
	Integrated laser firing control and power setting Cutter Gauge 25,23, 27				
	Accessories:				
	Two (2) pc Fragmetation Handpiece				
	Two (2) pcs Phaco Handpiece				
	Two 2) pcs I/A Handpiece				
	Two (2) pcs Sterilization Tray				
	Twelve (12) pcs Various Pacs				
	Stand Alone				
	II. Electrical Specification:				
	Equipment shall compatible with hospital power supply of 220 to 240VAC, single phase, 60Hz with grounding				
	With compatible Uninterruptible Power Supply (UPS) with voltage regulation function that can provide back-up power at least 30 minutes				
	III. Technical Specification:				
	a. General Requirements				
	FDA Certificate of Medical Device Notification (CMDN)				
	The equipment or devices must conform to the IEC 60601 Standards and any of the following Standards:				
	<ul style="list-style-type: none"> 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				
	Manufacturer's Certificate Brand must be in the local market for at least 5 years				
	Certification that there is established Service Center in Metro Manila or Philippines				
	Certificate for the Cost of Preventive Maintenance and Calibration Program for equipment after warranty period up to 7 years. Price validity of 7 years upon delivery of equipment				
	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				
	b. Warranty and Preventive Maintenance Services:				
	Five (5) years comprehensive warranty on parts and services for equipment				
	Three (3) years warranty on parts and services for UPS				
	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.				
	Preventive Maintenance and Calibration Services Schedule				
	Semi-Annual Preventive Maintenance Service during warranty period				
	Annual Calibration Service during warranty period and should be conducted by manufacturers qualified Service Engineers/Technicians during warranty period				
	Calibration Report from the Manufacturer or Verification Report from the Supplier				
	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.				
	Any corrective action requiring replacement of part/component(s) shall be conducted and completed within at most 3 days, during the warranty period.				
	Service support shall cover 24 hours/day, 7 days/week during the warranty period. Remote service should be provided within 12 hours and on-site service within 24 hours must be provided during the warranty period.				
	Pricelist for major spare parts, accessories and consumables that is valid for 5 years.				

	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.				
	IV. Installation, Acceptance, Testing and Commissioning				
	Installation, Acceptance, Testing & Commissioning of Equipment				
	Certification that the bidder shall be responsible for notification, transportation, delivery, installation and commissioning at no cost to the government.				
	Comprehensive on-site operations and applications training for at least three (3) days for the end-users, to be conducted by the supplier's clinical applications specialist on the installed unit after passing the performance testing. A follow-up/refresher session after at least 3 months shall be conducted. Training should cover operation and use of hardware and software (all applications) of all equipment and their accessories and peripherals.				
	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 is not later than CY 2023 onwards				
	Verification Test during Delivery				
	Electrical Safety Test				
	Provide two (2) sets of colored Technical Manual, Hardcopy and Softcopy (Flashdrive) in English Manual for all equipment including peripherals, UPS, etc.				
	a. User's Operational Manual b. Quality and Maintenance Manual c. Service and Technical Manual				
	Comprehensive technical training for hospital biomedical engineers and unit head on maintenance and troubleshooting of all equipment and their accessories and peripherals				
	Hands-on Training for Biomedical Unit, suppliers must perform an actual:				
	a. Operations b. Disassembly and Assembly c. Troubleshooting d. Recommended Maintenance as per manufacturer				
	V. Other Terms & Condition:				
	Bidder must undertake a site inspection. A Site inspection certificate shall be				REMOVED

	issued as proof of compliance. This certificate shall be submitted with the bid.				
	Preferably packaging is made of recyclable materials				
	Preferably with green eco-products specification (non-toxic materials, energy efficient, sustainability, etc.)				
	Certification that the Bid offer is in Philippine Peso to include taxes and duties, transportation to site, delivery, installation, and testing expenses on site by the bidder				
7	Gonioscopy Lens	1	unit	56,875.00	56,875.00
	Location: Ophthalmology Clinic				
	Personnel to use the equipment: Medical Specialist, Resident Doctors				
	4 x 64° mirror angles				
	1.0x image magnification				
	1.0x laser spot size				
	15 mm (flange) / 8.4 mm(no flange) contact diameter				
	One (1) year warranty on parts and services				One (1) year warranty
8	28D Lens	1	unit	31,315.00	31,315.00
	Location: Ophthalmology Clinic				
	Personnel to use the Instrument: Medical Specialist, Resident Doctors				
	53° / 69° field of view				
	2.27x image magnification				
	0.44x laser spot magnification				
	33 mm working distance				
	Available in autoclave sterilizable design or single use design				
	One (1) year warranty on parts and services				One (1) year warranty
9	78D Lens	1	unit	37,000.00	37,000.00
	Location: Ophthalmology Clinic				
	Personnel to use the instrument: Medical Specialist, Resident Doctors				
	81° / 97° field of view				
	0.93x image magnification				
	1.08x laser spot magnification				
	8 mm working distance				
	Provides great balance of magnification and field of view				
	Optimized for use within range of motion of all slit lamps				
	One (1) year warranty on parts and services				One (1) year warranty
10	90D Lens	1	unit	37,000.00	37,000.00
	Location: Ophthalmology Clinic				
	Personnel to use the equipment: Medical Specialist , Resident Doctors				
	74° / 89 ° field of view				
	0.76x image magnification				
	1.32x laser spot magnification				
	7 mm working distance				
	Small diameter ring in excellent for dynamic fundoscopy and easy manipulation within the orbit				
	One (1) year warranty on parts and services				One (1) year warranty

11	GLAUCOMA SURGICAL INSTRUMENTS	1	set	162,194.00	162,194.00	
	Location: Ophthalmology Clinic					
	Personnel to use the equipment: Medical Specialist					
	I. Instruments Specification:					
	One (1) piece Liberman Mechanical Speculum					
	One (1) piece Utility Forcep serrated					
	One (1) piece Mosquito Forcep					
	One (1) piece Castroviejo Needle Holder with or without lock					
	Two (2) pieces Barraquer Needle Holder delicate (Micro)					
	Two (2) pieces Wescott Scissor Blunt tip					
	One (1) piece Iris Scissor					
	One (1) piece Caliper straight					
	One (1) piece Colibri Forcep with tying flat form					
	One (1) piece Castroviejo colibri forcep 1x2 teeth					
	One (1) piece Kelly Punch					
	One (1) piece Mcpherson tying forcep				Two (2) pieces Mcpherson tying forcep	
	One (1) piece Mcpherson tying forcep				REMOVED	
	One (1) piece Cyclodialysis spatula					
					One (1) piece Sterilization Tray	
	Pterygium Set					
	One (1) piece Blade Holder / Scalpel Handle					
	One (1) piece Suturing Forceps 1x2 0.12mm					
	One (1) piece Adson Tissue Forcep 0.5mm. 1x2 teeth					
	One (1) piece Algerbrush Rust Ring Remover 0.5mm/ 1.0mm					
	One (1) piece Algerbrush 3.5mm					
	One (1) piece Cautery Ball					
	One (1) piece Utility Forceps					
	One (1) piece Iris Scissor					
	One (1) piece Sterilization Tray Double layer					
	IOL Delivery System					
	One (1) piece Monarch					
	One (1) year warranty on parts and services				One (1) year warranty	
				TOTAL:	40,248,717.00	

Other Concerns:

- Post-Qualification Evaluation: Technical Working Group (TWG) may ask for additional documents from supplier for validation.
- Minutes of the Pre-Bid and Bid Bulletin will be posted in the Philgeps and VMC Website (<https://vmc.doh.gov.ph/>)
- ITEMS will be subject for demo/evaluation.

A. Template in the Goods Offered in the Philippines and/or Abroad

- Column 1 – Should be in accordance with VMC's item number.
- Column 2 - Indicate the **item description of your offer** with BRAND. If no BRAND indicates **GENERIC OR NO BRAND**.
- **Column 3** – Country of Origin
- The Price Schedule should be filled completely or put zero if not applicable.

- The final unit price should be stated.
- In the Price Schedule, "*For Goods Offered from Abroad Form*" will be used if the origin of the item is from abroad, if manufactured in the Philippines, "*For Goods Offered from Within the Philippines Form*" shall be used. (Please use the attached Form/Template)

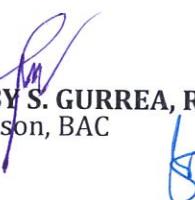
B. BID Opening will be on May 14, 2025, at 10:00 AM

The pre-bidding conference was adjourned at 2:00 pm.

Prepared by:


MS. KRISTINE JOY MANUEL
BAC Secretariat

Noted by:


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


Bid Form for the Procurement of Goods
[shall be submitted with the Bid]

BID FORM

Date : _____

Project Identification No. : _____

To: [name and address of Procuring Entity]

Having examined the Philippine Bidding Documents (PBDs) including the Supplemental or Bid Bulletin Numbers [insert numbers], the receipt of which is hereby duly acknowledged, we, the undersigned, offer to [supply/deliver/perform] [description of the Goods] in conformity with the said PBDs for the sum of [total Bid amount in words and figures] or the total calculated bid price, as evaluated and corrected for computational errors, and other bid modifications in accordance with the Price Schedules attached herewith and made part of this Bid. The total bid price includes the cost of all taxes, such as, but not limited to: [specify the applicable taxes, e.g. (i) value added tax (VAT), (ii) income tax, (iii) local taxes, and (iv) other fiscal levies and duties], which are itemized herein or in the Price Schedules,

If our Bid is accepted, we undertake:

- a. to deliver the goods in accordance with the delivery schedule specified in the Schedule of Requirements of the Philippine Bidding Documents (PBDs);
- b. to provide a performance security in the form, amounts, and within the times prescribed in the PBDs;
- c. to abide by the Bid Validity Period specified in the PBDs and it shall remain binding upon us at any time before the expiration of that period.

[Insert this paragraph if Foreign-Assisted Project with the Development Partner:]

Commissions or gratuities, if any, paid or to be paid by us to agents relating to this Bid, and to contract execution if we are awarded the contract, are listed below:

Name and address Amount and Purpose of
of agentCurrencyCommission or gratuity

(if none, state "None")]

Until a formal Contract is prepared and executed, this Bid, together with your written acceptance thereof and your Notice of Award, shall be binding upon us.

We understand that you are not bound to accept the Lowest Calculated Bid or any Bid you may receive.

We certify/confirm that we comply with the eligibility requirements pursuant to the PBDs.

The undersigned is authorized to submit the bid on behalf of [name of the bidder] as evidenced by the attached [state the written authority].

We acknowledge that failure to sign each and every page of this Bid Form, including the attached Schedule of Prices, shall be a ground for the rejection of our bid.

Name: _____

Legal capacity: _____

Signature: _____

Duly authorized to sign the Bid for and behalf of: _____

Date: _____

Bid Securing Declaration Form
[shall be submitted with the Bid if bidder opts to provide this form of bid security]

REPUBLIC OF THE PHILIPPINES)
CITY OF _____) S.S.

BID SECURING DECLARATION
Project Identification No.: *[Insert number]*

To: *[Insert name and address of the Procuring Entity]*

I/We, the undersigned, declare that:

1. I/We understand that, according to your conditions, bids must be supported by a Bid Security, which may be in the form of a Bid Securing Declaration.
2. I/We accept that: (a) I/we will be automatically disqualified from bidding for any procurement contract with any procuring entity for a period of two (2) years upon receipt of your Blacklisting Order; and, (b) I/we will pay the applicable fine provided under Section 6 of the Guidelines on the Use of Bid Securing Declaration, within fifteen (15) days from receipt of the written demand by the procuring entity for the commission of acts resulting to the enforcement of the bid securing declaration under Sections 23.1(b), 34.2, 40.1 and 69.1, except 69.1(f), of the IRR of RA No. 9184; without prejudice to other legal action the government may undertake.
3. I/We understand that this Bid Securing Declaration shall cease to be valid on the following circumstances:
 - a. Upon expiration of the bid validity period, or any extension thereof pursuant to your request;
 - b. I am/we are declared ineligible or post-disqualified upon receipt of your notice to such effect, and (i) I/we failed to timely file a request for reconsideration or (ii) I/we filed a waiver to avail of said right; and
 - c. I am/we are declared the bidder with the Lowest Calculated Responsive Bid, and I/we have furnished the performance security and signed the Contract.

IN WITNESS WHEREOF, I/We have hereunto set my/our hand/s this _____ day of *[month]* *[year]* at *[place of execution]*.

*[Insert NAME OF BIDDER OR ITS AUTHORIZED REPRESENTATIVE]
[Insert signatory's legal capacity]*

Affiant

Jurat

[Format shall be based on the latest Rules on Notarial Practice]

Omnibus Sworn Statement (Revised)
[shall be submitted with the Bid]

REPUBLIC OF THE PHILIPPINES)
CITY/MUNICIPALITY OF _____) S.S.

AFFIDAVIT

I, [Name of Affiant], of legal age, [Civil Status], [Nationality], and residing at [Address of Affiant], after having been duly sworn in accordance with law, do hereby depose and state that:

1. *[Select one, delete the other:]*

[If a sole proprietorship:] I am the sole proprietor or authorized representative of [Name of Bidder] with office address at [address of Bidder];

[If a partnership, corporation, cooperative, or joint venture:] I am the duly authorized and designated representative of [Name of Bidder] with office address at [address of Bidder];

2. *[Select one, delete the other:]*

[If a sole proprietorship:] As the owner and sole proprietor, or authorized representative of [Name of Bidder], I have full power and authority to do, execute and perform any and all acts necessary to participate, submit the bid, and to sign and execute the ensuing contract for [Name of the Project] of the [Name of the Procuring Entity], as shown in the attached duly notarized Special Power of Attorney;

[If a partnership, corporation, cooperative, or joint venture:] I am granted full power and authority to do, execute and perform any and all acts necessary to participate, submit the bid, and to sign and execute the ensuing contract for [Name of the Project] of the [Name of the Procuring Entity], as shown in the attached [state title of attached document showing proof of authorization (e.g., duly notarized Secretary's Certificate, Board/Partnership Resolution, or Special Power of Attorney, whichever is applicable);];

3. [Name of Bidder] is not "blacklisted" or barred from bidding by the Government of the Philippines or any of its agencies, offices, corporations, or Local Government Units, foreign government/foreign or international financing institution whose blacklisting rules have been recognized by the Government Procurement Policy Board, by itself or by relation, membership, association, affiliation, or controlling interest with another blacklisted person or entity as defined and provided for in the Uniform Guidelines on Blacklisting;

4. Each of the documents submitted in satisfaction of the bidding requirements is an authentic copy of the original, complete, and all statements and information provided therein are true and correct;

5. [Name of Bidder] is authorizing the Head of the Procuring Entity or its duly authorized representative(s) to verify all the documents submitted;

6. *[Select one, delete the rest:]*

[If a sole proprietorship:] The owner or sole proprietor is not related to the Head of the Procuring Entity, members of the Bids and Awards Committee (BAC), the Technical Working Group, and the BAC Secretariat, the head of the Project Management Office or the end-user unit, and the project consultants by consanguinity or affinity up to the third civil degree;

[If a partnership or cooperative:] None of the officers and members of *[Name of Bidder]* is related to the Head of the Procuring Entity, members of the Bids and Awards Committee (BAC), the Technical Working Group, and the BAC Secretariat, the head of the Project Management Office or the end-user unit, and the project consultants by consanguinity or affinity up to the third civil degree;

[If a corporation or joint venture:] None of the officers, directors, and controlling stockholders of *[Name of Bidder]* is related to the Head of the Procuring Entity, members of the Bids and Awards Committee (BAC), the Technical Working Group, and the BAC Secretariat, the head of the Project Management Office or the end-user unit, and the project consultants by consanguinity or affinity up to the third civil degree;

7. *[Name of Bidder]* complies with existing labor laws and standards; and
8. *[Name of Bidder]* is aware of and has undertaken the responsibilities as a Bidder in compliance with the Philippine Bidding Documents, which includes:
 - a. Carefully examining all of the Bidding Documents;
 - b. Acknowledging all conditions, local or otherwise, affecting the implementation of the Contract;
 - c. Making an estimate of the facilities available and needed for the contract to be bid, if any; and
 - d. Inquiring or securing Supplemental/Bid Bulletin(s) issued for the *[Name of the Project]*.
9. *[Name of Bidder]* did not give or pay directly or indirectly, any commission, amount, fee, or any form of consideration, pecuniary or otherwise, to any person or official, personnel or representative of the government in relation to any procurement project or activity.
10. **In case advance payment was made or given, failure to perform or deliver any of the obligations and undertakings in the contract shall be sufficient grounds to constitute criminal liability for Swindling (Estafa) or the commission of fraud with unfaithfulness or abuse of confidence through misappropriating or converting any payment received by a person or entity under an obligation involving the duty to deliver certain goods or services, to the prejudice of the public and the government of the Philippines pursuant to Article 315 of Act No. 3815 s. 1930, as amended, or the Revised Penal Code.**

IN WITNESS WHEREOF, I have hereunto set my hand this _____ day of _____, 20____ at _____, Philippines.

*[Insert NAME OF BIDDER OR ITS AUTHORIZED REPRESENTATIVE]
[Insert signatory's legal capacity]
Affiant*

Jurat
[Format shall be based on the latest Rules on Notarial Practice]

Price Schedule for Goods Offered from Abroad
[shall be submitted with the Bid if bidder is offering goods from Abroad]

For Goods Offered from Abroad

Name of Bidder _____ Project ID No. _____ Page ____ of ____

1	2	3	4	5	6	7	8	9
Item	Description	Country of origin	Quantity	Unit price CIF port of entry (specify port) or CIPnamed place (specify border point or place of destination)	Total CIFor CIPprice per item (col. 4 x 5)	Unit Price Delivered Duty Unpaid (DDU)	Unit price Delivered Duty Paid (DDP)	Total Price delivered DDP (col 4 x 8)
	Brand							

Name: _____

Legal Capacity: _____

Signature: _____

Duly authorized to sign the Bid for and behalf of: _____

Price Schedule for Goods Offered from Within the Philippines
[shall be submitted with the Bid if bidder is offering goods from within the Philippines]

For Goods Offered from Within the Philippines

Name of Bidder _____ Project ID No. _____ Page ____ of ____

1	2	3	4	5	6	7	8	9	10
Item	Description	Country of origin	Quantity	Unit price EXW per item	Transportation and all other costs incidental to delivery, per item	Sales and other taxes payable if Contract is awarded, per item	Cost of Incidental Services, if applicable, per item	Total Price, per unit (col 5+6+7+8)	Total Price delivered Final Destination (col 9) x (col 4)
	Brand								

Name: _____

Legal Capacity: _____

Signature: _____

Duly authorized to sign the Bid for and behalf of: _____

NFCC COMPUTATION FOR ELIGIBILITY CHECK

- A. Summary of the Applicant Supplier's/Distributor's/Manufacturer's assets and liabilities on the basis of the attached income tax return and audited financial statement, stamped "RECEIVED" by the Bureau of Internal Revenue or BIR authorized collecting agent, for the immediately preceding year and a certified copy of Schedule of Fixed Assets particularly the list of construction equipment.

	Year 20_____
1. Total Assets	
2. Current Assets	
3. Total Liabilities	
4. Current Liabilities	
5. Net Worth(1-3)	
6. Net Working Capital(2-4)	

- B. The Net Financial Contracting Capacity (NFCC) based on the above data is computed as follows:

NFCC= [(Current assets minus current liabilities) (15)] minus the value of all outstanding or uncompleted portions of the projects under ongoing contracts, including awarded contracts yet to be started, coinciding with the contract to be bid.

The values of the domestic bidder's current assets and current liabilities shall be based on the latest Audited Financial Statements (AFS) submitted to the BIR.

The Bidder shall attach the AFS to the NFCC Computation for Eligibility Check Form.

NFCC=P_____

Submitted by:

Name of Supplier/Distributor/Manufacturer

Signature of Authorized Representative

Date: _____

STATEMENT OF SINGLE LARGEST COMPLETED CONTRACT SIMILAR TO THE CONTRACT TO BE BID

This is to certify that _____ (company) _____ has the following completed contracts within Ten (10) years from the date of submission and receipt of bids.

Date of the Contract	Contracting Party	Name of Contract	Kind of Goods Sold	Amount of Contract	Date of Delivery/ End-user's Acceptance	Date of Official Receipt	Bidder is A) Manufacturer B) Supplier C) Distributor

Name and Signature of
Authorized Representative

Date

***Instructions:**

- a) Cut-off date as of:
 - (i) Up to the day before the deadline of submission of bids.
- b) In the column under "Dates", indicate the dates of Delivery/ End-user's Acceptance and Official Receipt.
- c) "Name of Contract". Indicate here the Nature/ Scope of the Contract for the Procuring Entity to determine the relevance of the entry with the Procurement at hand. Example: "Supply and Delivery of _____ for Valenzuela Medical Center"

STATEMENT OF: (I) ONGOING CONTRACTS AND; (II) AWARDED BUT NOT YET STARTED CONTRACTS

This is to certify that _____ has the following ongoing and awarded but not yet started contracts:

Date of the Contract	Contracting Party	Name of Contract	Kind of Goods Sold	Amount of Contract	Value of Outstanding Contracts	Bidder is A) Manufacturer B) Supplier C) Distributor

Name and Signature of
Authorized Representative

Date

***Instructions:**

- a) State all ongoing contracts including those awarded but not yet started (government and private contracts which may be similar or not similar to the project called for bidding) as of:
 - i. The day before the deadline of submission of bids.
- b) If there is no ongoing contract including awarded but not yet started as of the aforementioned period, state none or equivalent term.
- c) The total amount of the ongoing and awarded but not yet started contracts should be consistent with those used in the Net Financial Contracting Capacity (NFCC) in case an NFCC is submitted as an eligibility document.
- d) "Name of Contract". Indicate here the Nature/ Scope of the Contract for easier tracking of the entries/ representations. Example: "Supply and Delivery of _____ for Valenzuela Medical Center"