



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



MINUTES OF THE PRE-BID CONFERENCE

20 August 2025 @ 1:30pm

Public Bidding VMC No. 2025-070

Supply and Delivery of Various Medical Equipment - Batch 6

Present during the meeting were as follows:

BIDS & AWARDS COMMITTEE:

Engr. Zoraida S. Cuadra – Vice-Chairperson
Dr. Maria Concepcion Isberto – BAC Member
Mr. Roland Saoi – BAC Member
Dr. Jed Patrick Cruz – Provisional Member

BAC SECRETARIAT:

Ms. Ligaya Ubalde – Head
Ms. Kristine Joy Manuel
Ms. Angelita Dayego
Ms. Aileen C. Pacheco
Ms. Diana Pulido
Ms. Christallyne Castro
Ms. Kezia-Therese Medina

TWG, END-USERS & OBSERVERS:

Engr. Reynato Pascual – TWG (Clinical)
Ms. Avigail Ching – TWG (Clinical)
Ms. Maria Fatima Pastidio – TWG (Nursing)
Ms. Chezca Marie Gerodias – TWG (Nursing)
Mr. Dennis Santillan – TWG (Nursing)
Mr. Roderick Balagta – Observer, Procurement Section

Dr. Joyce Manahan – End-user, OB-Gyne
Mr. Eric Pajarillo – End-user, ED
Ms. Berna Marga Velasquez – End-user,
CSSU

PROSPECTIVE BIDDER/S:

1. Mr. Alexis Malicsi – NPK Medical Trading Inc.
2. Mr. Rolando Gonzalez Jr. – BioMerjs Trading Corp.
3. Ms. Antonette Mandate – Medical Gallery Trading Co.
4. Ms. Divine Grace Delgado – Carewell Biomedical Systems Co.
5. Mr. Rechie Osis – RP Medical
6. Mr. Dennis Alfonso – Josmef Enterprises

The conference started at 1:30pm and was presided by **Engr. Zoraida S. Cuadra**, Vice-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- Bid Security will be forfeited if withdrawn during the validity period.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"

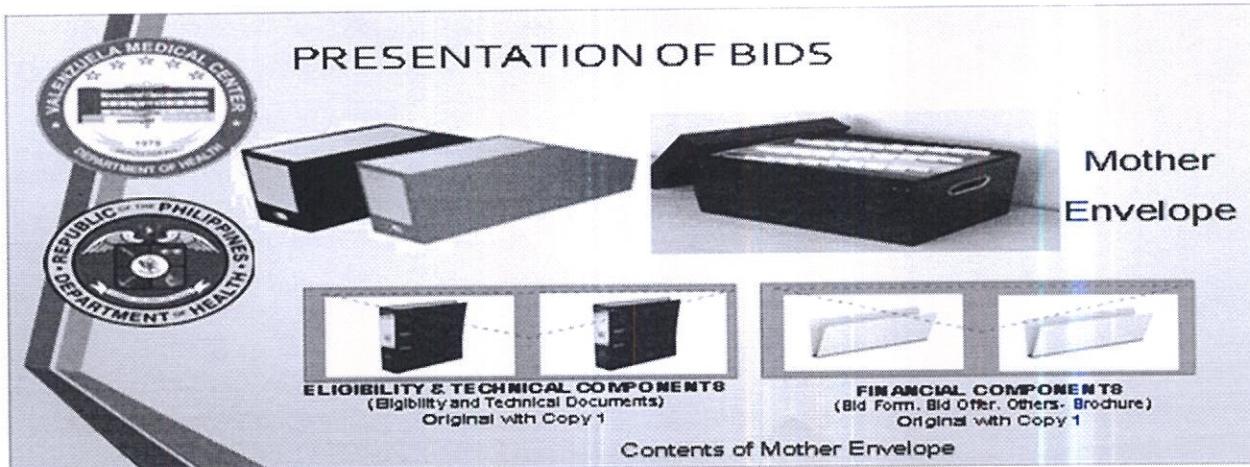


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Email Address: valgen_hosp@yahoo.com
Website: <http://vmc.doh.gov.ph>

- 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.
- 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 **2024 ITR** or its duly accredited and authorized institutions, for online submission, an email confirmation from BIR 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** – 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) from FDA
11. 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	FETAL MONITOR WITH BATTERY TWIN ANTEPARTUM	3	unit	696,000.00	2,088,000.00	
	Features:					
	Dual channel ultrasound fetal heart rate detection with audio					
	External monitoring of maternal contractions					
	Maternally sensed fetal movement					
	Colour 8.4" touchscreen display					
	Connections to Central Review System via serial port or ethernet					
	Connection to wireless transducer system, ultrasound and toco					
	USB-for upgrading and configuration					
	CARE Analysis -Dawes-Redman (100,000 normal records data base)					
	Wide beam technology ultrasound (locate and track algorithm)					
	Out of paper store capability up to 100 hours					
	Powerful internal memory (16GB)					
	Integral rechargeable battery					
	With two (2) ultrasound transducer (standard)					
	DISPLAY					
	Technology: full colour TFT Liquid Crystal display					
	Size: 8.4" diagonal 4:3 aspect ration					
	Resolution: SVGA, 800 x 600					
	Viewing Angle: Better Than 170 degree					
	FETAL PARAMETERS ULTRASOUND					
	Range: 30 to 240bpm					
	Accuracy: + 1 bpm over the range 100-180 bpm + 2 bpm outside range					
	Alarms: High and Low FHR: 30 to 240 bpm Signal loss: % loss in last 5 minutes					
	Mode: Directional pulsed doppler Repetition rate: 3.0 kHz					
	Display: FHR values Pulse indicator Confidence indicator Line graph					
	Print: Line graph					
	Repetition rate: 2.994 kHz					
	Frequency: 1.0 MHz (green)					
	Pressure: <30kPa					
	Iob: <1mW/cm ²					
	Ispta: <3mW/cm ²					
	Resolution: 12 bits					
	Safety: Type CF protection					
	Ingress Protection: IPX7 rating					
	Standards: IEC60601-2-37:2007					
	EXTERNAL UTERINE ACTIVITY (TOCO)					
	Range: 0-100 relative units					

	Sensitivity: 80% (+5%) scale reading equivalent to 100g 100% FSD equivalent to 125 g				
	Offset range: + 100g				
	Baseline: Manual and auto zero facility to 0, 10 or 20%				
	Display: TOCO values Line graph				
	Print: Line graph				
	Safety: Type CF protection				
	Ingress Protection: IPX7				
	ALARMS & ALERTS				
	Audible and visual notification is provided for all fetal alarms. Alarms limits are all user adjustable *High heart rate *Low heart rate *Signal loss *Dual rate detection				
	FETAL MOVEMENT				
	Recorded with either the maternally sensed marker, or automatically using Actogram. This records the fetal limb and trunk movements by detecting low frequency Doppler signals through the 1.0 MHz ultrasound transducer				
	PRINTER				
	Print head: 128 mm thick film				
	Resolution: 8 dots per mm				
	Printer speeds: 1,2, or 3cm per minute (user selectable)				
	Fast forward: 10cm/minute				
	FHR scales: 30-240 bpm or 50-210 bpm (user selectable)				
	Annotation: Hospital name, time, date, paper speed, monitoring modes, signal loss				
	High speed: Review and print catch-up at up to 20cm/minute				High speed: Review and print catch-up at up to 10-20cm/minute
	PAPER				
	Plain paper: Thermal paper, z-fold 45m length (75 hours @ 1cm/minute)				
	Pre-printed paper: HP/Philips/Agilent Corometrics Spacelabs				
	SCALES & GRIDS				
	FHR:30-240 bpm 50-210 bpm				
	TOCO: 0-100% 0-13.3 kPa				
	OUT OF PAPERSTORE				
	A single recorded session has a maximum time limit of 100 hours				
	CONNECTORS				
	Location: ER, WARD, OPD, LR				
	Front panel Ultrasound:1.0 MHz ultrasound transducer (2 connectors) TOCO: Strain gauge toco-dynamometer				
	Rear panel: *IEC-320 C14-mains power * Fetal Event Marker socket: 1/4 inch (6.35mm) jack plug connection				

	<ul style="list-style-type: none"> *Equipotential Earth Point: provides coomon eathing point for connected equipment *RS232: CRS interface via 9 way D-Type connector Auziliary: for wireless telemetry system USB Ports: External keyboard, barcode reader, upgrader memory stick Ethernet Port: Future CRS 				
	<ul style="list-style-type: none"> Power *Supply Voltage:100-240V auto sensing *Fuse Type: 2x T3.15AH 250V *Power Input: 60Hz *Consumption:8-133 VA 				
	<ul style="list-style-type: none"> Battery *Capacity: 4400mAh *Type: Lithium Ion *Use: Up to 4 hours depending on operating mode *Charging : approx. 4 hours 				
	<ul style="list-style-type: none"> Physical Height:23.4cm(9.2 in) Width: 32.0 com (12.6) Length: 23.0 cm (9.0) Weight: 6 kg (13.2 lbs) 				
	ENVIRONMENTAL				
	<ul style="list-style-type: none"> Operating * Temperature range: +10 C to +40 C * Relative Humidity: 10% to 90% (non condensing) *Pressure: 860mb to 1060 mb 				
	<ul style="list-style-type: none"> STORAGE * Temperature range: -10 C to +40 C * Relative Humidity:93 % maximum *Pressure: 860mb to 1060mb 				
	EQUIPMENT CLASSIFICATION <ul style="list-style-type: none"> *Type of Protection against electric shock: Class1 * Mode of operation: Continous * Degree of protection against harmful ingress of particles/or water: IP30 Degree of safety of application in the presence of a flammable anaesthetic: Equipment not suitable for use in the presence of a flammable Anaesthetic mixture *with air, oxygen or nitrous oxide 				
	Warranty: Two (2) years warranty on parts and service under normal use and condition except on expendable parts.				
	II. Electrical Specification:				
	Can be mains operated and battery				
	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				

	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				
	The equipment or devices must conform to 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:				
	Two (2) years 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.				
	Preventive Maintenance and Calibration Services Schedule				
	Semi-Annual Preventive Maintenance and Annual Calibration Service during warranty period				
	Certification that the bidder will provide a Loancer/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.				
	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				
	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	PEDIATRIC CRIB	17	unit	42,245.67	718,176.39
	I. Equipment Specification:				
	Bed frame: made of stainless steel				
	CRIB DIMENSION:				
	LENGTH: 130 cm				
	WIDTH: 75 cm				
	HEIGHT: not exceeding 150cm from floor to top, not exceeding 55 cm from floor to base.				
	Headrests: Stainless Steel				Headrests: Stainless Steel (Adjustable)
	Footrests: Stainless Steel				
	Side Railings: Stainless Steel				
	Railing Height: min. 500mm				
	Loading Capacity: min. 120 kgs				
	Stainless steel, adjustable IV pole, 4 hooks				
	With nameplate holder				
	With urine bag hooks				
	4" Heavy Duty Caster wheels, double ball bearing (2 fixed, 2 with locking break)				4-6" Heavy Duty Caster wheels, double ball bearing (2 fixed, 2-4 with locking break)
	4" thick high density and heavy duty foam mattress with leatherette cover, fire retardant				
	Function:				
	-Back panel can be lifted up by operating the handle at foot end				
	-Side rails, one side can be lowered with leveler to adjust height				
	III. Technical Specification:				
	*Certificates that need to be presented during Post-Qualification				
	a. General Requirements				
	The equipment or devices must conform to 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				

	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				
	*Certificates to be submitted upon Delivery				
	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				
	b. Warranty and Preventive Maintenance Services:				
	One (1) year comprehensive warranty on parts and services for equipment				
	Certificate of availability of spare parts within five (5) years				
	IV. Installation, Acceptance, Testing and Commissioning				
	*Procedures to be performed upon Delivery				
	Acceptance, Testing & Commissioning of Equipment				
	The unit to be delivered must show proof that its manufacturing date is not later than CY 2024 onwards				
	Provide two (2) sets of colored Technical Manual, Hardcopy and Softcopy (Flashdrive) in English Manual				
	a. User's Operational Manual b. Quality and Maintenance Manual (if applicable) c. Service and Technical Manual (if applicable)				
	V. Other Terms & Condition:				
	Preferably with green eco-products specification (made of recyclable and non-toxic materials, sustainability, etc.)				
3	PATIENT TRANSPORT STRETCHER	6	unit	136,000.00	816,000.00
	I. Equipment Specification:				
	Bed Frame: Carbon Steel with electrostatic powder coat finish and oxygen cylinder holder				Size: W-80cm - 90cm (including bed rails) x L-210cm
	Mounted, heavy duty 5-6" diameter double caster wheels, controlled by pedal central lock and with retractable center fifth wheel				
	With X-Ray Cassette holder and with X-Ray translucent platform				
	Capacity: At least 200kg.				
	4" thick high density and heavy duty foam mattress with leatherette cover, fire retardant				
	IV pole with two (2) pcs sockets on corners with height adjustable IV pole with two (2) to four (4) hooks				Steel IV pole with two (2) pcs sockets on corners with height adjustable IV pole with two (2) to four (4) plastic or steel hooks

	Adjustable by manual steel crank handle or hydraulic				
	Stainless steel Tuck-away side rails				
	Accessories: removable restraint strap and patient transfer mat, with removable 2-inch thick mattress with leatherette cover (Black or Blue)				
	II. Technical Specification:				
	a. General Requirements				
	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				
	The equipment or devices must conform to 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				
	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 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				
	b. Warranty:				
	Comprehensive Warranty: 2 years				
	Certification of Uptime of 95% and Downtime of 5%				
	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.				
	III. Acceptance, Testing and Commissioning				
	Acceptance, Testing & Commissioning of Equipment				
	Training of End-User (Operation) and Biomedical Unit (Operation and Troubleshooting) 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.					
	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					
	IV. Other Terms & Condition:					
	Preferably with green eco-products specification (non-toxic materials, energy efficient, sustainable, etc.)					
4	PATIENT TRANSPORT STRETCHER	10	unit	136,000.00	1,360,000.00	
	I. Equipment Specification:					
	Bed Frame: Carbon Steel with electrostatic powder coat finish and oxygen cylinder holder					Size: W-80cm - 90cm (including bed rails) x L-210cm
	Mounted, heavy duty 5-6" diameter double caster wheels, controlled by pedal central lock and with retractable center fifth wheel					
	With X-Ray Cassette holder and with X-Ray translucent platform					
	Capacity: At least 200kg.					
	4" thick high density and heavy duty foam mattress with leatherette cover, fire retardant					
	IV pole with two (2) pcs sockets on corners with height adjustable IV pole with two (2) to four (4) hooks					Steel IV pole with two (2) pcs sockets on corners with height adjustable IV pole with two (2) to four (4) plastic or steel hooks
	Adjustable by manual steel crank handle or hydraulic					
	Stainless steel Tuck-away side rails					
	Accessories: removable restraint strap and patient transfer mat, with removable 2-inch thick mattress with leatherette cover (Black or Blue)					
	II. Technical Specification:					
	a. General Requirements					
	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					
	The equipment or devices must conform to 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					
	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 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					
	b. Warranty: Comprehensive Warranty: 2 years Certification of Uptime of 95% and Downtime of 5%					
	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					
	III. Acceptance, Testing and Commissioning					
	Acceptance, Testing & Commissioning of Equipment					
	Training of End-User (Operation) and Biomedical Unit (Operation and Troubleshooting) 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.					
	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					
	IV. Other Terms & Condition:					
	Preferably with green eco-product specification (packaging is made of recyclable materials, made with non-toxic materials, energy efficient, sustainable, etc.)					
5	AUTOCLAVE, HORIZONTAL	1	unit	1,992,340.20	1,992,340.20	
	I. Equipment Specification:					
	Capacity: 200L					
	Door: Manual					

	Display: 7-inch LCD touch screen				
	Designed Pressure: -0.1/0.3MPa				
	Working Pressure: 0.23MPa				
	Max Designated Temp.: 200°C				
	Working Temp.: 105-134°C				
	Temp. Precision: 0.1°C				
	Sterilization Time: 0-99min				
	Drying Time: 0-99min				
	Vacuum System: Water circulating vacuum pump				
	Chamber Material: Stainless Steel S30408				
	Rack: 1 mesh frame with 1 cover				
	Accessories: Manufacturer's Standard including stainless steel frame with cover, mobile steel frame with cover, mobile sensor, printer				
	With one (1) free Biological indicator unit				
	Features:				
	-LCD Display Information: Temperature, Pressure,				
	Operating Status, Fault Alarm, Cause Analysis, and Solutions				
	-Air in the chamber is sterile and avoids re-contamination				
	-With quick-release side and top covers				
	-Micro-controller programmable control technology				
	-Full protective door cover				
	-With self-expanding sealing ring				
	-With built-in high speed steam generator				
	-With pressure, mechanical and electronic safety interlock device, over-temperature over-pressure protection				
	II. Electrical Specification:				
	Equipment shall compatible with hospital power supply of 220 to 240VAC, single phase, 60Hz with grounding				
	III. Technical Specification:				
	*Certificates that need to be presented during Post-Qualification				
	a. General Requirements				
	The equipment or devices must conform to the IEC 60601 Standards or 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 5 years. Price validity of 5 years upon delivery of equipment				
	*Certificates to be submitted upon Delivery				
	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:				
	Two (2) years 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 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				
	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 (if applicable)				
	IV. Installation, Acceptance, Testing and Commissioning				
	*Procedures to be performed upon Delivery				
	Supply, Delivery, Installation, Acceptance, Testing & Commissioning of Equipment				

	The supplier shall shoulder all installation including but not limited to architectural, civil, electrical, electronics, mechanical, plumbing, sanitary and other appropriate modifications to meet the manufacturer's requirements and to comply with relevant requirements				
	All electrical wiring/cabling, and electrical devices from the electrical room to the equipment room shall be supplied, installed and tested by the supplier.				
	Complete plumbing works including Water System (Reverse Osmosis Machine) dedicated to equipment shall be supplied, installed and tested by the supplier				
	All design and materials to be used shall be approved by the Engineering and Facilities Management Section (EFMS) head prior to installation				
	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 2024 onwards				
	Conduct Verification Test during Delivery				
	Conduct Electrical Safety Test				
	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				
	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 with green eco-products specification (made of recyclable and non-toxic materials, energy efficient, sustainability, etc.)				
			TOTAL:	6,974,516.59	

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 as part of Post Qualification Evaluation.
- Bidders are advised to use two (2) decimal places in setting up their bid prices.
- New forms are already available and downloadable thru GPPB Website but old forms are still acceptable within the transition period of the NGPA (RA 9184 to RA 12009).

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/Source of Domestic Product, as certified by the Relevant Agency
- 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 September 2, 2025, at 10:00 AM

The pre-bidding conference was adjourned at 4:00 pm.

Prepared by:


MS. KRISTINE JOY MANUEL
BAC Secretariat

Noted by:


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

FORMS

Bid Form for Procurement of Goods

[Note: The duly accomplished form shall be submitted with the Bid]

BID FORM

Project Identification No.: [Insert number]

To: [Name of Procuring Entity]

Having examined the Philippine Bidding Documents (PBD) including the Supplemental Bid Bulletin Numbers [insert numbers], the receipt of which is hereby duly acknowledged, we, the undersigned, declare that:

- a) I/We have no reservation to the PBD, including the Supplemental Bid Bulletins, for the Procurement Project **[Project Title]**;
- b) Select one, delete the other
 - I/We undertake to deliver the Goods in accordance with the delivery schedule in the Schedule of Requirements;
 - I/We offer to execute the Works for this Contract in accordance with the PBD;
- c) The total price of our Bid in words and figures, excluding any discount offered below, is **[insert information]**;
- d) The discounts offered and the methodology for their application are: **[insert information]**;
- e) 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 **[Select one, delete the other]:** the Price Schedules/ Detailed Estimates];
- f) This Bid shall remain valid within a period stated in the PBD, and it shall be binding upon me/us at any time before the expiration of that period;
- g) If our bid is accepted, I/We commit to provide a performance security in the form, amounts, and within the times prescribed in the PBD.

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 the Bidder.

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

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

The undersigned is authorized to submit the bid on behalf of **[Name of the Bidder]** as evidenced by the attached **[State the Written Authority]**.

I/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.

Duly authorized to sign the Bid for and behalf of:

[Insert Bidder's Name]

[Signature over Printed Name]

[Position/Designation]

[Date]

Price Schedule for Goods

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

Pricing Details for Goods Offered from Within the Philippines

1	2	3	4	5	6	7	8	9	10
Item	Description	Source of Domestic Product, as certified by the Relevant Agency	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)

Summary of Bid Prices

The Procuring Entity may modify the table below as necessary to comply with the requirements of the Procurement Project.

1	2	3	4
Item No.	Item	Particulars / Description	Total Amount

Name: _____

Signature: _____

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

Price Schedule for Goods

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

Pricing Details for Goods Offered from Abroad

1	2	3	4	5	6	7	8	9
Item	Description	Country of origin	Quantity	Unit price CIF port of entry (specify port) or CIP named place (specify border point or place of destination)	Total CIF or CIP price 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)

Summary of Bid Prices

The Procuring Entity may modify the table below as necessary to comply with the requirements of a specific Project.

1	2	3	4
Item No.	Item	Particulars / Description	Total Amount

Name: _____

Signature: _____

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

Omnibus Sworn Statement Form

[Note: The duly accomplished form shall be submitted with the Bid]

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

OMNIBUS SWORN STATEMENT

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

1) Select one, delete the others:

- If sole proprietorship: I am the sole proprietor or authorized representative of [Name of Bidder] with office address at [Address of Bidder];
- If 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];
- If individual consultant not registered under a sole proprietorship, in case of Consulting Services: I am the individual consultant or authorized representative of [Name of Bidder] with office address at [Address of Bidder];

2) Select one, delete the others:

- If 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 [Project Title] of the [Name of the Procuring Entity][insert "as supported by the attached duly notarized Special Power of Attorney" for authorized representative];
- If 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 [Project Title] of the [Name of the Procuring Entity], as supported by the attached duly notarized Special Power of Attorney, Board/Partnership Resolution, or Secretary's Certificate, whichever is applicable;
- If individual consultant not registered under a sole proprietorship, in case of Consulting Services: As the individual consultant 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 [Project Title] of the [Name of the Procuring Entity], as supported by the attached duly notarized Special Power of Attorney for authorized representative;

- 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;
- 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 others:**

- *If sole proprietorship* : The **[Name of Bidder]** and its spouse are not related by consanguinity or affinity up to the third civil degree to the Head of the Procuring Entity, Procurement Agent (if engaged), End-User or Implementing Unit, project consultants, head of the Project Management Office, or the members of the Bids and Awards Committee (BAC), the Technical Working Group, and the BAC Secretariat;
- *If partnership* : The partnership itself and the partners of **[Name of Bidder]** are not related by consanguinity or affinity up to the third civil degree to the Head of the Procuring Entity, Procurement Agent (if engaged), End-User or Implementing Unit, project consultants, head of the Project Management Office, or the members of the Bids and Awards Committee (BAC), the Technical Working Group, and the BAC Secretariat;
- *If cooperative*: The cooperative itself and members of the board of directors, general manager, or chief executive officer of **[Name of Bidder]** are not related by consanguinity or affinity up to the third civil degree to the Head of the Procuring Entity, Procurement Agent (if engaged), End-User or Implementing Unit, project consultants, head of the Project Management Office, or the members of the Bids and Awards Committee (BAC), the Technical Working Group, and the BAC Secretariat;
- *If corporation, or joint venture*: The corporation or joint venture itself, and officers, directors, and controlling stockholders of **[Name of Bidder]** are not related by consanguinity or affinity up to the third civil degree to the Head of the Procuring Entity, Procurement Agent (if engaged), End-User or Implementing Unit, project consultants, head of the Project Management Office, or the members of the Bids and Awards Committee (BAC), the Technical Working Group, and the BAC Secretariat;
- *If individual consultant not registered under a sole proprietorship, in case of Consulting Services*: The individual consultant and its spouse are not related by consanguinity or affinity up to the third civil degree to the Head of the Procuring Entity, Procurement Agent (if engaged), End-User or Implementing Unit, project consultants, head of the Project Management Office, or the members of the Bids and Awards Committee (BAC), the Technical Working Group, and the BAC Secretariat;

- 7) It is understood that failure to faithfully disclose its relationship with the HoPE, members of the BAC, the TWG, and the BAC Secretariat, the head of the PMO or the end-user unit or implementing unit, and the project consultants of the Procuring Entity, or of the procurement agent by consanguinity or affinity up to the third civil degree, as well as its submission of beneficial ownership information containing false entries shall be subject to blacklisting under Section 100 of the IRR of RA No. 12009, without prejudice to criminal and civil liabilities under applicable laws, including their accessory penalties, if any.

[Select one, delete the rest:]

- *In case of corporations*: **[Name of Bidder]** declares its beneficial ownership consistent with its updated General Information Sheet or Beneficial Ownership Declaration Form or any other document duly submitted to the SEC in accordance with its annual reportorial requirements.
- *In case of Foreign Bidders*: **[Name of Bidder]** submitted an appropriate equivalent document in English issued by the country of the bidder concerned in accordance with Section 20.2.9.2 of the IRR of RA No. 12009.

- 8) **[Name of Bidder]** complies with existing labor laws and standards; and

9) **[Name of Bidder]** is aware of and has undertaken the following responsibilities as a Bidder:

- a) Carefully examine all of the Bidding Documents;
- b) Acknowledge all conditions, local or otherwise, affecting the implementation of the Contract;
- c) Made an estimate of the facilities available and needed for the contract to be bid, if any; and
- d) Inquire or secure Supplemental Bid Bulletin(s) issued for the **[Project Title]**.

10) **[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.

11) In case advance payment was made or given to **[Name of Bidder]**, failure to perform or deliver any of the obligations and undertakings in the contract shall be sufficient grounds to constitute criminal liability under existing laws.

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

Duly authorized to sign the Bid for and behalf of:

[Insert Bidder's Name]

[Affiant's Signature over Printed Name]

[Position/Designation]

[Date]

JURAT

SUBSCRIBED AND SWORN to before me this ____ day of **[month] [year]** at **[place of execution]**, Philippines. Affiant/s is/are personally known to me and was/were identified by me through competent evidence of identity as defined in the 2004 Rules on Notarial Practice (A.M. No. 02-8-13-SC). Affiant/s exhibited to me his/her **[insert type of government identification card used]**, with his/her photograph and signature appearing thereon, with no. _____.
WITNESS MY HAND AND SEAL this ____ day of **[month] [year]**.

NAME OF NOTARY PUBLIC

Notarial Commission No. _____

Notary Public for _____ until _____

Roll of Attorneys No. _____

PTR No. ___, [date issued], [place issued]

IBP No. ___, [date issued], [place issued]

Doc. No. _____

Page No. _____

Book No. _____

Series of _____.

Bid Securing Declaration Form

[The duly accomplished form shall be submitted with the Bid if bidder opts to provide this type of bid security]

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

BID SECURING DECLARATION Project Identification No.: [Number]

To: *[Insert name 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;

[Insert paragraph for Unsolicited Offer with Bid Matching]

I/We understand that upon conferment of the original offeror status under Section 30.6 of the Implementing Rules and Regulations (IRR) of Republic Act (RA) No. 12009, the offeror shall submit a Bid Securing Declaration within ten (10) days from the receipt of the certificate of conferment;

- 2) **Select one, delete the other:**

- I/We accept that: (a) I/we will be automatically disqualified from bidding for any procurement contract with any Procuring Entity upon receipt of your Blacklisting Order; and, (b) I/we will pay the applicable fine provided under 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 52.2 (a), 63.2, 69.1 and 100, except 100.3 (c), of the IRR of Republic Act No. 12009; without prejudice to other legal action the government may undertake; and

(For Unsolicited Offer with Bid Matching)

- I/We accept that: I/we will be automatically disqualified from any procurement opportunity of the Procuring Entity for a period of one (1) year on the first offense, two (2) years on the second offense, and perpetually on the third offense 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:

Upon expiration of the bid validity period, or any extension thereof pursuant to your request;

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;

[Insert this paragraph for Unsolicited Offer with Bid Matching]

Upon contract award and the LCCRB is not the original offeror; or

I am/we are declared the bidder with the *[Insert Award Criterion]* 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].

Duly authorized to sign the Bid for and behalf of:

[Insert Bidder's Name]

[Signature over Printed Name]

[Position/Designation]

[Date]

JURAT

SUBSCRIBED AND SWORN to before me this _____ day of [month] [year] at [place of execution], Philippines. Affiant/s is/are personally known to me and was/were identified by me through competent evidence of identity as defined in the 2004 Rules on Notarial Practice (A.M. No. 02-8-13-SC). Affiant/s exhibited to me his/her [insert type of government identification card used], with his/her photograph and signature appearing thereon, with no. _____.

WITNESS MY HAND AND SEAL this _____ day of [month] [year].

NAME OF NOTARY PUBLIC

Notarial Commission No. _____

Notary Public for _____ until _____

Roll of Attorneys No. _____

PTR No. ___, [date issued], [place issued]

IBP No. ___, [date issued], [place issued]

Doc. No. _____

Page No. _____

Book No. _____

Series of _____. _____.

Annex C

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"