MINUTES OF THE PRE-BID CONFERENCE

SUPPLY AND DELIVERY OF VARIOUS LABORATORY SUPPLIES OF CY 2025
PUBLIC BIDDING NO. VMC-2026-018

November 13, 2025, 1:00 PM

The following were present during the conference:

BIDS & AWARDS COMMITTEE:
Ms. Shirlene V. Vianzon – Chairperson
Mr. Edsel S. Martin – Vice-Chairperson
Mr. Juan B. Sapasap, Jr. – BAC Member
Ms. Cherryl Ann L. Toyocan – BAC Member (via zoom)
Ms. Mary Grace T. Santillan – Provisional Member
Ms. Catherine F. Sofia – Provisional Member

BAC SECRETARIAT:
Ms. Ligaya Ubalde - Head, BAC Secretariat
Kristine Joy R. Manuel
Ms. Angelita B. Dayego
Ms. Aileen Pacheco
Ms. Kezia-Therese M. Guevarra
Mr. Lester John Jake Divino
Ms. Christallyne Castro - In-Charge
Ms. Aileen Cali
Ms. Diana Pulido

OBSERVER/TWG/END-USER:
Ms. Maria Josefina A. Bartolo-Cariaga, TWG- Laboratory Supplies
Ms. Celestine N. Lim, TWG- Laboratory Supplies
Mr. Roderick R. Balagtas – Observer, Procurement Section
Ms. Rufina Vadil – Observer, Budget Section
Mr. Raymund Joe Macuana– Observer, Accounting Section

PARTICIPANTS / PROSPECTIVE BIDDERS:

1. Mr. Kharlo Emerenciana – Edcor Medical and Laboratory Supplies
2. Mr. Akeem Villagracia – Diagnostika Philippines, Inc.
3. Ms. Danica Ley Figueroa – Trace Biomedical Corp.
4. Ms. Amor Domdom – Lifesource Enterprises
5. Mr. Norvyn T. Rubin – Trulaboratories Corporation
6. Ms. Karen L. Marinay – Medical Center Trading Corporation
7. Ms. Louella Daquigan – Zafire Distributors, Inc.
8. Mr. Gil Kenneth Caballero – Whelsycare Medical Device & Laboratory Reagent Trading
9. Mr. Aldrix Panopio – VG & GE Trading Diagnostic, Inc.
10. Mr. Roniel Dizon – Singapore Diagnositc
11. Mr. Renz Ticsay – QuidelOrtho Philippines, Inc.
12. Ms. Madel Echivarri – Distribution Solutions Philippines, Inc.
13. Ms. Kristine Roca – ROCHE Philippines, Inc.
14. Mr. Juliven Atienza – Lifeline Diagnostic Supplies, Inc.
15. Mr. Jetro Escoto – Zafire Distributors, Inc.
16. Ms. Ayessa Soro – Scientific Biotech Specialties, Inc.
17. Mr, Marco Cachola – TNC Everlight Philippines, Inc.
18. Mr. Jasper Daniel – Allied Hospital Supply International Corp.
19. Ms. Karissa Canlas – Pre-Ans Enterprises
20. Ms. Emilyn Rodriguez – IMS Global Enterprises, Inc.
21. Nathalie Loquias – Murex Diagnostic Products Specialists
22. Ms. Annie Alberto – Metro Drug, Inc.
23. Ms. Jemimah Tarayao – Zafire Distributors, Inc.

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

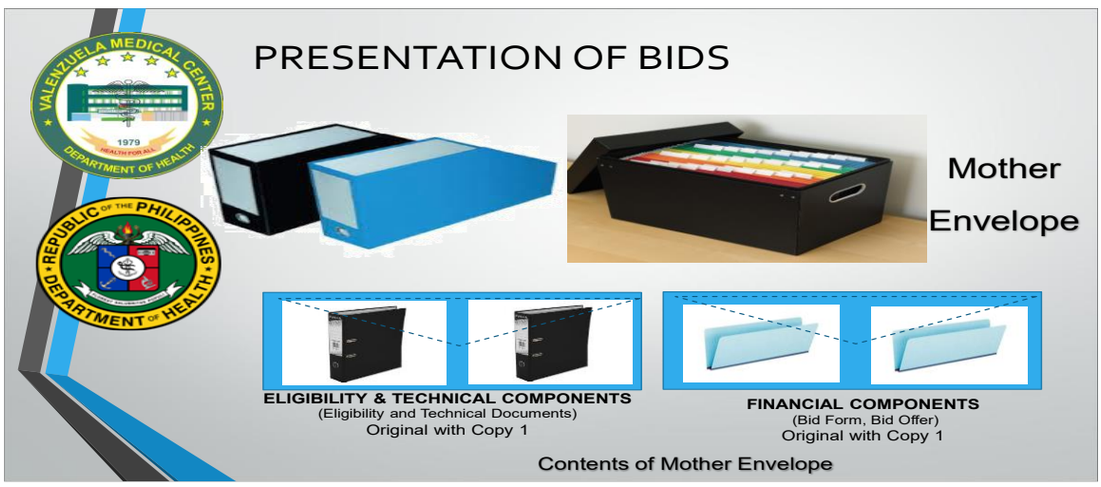
The conference started at 10:00am and was presided by **Ms. Shirlene V. Vianzon**, Chairperson of Bids & Awards Committee (BAC), held at the BAC Office, 2nd Floor, Annex Building, Valenzuela Medical Center, Padrigal St., Karuhatan, Valenzuela City. She acknowledged the presence of all representatives 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 it 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.
- 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.)
- 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.
- Notice of Award will be **emailed** to winning bidders. The following day will be counted as 1st day of receipt.
- 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 the New IRR of RA No. 12009, within the relevant period as provided in the Bidding Documents:

- Amount of the completed contract should be fifty percent (50%) of the total ABC based on category by lot.

CATEGORY BY LOT

- Lot 1: Clinical Microscopy Section **(Item Nos. 1-6, SLCC: 1,471,082.30)**
- Lot 2: Drug Testing **(Item Nos. 7-8, SLCC: 65,467.50)**
- Lot 3: Hematology Section **(Item Nos. 9-21, SLCC: 4,639,945.60)**
- Lot 4: Clinical Chemistry Section **(Item Nos. 22-45, SLCC: 13,157,068.85)**
- Lot 5: Microbiology Section **(Item Nos. 46-78, SLCC: 2,271,583.50)**
- Lot 6: Blood Bank Section **(Item Nos. 79-102, SLCC: 9,857,660.42)**
- Lot 7: Serology **(Item Nos. 103-130, SLCC: 4,903,811.00)**
- Lot 8: Histopathology **(Item Nos. 131-157, SLCC: 506,294.75)**
- Lot 9: Molecular Laboratory **(Item Nos. 158-162, SLCC: 681,625.00)**

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);
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. Certificate of Product Registration (CPR) from FDA.

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 Securities and Exchange Commission (SEC), Department of Trade and Industry (DTI) for sole proprietorship, or Cooperative Development Authority (CDA) for cooperatives or its equivalent document.
3. Mayor's or Business permit 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 one (1) Government or Private Hospital/Agency except from Valenzuela Medical Center ***(with at least Satisfactory Rating)*** - CY 2024 to present
7. Certificate of Good Performance with **SATISFACTORY rating** from laboratory section for existing suppliers for CY 2025
8. Special Power of Attorney (SPA) for authorized representative if OSS is Sole Proprietorship

9. For SLCC - Proof of evidence for Single Largest Completed Contract (SLCC) – Purchase Order or Notice of Award or Contract Agreement
10. Certificate from the manufacturer to distribute their products or Exclusive Distributorship or any equivalent document
11. License to Operate (LTO) from FDA
12. Certificate of Stocks Availability from Bidders (Notarized)
13. Bid Security

(For Lot 1: Clinical Microscopy Section)

- In addition to Bid Securing Declaration, bidder must submit at least two (2) of the ff.:
  1. The amount of not less than 58,843.29, if bid security is in cash.
  2. The amount of not less than 58,843.29, if bid security is in cashier's check.
  3. The amount of not less than 58,843.29, if bid security is in manager's check.
  4. The amount of not less than 147,108.23 if bid security is in bank draft.
  5. The amount of not less than 147,108.23 if bid security is in guarantee.
  6. The amount of not less than 147,108.23 if bid security is irrevocable LoC. or
  7. The amount of not less than 147,108.23 if bid security is Surety Bond.

NOTE: BID SECURITY FOR NUMBERS 1, 2 AND 3 IS SUBJECT FOR REFUND

(For Lot 2: Drug Testing)

- In addition to Bid Securing Declaration, bidder must submit at least two (2) of the ff.:
  1. The amount of not less than 2,618.70, if bid security is in cash.
  2. The amount of not less than 2,618.70, if bid security is in cashier's check.
  3. The amount of not less than 2,618.70, if bid security is in manager's check.
  4. The amount of not less than 6,546.75 if bid security is in bank draft.
  5. The amount of not less than 6,546.75 if bid security is in guarantee.
  6. The amount of not less than 6,546.75 if bid security is irrevocable LoC. or
  7. The amount of not less than 6,546.75 if bid security is Surety Bond.

NOTE: BID SECURITY FOR NUMBERS 1, 2 AND 3 IS SUBJECT FOR REFUND

(For Lot 3: Hematology Section)

- In addition to Bid Securing Declaration, bidder must submit at least two (2) of the ff.:
  1. The amount of not less than 185,597.82, if bid security is in cash.
  2. The amount of not less than 185,597.82, if bid security is in cashier's check.
  3. The amount of not less than 185,597.82, if bid security is in manager's check.
  4. The amount of not less than 463,994.56 if bid security is in bank draft.
  5. The amount of not less than 463,994.56 if bid security is in guarantee.
  6. The amount of not less than 463,994.56 if bid security is irrevocable LoC. or
  7. The amount of not less than 463,994.56 if bid security is Surety Bond.

NOTE: BID SECURITY FOR NUMBERS 1, 2 AND 3 IS SUBJECT FOR REFUND

(For Lot 4: Clinical Chemistry Section)

- In addition to Bid Securing Declaration, bidder must submit at least two (2) of the ff.:
  1. The amount of not less than 526,282.75, if bid security is in cash.
  2. The amount of not less than 526,282.75, if bid security is in cashier's check.
  3. The amount of not less than 526,282.75, if bid security is in manager's check.
  4. The amount of not less than 1,315,706.89 if bid security is in bank draft.
  5. The amount of not less than 1,315,706.89 if bid security is in guarantee.
  6. The amount of not less than 1,315,706.89 if bid security is irrevocable LoC. or
  7. The amount of not less than 1,315,706.89 if bid security is Surety Bond.

NOTE: BID SECURITY FOR NUMBERS 1, 2 AND 3 IS SUBJECT FOR REFUND

(For Lot 5: Microbiology Section)

- In addition to Bid Securing Declaration, bidder must submit at least two (2) of the ff.:
  1. The amount of not less than 90,863.34, if bid security is in cash.
  2. The amount of not less than 90,863.34, if bid security is in cashier's check.
  3. The amount of not less than 90,863.34, if bid security is in manager's check.
  4. The amount of not less than 227,158.35 if bid security is in bank draft.
  5. The amount of not less than 227,158.35 if bid security is in guarantee.
  6. The amount of not less than 227,158.35 if bid security is irrevocable LoC. or
  7. The amount of not less than 227,158.35 if bid security is Surety Bond.

NOTE: BID SECURITY FOR NUMBERS 1, 2 AND 3 IS SUBJECT FOR REFUND

(For Lot 6: Blood Bank Section)

- In addition to Bid Securing Declaration, bidder must submit at least two (2) of the ff.:
  1. The amount of not less than 394,306.42, if bid security is in cash.
  2. The amount of not less than 394,306.42, if bid security is in cashier's check.
  3. The amount of not less than 394,306.42, if bid security is in manager's check.

4. The amount of not less than 985,766.04 if bid security is in bank draft.
5. The amount of not less than 985,766.04 if bid security is in guarantee.
6. The amount of not less than 985,766.04 if bid security is irrevocable LoC. or
7. The amount of not less than 985,766.04 if bid security is Surety Bond.

NOTE: BID SECURITY FOR NUMBERS 1, 2 AND 3 IS SUBJECT FOR REFUND

(For Lot 7: Serology)

- In addition to Bid Securing Declaration, bidder must submit at least two (2) of the ff.:
  1. The amount of not less than 196,152.44, if bid security is in cash.
  2. The amount of not less than 196,152.44, if bid security is in cashier's check.
  3. The amount of not less than 196,152.44, if bid security is in manager's check.
  4. The amount of not less than 490,381.10 if bid security is in bank draft.
  5. The amount of not less than 490,381.10 if bid security is in guarantee.
  6. The amount of not less than 490,381.10 if bid security is irrevocable LoC. or
  7. The amount of not less than 490,381.10 if bid security is Surety Bond.

NOTE: BID SECURITY FOR NUMBERS 1, 2 AND 3 IS SUBJECT FOR REFUND

(For Lot 8: Histopathology)

- In addition to Bid Securing Declaration, bidder must submit at least two (2) of the ff.:
  1. The amount of not less than 20,251.79, if bid security is in cash.
  2. The amount of not less than 20,251.79, if bid security is in cashier's check.
  3. The amount of not less than 20,251.79, if bid security is in manager's check.
  4. The amount of not less than 50,629.47 if bid security is in bank draft.
  5. The amount of not less than 50,629.47 if bid security is in guarantee.
  6. The amount of not less than 50,629.47 if bid security is irrevocable LoC. or
  7. The amount of not less than 50,629.47 if bid security is Surety Bond.

NOTE: BID SECURITY FOR NUMBERS 1, 2 AND 3 IS SUBJECT FOR REFUND

(For Lot 9: Molecular Laboratory)

- In addition to Bid Securing Declaration, bidder must submit at least two (2) of the ff.:
  1. The amount of not less than 27,265.00, if bid security is in cash.
  2. The amount of not less than 27,265.00, if bid security is in cashier's check.
  3. The amount of not less than 27,265.00, if bid security is in manager's check.
  4. The amount of not less than 68,629.47 if bid security is in bank draft.
  5. The amount of not less than 68,629.47 if bid security is in guarantee.
  6. The amount of not less than 68,629.47 if bid security is irrevocable LoC. or
  7. The amount of not less than 68,629.47 if bid security is Surety Bond.

NOTE: BID SECURITY FOR NUMBERS 1, 2 AND 3 IS SUBJECT FOR REFUND

PR# 25-10-1665

ITEM NO.	UOM	ITEM DESCRIPTION	QTY.	ABC	TOTAL AMOUNT	AMENDMENT
		CLINICAL MICROSCOPY SECTION				
1	bot	Lugol's Iodine, 500ml	2	3,340.00	6,680.00	
2	pc	Pregnancy Test, cassette type, (Specification:Rapid, Chromatographic Immunoassay of Human Chorionic Gonadotropin(HCG) for urine and serum sample. Can detect lowest concentration of 25 mIU/mL)	4,303	50.00	215,150.00	
3	pc	Fecal Occult Blood,Fecal (Specification : IMMUNOCHEMICAL TEST Rapid chromatographic immunoassay for the qualitative detection of FOB in stool, individually pouched, can detect 25-50 ng/ml. Dietary restrictions are not necessary.)	515	136.44	70,266.60	
4	bot	Urine Reagent Strips, (Specification: 10 Parameters,	13	550.00	7,150.00	

ITEM NO.	UOM	ITEM DESCRIPTION	QTY.	ABC	TOTAL AMOUNT	AMENDMENT
		Gravity, Glucose, Protein, Blood [Erythrocytes, Hemoglobin], Leukocytes, Nitrite, Ketone, Bilirubin, Urobilinogen)100`s				
		Specification for No. 5 :For Fully-Automated Urine Physical, Chemical and Sediment Analyzer Reagent (free use of machine/tie-up) : Fully Automated Urine Physical and Chemistry Analyzer Specification: Should utilize Reflectance Photometry for test strip analysis and Refractometry for measuring Specific Gravity. It must achieve a minimum throughput of 200 samples per hour, with an aspiration volume of no more than 1mL of urine sample. Integration with a urine sediment analyzer is required. Additionally, consideration will be given to systems that offer extra parameters, provided they are supported by validated protocols. Fully Automated Urine Sediment Analyzer Specification: Digital Imaging analyzer capable of processing STAT samples, with an aspiration volume of no more than 1mL of urine and a minimum throughput of 100 samples per hour. The system should be able to identify and differentiate various formed elements in urine, including Red Blood Cells (RBC), White Blood Cells (WBC) and WBC clumps, Epithelial cells (squamous, non-squamous, transitional, and renal), casts (hyaline and pathologic), bacteria, crystals, yeasts, sperm cells, and mucus threads. Strongly preferred capable of analyzing body fluids (with FDA certification) and provide information on RBC morphology, specifically distinguishing between isomorphic and dysmorphic RBC. Other requirements: Provide consumables including disposable khan tubes or its equivalent, urine chemistry and sediments controls, and distilled water (if needed). Complete set of computer and printer with				

ITEM NO.	UOM	ITEM DESCRIPTION	QTY.	ABC	TOTAL AMOUNT	AMENDMENT
		scanner (including ink). Result printout customizable to end-user's format; Can generate printout of summary of reports run daily as specified by end-user; LIS ready, the winning bidder should shoulder any expenses related to LIS connectivity. LOT BID No. 5				
5	test	Fully Automated Urinalysis (Complete Set of Reagents, controls, calibrators, consumable and supplies and free use of machine)	18,000	145.00	2,610,000.00	
		DOTS SPUTUM MICROSCOPY				
6	box	Glass slides, frosted(Specification : 72's, optically clear, perfectly flat, well-polished, no rough or uneven edges, resistant to heat, scratches breakages and corrosion by chemicals, vacuum packaging, 26mmx76mm)	436	75.50	32,918.00	
7	pc	**Drug Testing Kit Met/THC** Specification : Rapid, Lateral flow chromatographic, immuno assay for qualitative detection of methamphetamine and cannabinoids tetrahydroxy cannabinol) FDA registered , with Specimen Container, polyethylene , sterile,disposable, wide mouth, screw cap, leak-proof, 60ml.	243	45.00	10,935.00	
8	pc	**Drug test liquid control** (Positive and Negative Control) /FDA registered at least 5ml	12	**10,000.00**	120,000.00	
		HEMATOLOGY SECTION				
		For Fully-automated Hematology Analyzer (6-part diff) Reagent(free use of machine with complete set of computer and printer ; Result printout customizable to end-user's format; Can generate printout of summary of reports run daily as specified by end-user;LIS ready, can test other body fluids)the winning bidder should shoulder any expenses related to LIS connectivity. Throughput : 80-100 samples/hour in WB mode with autoloader, closed system. Should ran stat mode. Aspiration volume : minimum of at least 70-90ul in				**For Fully-automated Hematology Analyzer (6-part diff) Reagent(free use of machine with complete set of computer and printer ; Result printout customizable to end-user's format; Can generate printout of summary of reports run daily as specified by end-user;LIS ready, can test other body fluids)the winning bidder should shoulder any expenses related to LIS connectivity. Throughput : at least**

ITEM NO.	UOM	ITEM DESCRIPTION	QTY.	ABC	TOTAL AMOUNT	AMENDMENT
		consideration of pedia and neonatal patients. Should generate 6 part differential including immature cells (must be FDA certified). Should be capable of running other body fluids (must be FDA certified) with volume aspiration of at least 70-90 ul. Should generate auto NRBC correction. Automated Reticulocyte count capable. Capable of platelet flourescence. Control's shelf life must be at least 1 month upon delivery. Should be able to generate printable LJ chart of controls. Shall provide back up machine (6 part differential) that has sample aspiration of not more than 25-30 ul for neonatal patient. Should be capable of body fluids analysis (Must be FDA certified). All information and specification must have supporting documents. Automated registry of controls. Principle : Combination of Fluorescence Flow Cytometry and Electrical Impedance method with Quality Control Flagging. 24/7 after service required upon machine malfunction or failure. Set of reagents must consists of Diluent, Lysing reagents, staining reagents, controls. At least 500 test/set excluding controls and calibration running. LOT BID item no. 9				80-100 samples/hour in WB mode with autoloader, closed system. Should ran stat mode. Aspiration volume : minimum of at least 70-90ul in consideration of pedia and neonatal patients. Should generate 6 part differential including immature cells (must be FDA certified). Should be capable of running other body fluids (must be FDA certified) with volume aspiration of at least 70-90 ul. Should generate auto NRBC correction. Automated Reticulocyte count capable. Capable of platelet flourescence. Control's shelf life must be at least 1 month upon delivery. Should be able to generate printable LJ chart of controls. Shall provide back up machine (6 part differential) that has sample aspiration of not more than 25-30 ul for neonatal patient. Should be capable of body fluids analysis (Must be FDA certified or its equivalent). All information and specification must have supporting documents. Automated registry of controls. Principle : Combination of Fluorescence Flow Cytometry and Electrical Impedance method with Quality Control Flagging. 24/7 after service required upon machine malfunction or failure. Set of reagents must consists of Diluent, Lysing reagents, staining reagents, controls. At least 500

ITEM NO.	UOM	ITEM DESCRIPTION	QTY.	ABC	TOTAL AMOUNT	AMENDMENT
						test/set excluding controls and calibration running. LOT BID item no. 9
9	test	**Complete Blood Count** :Set of Reagents, (Diluent, Lysing Reagents, Controls, et al with computer, printer and peripherals,excluding controls and calibration running) 2-machine installation	52,500	110.00	5,775,000.00	
10	set	**Differential or Wright's Stain for Blood Smear** Specification : Composition: ✅ Volume: 250mL ✅ Solutions: 1 to 4 Solutions ✅ Application: Hematology, Clinical Laboratory, Blood Smear Staining. ✓ Fast Results – Delivers high-quality staining in less than 1 minute ✓ Brilliant & Reproducible – Ensures clear, stable, and consistent stains ✓ Highly Stable Solutions – Prevents degradation for long-term usability ✓ Comparable to Standard Pappenheim Staining – Achieves high diagnostic accuracyPackaging: Approximately 500 tests , Stability: Must be stable for a minimum of 12 months at room temperature. Other Inclusions: Stained smear must appear pinkish gray. Microscopically, erythrocytes stain must appear pink to orange. Eosinophils must appear bright orange. Platelets must appear purple granules. Neutrophils cytoplasm must appear light pink. And lymphocytes cytoplasm must appear light to medium blue.	16	3,710.00	59,360.00	
11	box	Blood Lancet, g21,twist,200's per box	90	207.83	18,704.70	
12	pack	**Evacuated Tube - EDTA, K2, 100's, 2 ml** (Specification: 1) sterile, 2) Each tube is clearly labeled indicating: additive content, approximate blood draw, indicator, expiration and lot #, whether tube interior is sterile or non-sterile and field for patient name, age and ward 3) Safety - with safety closure that eliminates the risk of contact with blood left on stopper, easy and efficient re-insertion	850	450.00	382,500.00	

ITEM NO.	UOM	ITEM DESCRIPTION	QTY.	ABC	TOTAL AMOUNT	AMENDMENT
		of the closure, closure does not pop-out when returning cap (when transfering blood from syringe) 4) Tube made of plastic, precisely controlled vacuum, ease of penetration (to rubber stopper) 5) can stand centrifugation for 10mins at 3500rpm 6) Sealed properly using the desealer of TLA 7)must be compatible with the existing analyzers of Clinical Chemistry and Hematology)				
13	pack	**Evacuated Tube - Na Citrate**, (Specification: 3.2%, blue top, 100's, 1.8 ml max bld vol., disposable , sterile, plastic, for single use, with label indicating additive content, approximate blood draw, expiration and lot #, field for patient name, age and ward)	130	466.00	60,580.00	
14	box	**Needle, multisample, flash back blood collection,** Specification : 100's/box, sterile, gauge 21, multiple sample draw, transparent/clear hub, with adaptor per box.	15	1,237.50	18,562.50	-
15	pack	**Micro blood collecting tubes, EDTA, K2**(Specification: unbreakable plastic tube with K2 EDTA additive , at least 50pcs per pack,fill blood volume 0.5ml, 50's/pack)	400	343.33	137,332.00	
16	bot	**Reticulocyte Stain, 100 ml** (Specification: Brilliant cresyl blue in normal saline solution to examine reticulocytes in blood film.Intense blue solution.Clear without any particles. Stable below 20 ^{0}C-30^{0}C temperature)	1	3,472.00	3,472.00	
17	bot	**WBC Diluting Fluid, 1L**	1	1,100.00	1,100.00	
		For Fully-Automated Coagulation Analyzer Reagent free use of machine, with complete set of computer and printer and provision of ink and other consumables. Printout customizable to end-user's format; including controls, cuvettes/sample wells (including beads and distilled water if needed);Shall generate printout of summary of reports run daily as specified by end-user;LIS ready, the winning bidder should shoulder any expenses related to LIS connectivity. Provision of consumable (e.g. cuvettes,				

ITEM NO.	UOM	ITEM DESCRIPTION	QTY.	ABC	TOTAL AMOUNT	AMENDMENT
		distilled water if needed pipettors for reagent constitution, Thermal paper for manual printing of result). Principle : Clotting detection (Scattered Light Detection), Chromigenic (colorimetric method) Throughput : 40-60 test per hour, Volume : mst be at least 50 ul plasma, control's shelf life must be 4-5 days after reconstitution. with inserts provided. Reagent stability on board mst not be less than 3 days with insert provided. Shall generate printable LJ chart for controls. Immunoassay - latex ; Photo-optical Detection Method.24/7 after service required upon machine malfunction or failure. Lot Bid for item no. 18-21				
18	box	Prothrombin Time Reagent, at least 360 test /box	36	37,835.00	1,362,060.00	
19	box	Partial Thromboplastin Time Reagent+Calcium Chloride, at least 340 test / box	36	39,330.00	1,415,880.00	
20	box	Normal Plasma Control for Coagulation Reagent	5	4,534.00	22,670.00	
21	box	Pathologic Plasma Control for Coagulation Reagent	5	4,534.00	22,670.00	
		CLINICAL CHEMISTRY SECTION				
		For Fully Automated Integrated Floor Type Chemistry Analyzer (Reagent Tie- Up ;free use of machine with complete set of computer and printer including controls, calibrators and other consumables of the machine) (Througput : at least 400-800 tests per hour)(With or without Water supply ; if with water : Deionized Water should be shouldered by the supplier; Machine should not exceed 10L of water requirement per hour)(Preventive Maintenance and calibration should be done on a monthly basis)(Sample Volume : 2.5 to 11ul ; Sample Type: Serum, Plasma, Urine, CSF & Whole Blood) (Onboard sample capacity : continuous access & STAT interrupt capability) (With sample clot detection) (With real time Quality Control and calibration available) (Automatic : re-run, manual reruns , and				For Fully Automated Integrated Floor Type Chemistry Analyzer (Reagent Tie- Up ;free use of machine with complete set of computer and printer including controls, calibrators and other consumables of the machine) (Througput : at least 400-800 tests per hour)(With or without Water supply ; if with water : Deionized Water should be shouldered by the supplier; Machine should not exceed 10L of water requirement per hour)(Preventive Maintenance and calibration should be done on a monthly basis)(Sample Volume : 2.5 to 11ul ; Sample Type: Serum, Plasma, Urine, CSF & Whole Blood) (

ITEM NO.	UOM	ITEM DESCRIPTION	QTY.	ABC	TOTAL AMOUNT	AMENDMENT
		repeats)(Machine can measure directly the unconjugated bilirubin, reagents must be ready to use and should not require manual reconstitution or mixing of different reagent components, Machine should have clot correction, bubble detection, level sensing, and short sample detection, Machine should be able to scan barcoded samples) Time for single result : 2.5 minutes to 9 minutes. Result printout customizable to end-user's format; Can generate printout of summary of reports run daily as specified by end-user;LIS ready, the winning bidder should shoulder any expenses related to LIS connectivity. With recent excellent EQAS result from their installed institution. LOT BID (item no. 22-40)				Onboard sample capacity : continuous access & STAT interrupt capability) (With sample clot detection) (With real time Quality Control and calibration available) (Automatic : re-run, manual reruns , and repeats)(Machine can measure directly the unconjugated bilirubin, reagents must be ready to use and should not require manual reconstitution or mixing of different reagent components, Machine should have clot correction, bubble detection, level sensing, and short sample detection, Machine should be able to scan barcoded samples) Time for single result : 2.5 minutes to 10 minutes. Result printout customizable to end-user's format; Can generate printout of summary of reports run daily as specified by end-user;LIS ready, the winning bidder should shoulder any expenses related to LIS connectivity. With recent excellent EQAS result from their installed institution. LOT BID (item no. 22-40)
22	test	Albumin	7,500	68.51	513,825.00	
23	test	Alkaline Phosphatase	6,000	91.31	547,860.00	
24	test	ALT/SGPT	16,500	103.76	1,712,040.00	
25	test	AST/SGOT	16,500	103.69	1,710,885.00	
26	test	Bilirubin, Total and Direct	3,900	147.33	574,587.00	
27	test	Calcium	1,500	97.36	146,040.00	
28	test	Amylase	2,200	155.11	341,242.00	
29	test	Cholesterol, Total	12,000	81.05	972,600.00	
30	test	Creatinine	27,900	78.79	2,198,241.00	
31	test	Glucose	12,000	62.25	747,000.00	
32	test	HDL Cholesterol and LDL	12,000	207.98	2,495,760.00	
33	test	LDH	6,000	97.35	584,100.00	

ITEM NO.	UOM	ITEM DESCRIPTION	QTY.	ABC	TOTAL AMOUNT	AMENDMENT
34	test	Phosphorous	3,000	164.00	492,000.00	
35	test	Total Protein	6,600	75.53	498,498.00	
36	test	Triglycerides	12,000	117.62	1,411,440.00	
37	test	Urea Nitrogen	25,500	69.77	1,779,135.00	
38	test	Uric Acid	9,600	132.72	1,274,112.00	
39	test	Protein OTHER body fluids (urine/csf)	3,300	180.00	594,000.00	
40	test	Lipase	2,200	424.08	932,976.00	
		For Electrolytes Machine Analyzer (Sodium, Potassium, Chloride, Ionized Calcium and Ionized Magnessium) reagent tie-up and complete set of computer and printer including controls, calibrators and other consumables like electrode,the winning bidder should shoulder any expenses related to LIS connectivity. LOT BID for item no. 41				
41	test	**Electrolytes Reagents** (Sodium, Potassium, Chloride, Ionized Calcium and Ionized Magnesium) Specification : Complete Set of Reagents, controls, and supplies (consummables); with auto sampler and STAT position for emergency assay; For whole blood, plasma, serum and urine samples.	34,000	168.00	5,712,000.00	
		For HBA1c Machine Analyzer reagent free use of machine with complete set of computer and printer; Result printout customizable to end-user's format; Can generate printout of summary of reports run daily as specified by end-user;LIS ready, the winning bidder should shoulder any expenses related to LIS connectivity. Lot bid 42				
42	test	**HBA1C** (Specification : Complete Set of Reagents, controls, calibrators and supplies; High Performance Liquid Chromatography (HPLC) Method; NGSP and IFCC Certified; Fully automated; User-friendly; flexible (different sample tubes can be loaded in any order in any rack); with sample loader and STAT position for emergency assay; compact size ,Bench top)	7,200	141.00	1,015,200.00	
43	roll	Parafilm	1	2,900.00	2,900.00	
44	pack	Pipette tips - blue, 500's	10	393.67	3,936.70	
45	bot	OGTT 75 grams	420	128.00	53,760.00	

ITEM NO.	UOM	ITEM DESCRIPTION	QTY.	ABC	TOTAL AMOUNT	AMENDMENT
		MICROBIOLOGY SECTION				
46	set	Gram's Stain Reagent, 4x 250ml bottle/set	13	2,833.00	36,829.00	
47	bot	Potassium hydroxide (KOH), 10% , 500ml bottle	1	760.00	760.00	
48	pc	Inoculating loop, 1ul sterile, individually wrapped, disposable	2,800	4.85	13,580.00	
49	pc	Inoculating loop, 10ul sterile, individually wrapped, disposable	7,000	4.85	33,950.00	
50	bot	0.45% Saline Solution , 500 ml	54	1,080.00	58,320.00	
		Fully-automated system for Identification and Susceptibility Testing of Bacteria and Yeasts, and Blood Culture reagent free use of machine, includes analyzers and computer with peripherals, printer;Result printout customizable to end-user's format; Can generate printout of summary of reports run daily as specified by end-user;LIS ready and the winning bidder should shoulder any expenses related to LIS connectivity. LOT BID 51-52				
51	bot	Standard Pediatric Blood Culture Media with ARD	2,750	329.67	906,592.50	
52	bot	Standard Adult Blood Culture Media with ARD	4,550	329.67	1,499,998.50	
53	plate	Prepared plated media- Blood Agar Plate (Trypcase soy w/ sheep's blood)	3,900	83.33	324,987.00	
54	plate	Prepared plated media- Chocolate Agar Plate w/ polyvitex	3,600	83.33	299,988.00	
55	plate	Prepared plated media- Mac Conkey Agar Plate with crystal violet	3,900	83.33	324,987.00	
56	cart	Bacitracin disc (Taxo A), 50's disc/cart 10IU/IE/ul	2	1,375.00	2,750.00	
57	cart	Optochin disc (Taxo P), 50's disc/cart, 5mcg	2	1,162.50	2,325.00	
58	cart	Taxo V	1	1,600.00	1,600.00	
59	cart	Taxo X	1	1,600.00	1,600.00	
60	cart	Taxo X+V	1	1,600.00	1,600.00	
61	bot	Coagulase test (Rabbit Plasma / Fetal Bovine)	5	3,750.00	18,750.00	
62	pack	Oxidase Disc / strip,50's/box	12	6,850.00	82,200.00	
63	cart	Nitrocefin disk /Cefinase Paper Disc, 50 discs	1	5,200.00	5,200.00	
64	bot	Sheep Blood defibrinated, 100ml	2	1,500.00	3,000.00	
65	bot	Dehydrated Culture Media, 500 grams- Blood Agar Base	1	5,500.00	5,500.00	
66	bot	Dehydrated Culture Media, 500 grams- Brain Heart Infusion Brot	1	5,500.00	5,500.00	

ITEM NO.	UOM	ITEM DESCRIPTION	QTY.	ABC	TOTAL AMOUNT	AMENDMENT
67	bot	Dehydrated Culture Media, 500 grams- Bile Esculin	1	5,500.00	5,500.00	
68	bot	Dehydrated Culture Media, 500 grams- Citrate Media	1	5,500.00	5,500.00	
69	bot	Dehydrated Culture Media, 500grams- Gonococcal Agar	1	6,500.00	6,500.00	
70	bot	Dehydrated Culture Media, 500 grams- Haemophilus Test Medium	1	10,000.00	10,000.00	
71	rack	Pipette Tips Filter , blue, racked, sterile, 1000ul, 96's/rack	12	1,100.00	13,200.00	
72	rack	Pipette Tips, filter, 200ul, yellow, racked, sterile, 96's/rack	13	1,100.00	14,300.00	
		Fully-automated system for Identification and Susceptibility Testing of Bacteria and Yeasts, and Blood Culture reagent free use of machine, includes analyzers and computer with peripherals, printer;Result printout customizable to end-user's format; Can generate printout of summary of reports run daily as specified by end-user;LIS ready and the winning bidder should shoulder any expenses related to LIS connectivity. LOT BID: 73-75				
73	box	Automated ID for Gram Negative Bacilli, 20 cards/box	44	7,400.00	325,600.00	
74	box	Automated Susceptibility Tests for Gram Negative Bacilli, 20 cards/box	44	7,400.00	325,600.00	
75	box	Automated ID for Gram Positive Cocci, 20 cards/box	4	7,400.00	29,600.00	
76	pc	Amies's Transport Media, sterile transport medium swab	1500	102.50	153,750.00	
77	bot	Additive/supplement to Thayer Martin Medium for Cultivatin of nutritionally fastidious microorganism (Isovitalex or Vitox supplement)	1	7,100.00	7,100.00	
78	pc	Anaerobic Gaspak Jar/Candle Jar	1	16,500.00	16,500.00	
		BLOOD BANK SECTION				
79	vial	Anti-A, monoclonal (IgM/IgG), 10ml/vial, color:blue , Potency titer must not be lower than 1:1000	156	555.00	86,580.00	
80	vial	Anti-B, monoclonal (IgM/IgG), 10ml/vial, color:yellow , Potency titer must not be lower than 1:1000	154	555.00	85,470.00	
81	vial	Anti-D, monoclonal (IgM/IgG), 10ml/vial, colorless , Potency titer must not be lower than 1:1000	209	600.00	125,400.00	

ITEM NO.	UOM	ITEM DESCRIPTION	QTY.	ABC	TOTAL AMOUNT	AMENDMENT
		BLOOD BANK AND BLOOD TRANSFUSION SERVICE				
82	vial	Anti-A,B, monoclonal (IgM/IgG), 10ml/vial, colorless , Potency titer must not be lower than 1:1000	10	1,083.75	10,837.50	
83	vial	Anti-Human Globulin - 10ml/vial	1	782.75	782.75	
84	vial	RAM/LISS - 10ml/vial	1	782.75	782.75	
85	pc	Blood Bags, Plastic, triple, CPDA-1,sharp, silicon coated needle gauge sixteen (16), with serial number on the pilot tube, with needle guard or its equivalent Capacity-450 mL, sterile with diversion pouch	4,860	550.00	2,673,000.00	
86	bot	Giemsa Stain solution, 500 ml/bottle	1	2,812.00	2,812.00	
87	box	Blood Transfer bags, 300ml, 25's/box	5	3,600.00	18,000.00	
88	pack	**Evacuated Tube-Plain, plastic, with gel clot activator 100's, 4ml** (Specification: 1) sterile, 2) Each tube is clearly labeled indicating: additive content, approximate blood draw, indicator, expiration and lot #, whether tube interior is sterile or non-sterile and field for patient name, age and ward 3) Safety - with safety closure that eliminates the risk of contact with blood left on stopper, easy and efficient re-insertion of the closure, closure does not pop-out when returning cap (when transfering blood from syringe) 4) Tube made of plastic, precisely controlled vacuum, ease of penetration (to rubber stopper) 5) can stand centrifugation for 10mins at 3500rpm	862	1,081.50	932,253.00	
89	pack	Cryo Tube, Internally threaded, polypropylene vials for cryogenic storage of cells, blood, serum, and other biological specimens 1 - 2 ml capacity; cap incorporates a special silicone gasket, 50's/ pack	3	1,000.00	3,000.00	
		Reagent free use of machine for Hemoglobiometer - 2 machine installation (Portable): LOT BID : 90				
90	box	Microcuvettes for Hemoglobinometer, (For hemoglobin) 200's/box with free controls	68	6,000.00	408,000.00	
91	pack	Pipette tips - yellow, 1000's	10	652.50	6,525.00	
92	box	Tube Wafer 72's	2	22,000.00	44,000.00	

ITEM NO.	UOM	ITEM DESCRIPTION	QTY.	ABC	TOTAL AMOUNT	AMENDMENT
93	pc	Test Tube 5ml	2,500	4.10	10,250.00	
		Reagent free use of machine for Fully-Automated Blood Typing, Cross-matching and Antibody Screening with complete set peripherals Bar code scanner and barcode printer(LOT BID), LIS ready and the winning bidder should shoulder any expenses related to LIS connectivity., with complete set of computer and printer, result printout customizable to end user's format, with current and valid certificates of product registration (FDA). Fully automated immunohematological gel card technology.LOT BID 94-97)				
94	test	Complete set of reagents and consumables for Cross-matching	5,832	124.12	723,867.84	
95	test	Complete set of reagents and consumables for Patient Antibody Screening,	3,168	155.00	491,040.00	
96	test	Complete set of reagents and consumables for Donor Antibody Screening	6,480	108.00	699,840.00	
97	test	Complete set of reagents and consumables for Blood Typing	16,128	250.00	4,032,000.00	
		Reagent free use of machine, Fully automated Immunology analyzer, With current and valid certifcate of product registration (FDA). LIS ready. Reagent expiry at least 18 months upon delivery. With free consumables including distilled water for wash buffer solutions.Lot Bid 98				
98	test	Malaria Ag (EIA)	8,960	134.00	1,200,640.00	
		BLOOD BANK SECTION				
		Reagent free use of machine, Shall provide for FREE latest model Fully automated Immunology analyzer (HBsAg, Anti-HCV, HIV Ag & Ab, Syphilis), floor standing for Blood Bank Section. With current certifcate of product registration (FDA) and SACCL evaluation (lot bid). Throughput of at least 200 test per hour, sample volume 10-50uL, capacity of up to 100 samples with continuous loading. Reagent capacity of at least 18 reagents. With complete set of computers and printer, result print out customizable to end user				

ITEM NO.	UOM	ITEM DESCRIPTION	QTY.	ABC	TOTAL AMOUNT	AMENDMENT
		format, can generate printout summary of reports run daily as specified. The winning bidder should shoulder any expenses related to LIS connectivity, with controls and consumables. The fully automated equipment necessary FLOOR TYPE for the testing, data processing and analysis, generation of results, and efficient system operation with the following features: a. Batch and random-access analyzer using Enhanced Chemiluminescence Assay or Chemiluminescent Microparticle Immunoassay (CMIA) as test methodology, using samples such as: serum, plasma or whole blood, using Levy –Jennings, and Westgard rules for quality controls; b. Quantitative detection of Transfusion Transmissible Infection (TTI) markers for Hepatitis B, HIV, HCV, and Syphilis (e.g. HBsAg, HCV antibody and Antigen, HIV 1 and 2 antibodies, and anti-TP) in serum or plasma samples; c. Preferred reagents shall have specificity and sensitivity levels of not less than 99.5%. d. Must be capable of sample and reagent identification by barcode scanner e. All assays shall have a current or latest proof of Kit Evaluation from STD AIDS Cooperative Central Laboratory (SACCL) or from the National Reference Laboratory for Transfusion Transmissible Infections (Research Institute for Tropical Medicine); f. Uses e-connectivity technology g. Uses single-use tips and single use cuvettes or microwells or its equivalent;				
		h. Must be able to detect for clots, bubbles and short samples; i. No preparation, no mixing or reconstitution required for reagents; j. On board stability of reagents should be two months or more; k. No manual primes, purges, washes, or tubing maintenance;				

ITEM NO.	UOM	ITEM DESCRIPTION	QTY.	ABC	TOTAL AMOUNT	AMENDMENT
		l. No water or drain required and with self-contained onboard waste management; m. No daily calibrations or calibration checks (stable calibration for up to 20 days or more) n. Capable of running small sample volume (not more than 100 uL per assay) o. Dimension of analyzer shall fit in the allotted space in the designated area in the Blood Bank section.LOT BID : 99-102				
99	test	HBsAg (CLIA or CMIA)	10,000	126.30	1,263,000.00	
100	test	HCV, Ab (CLIA or CMIA)	9,600	352.07	3,379,872.00	
101	test	HIV 1 & 2 or HIV Ag/Ab combination (CLIA or CMIA)	10,200	193.00	1,968,600.00	
102	test	Syphilis TP, highly sensitive and specific test for treponemal antibodies (CLIA or CMIA)	9,600	161.33	1,548,768.00	
		SEROLOGY				
		Reagent free use of machine for Fully Automated Immunology Analyzer Floor Type for Serology section, with controls and other consumables of the machine,with current and valid certificates of product registration (FDA) Result printout customizable to end-user's format; Can generate printout summary of reports run daily as specified by end-user;LIS ready and the winning bidder should shoulder any expenses related to LIS connectivity. with complete set of computers and printer.(LOT BID):103-122 1. Shall provide for FREE latest model of fully automated equipment necessary FLOOR TYPE for the testing, data processing and analysis, generation of results, and efficient system operation with the following features: a. Batch and random-access analyzer using Enhanced Chemiluminescence Assay or Chemiluminescent Microparticle Immunoassay (CMIA) as test methodology, using samples such as: serum, plasma or whole blood, using Levy –Jennings, and Westgard rules for quality controls; b. Must be capable of sample and reagent identification by				Reagent free use of machine for Fully Automated Immunology Analyzer Floor Type for Serology section, with controls and other consumables of the machine,with current and valid certificates of product registration (FDA) Result printout customizable to end-user's format; Can generate printout summary of reports run daily as specified by end-user;LIS ready and the winning bidder should shoulder any expenses related to LIS connectivity. with complete set of computers and printer.(LOT BID):103-122 1. Shall provide for FREE latest model of fully automated equipment necessary FLOOR TYPE for the testing, data processing and analysis, generation of results, and efficient system operation with the

ITEM NO.	UOM	ITEM DESCRIPTION	QTY.	ABC	TOTAL AMOUNT	AMENDMENT
		barcode scanner c. All assays shall have a current and valid certificate of product registration from the Food and Drug Administration (FDA). d. Uses e-connectivity technology: e. Uses single-use tips and single use cuvettes or microwells or its equivalent; f. Must be able to detect for clots, bubbles and short samples; g. No preparation, no mixing or reconstitution required for reagents; h. On board stability of reagents should be two months or more; i. No manual primes, purges, washes, or tubing maintenance; j. No water or drain required and with self-contained onboard waste management; k. No daily calibrations or calibration checks (stable calibration for up to 20 days or more) l. Capable of running small sample volume (not more than 100 uL per assay) m. Dimension of analyzer shall fit in the allotted space in the designated area in the SEROLOGY section. n. throughput of up to 100-130 test/hour				following features: a. Batch and random-access analyzer using Enhanced Chemiluminescence Assay or Chemiluminescent Microparticle Immunoassay (CMIA) as test methodology, using samples such as: serum, plasma or whole blood, using Levy –Jennings, and Westgard rules for quality controls; b. Must be capable of sample and reagent identification by barcode scanner c. All assays shall have a current and valid certificate of product registration from the Food and Drug Administration (FDA). d. Uses e-connectivity technology: e. Uses single-use tips and single use cuvettes or microwells or its equivalent; f. Must be able to detect for clots, bubbles and short samples; g. No preparation, no mixing or reconstitution required for reagents; h. On board stability of reagents should be two months or more; i. with or without manual primes, purges, washes, or tubing maintenance; j. with or without water or drain required and with self-contained onboard waste management (if with water, liquid should be provided by the bidder); k. No daily calibrations or calibration checks (stable calibration for up to 20 days or more)

ITEM NO.	UOM	ITEM DESCRIPTION	QTY.	ABC	TOTAL AMOUNT	AMENDMENT
						l. Capable of running small sample volume (not more than 100 uL per assay) m. Dimension of analyzer shall fit in the allotted space in the designated area in the SEROLOGY section. n. throughput of up to 100-130 test/hour
		2. Shall continously provide and replenish consumables including controls and calibrators ; 3. Shall assure replacement of unused reagents and consumables (delivered within six months of expiration date) before expiration and will provide a guarantee letter of replacement; 4. Shall replace reagents and consumables with three-months remaining shelf life before the expiration date; 5. Shall provide FREE transport, installation (including minor civil works) and regular maintenance and calibration of all equipment and replacement of parts or units duly coordinated with concerned laboratory staff and EFM personnel; 6. Supplier shall issue a commitment statement that it shall provide FREE training and seminar and/or re-training of laboratory staff and hospital biomedical technicians to make them competent in the use of the equipment; shall issue certificates of training to the personnel concerned; 7. Shall perform verification/validation of equipment in conjunction with laboratory staff, and provision of FREE reagents and consumables for that purpose; 8. Has the capacity to upgrade the equipment provided when necessary or when the need arises upon request by end user without additional costs to procured reagents; 9. Shall provide connection for Laboratory Information System (LIS) of choice of end				

ITEM NO.	UOM	ITEM DESCRIPTION	QTY.	ABC	TOTAL AMOUNT	AMENDMENT
		user and/or Hospital Information System (HIS) - shall be shouldered by the winning bidder. 10. Analyzer and accessory equipment must be brand new (equipment not more than three years old are acceptable); 11. Shall provide for FREE printer with scanner, paper and ink including maintenance and replacement if malfunctioning within forty-eight hours from notice; 12. Shall provide for FREE compatible uninterruptible power supply (UPS) and automatic voltage regulator (AVR) for the analyzer as well as corresponding computer hardware and softwareintegrated with the analyzer; 13. Supplier or distributor shall be responsible for securing the equipment and its accessories from rodents and other pests and shall immediately repair or replace the equipment and its accessories should damage occur due to these pests within forty-eight (48) hours from notice; 14. Shall provide FREE regular or scheduled preventive maintenance or calibration and ASAP repair or replacement of parts or units of the equipment for FREE duly coordinated with concerned laboratory staff and EFMS Biomedical personnel;				
		15. Shall ensure prompt response by designate engineer/s for correction of equipment failure or malfunction;16. Shall provide a back-up machine for FREE, including transportation and installation, in case of machine breakdown which cannot be resolved within 48 hours from notice;17. Shall shoulder the cost or reimburse the expenses for examinations sent out related to equipment failure or defects in the reagents or consumables which cannot be resolved within forty-eight hours from notice;18. Shall replenish				

ITEM NO.	UOM	ITEM DESCRIPTION	QTY.	ABC	TOTAL AMOUNT	AMENDMENT
		consumed reagents and consumables incidental to repeated test runs due to equipment failure or malfunction including that of the UPS and AVR, defects in reagent or consumables; or due to trial runs during trainings or retraining; or during scheduled preventive maintenance or calibration of equipment;19. Shall provide updated Material Safety Data Sheet (MSDS) for chemical reagents upon delivery of the item;20. Manufacturing or reagents and equipment are covered by relevant ISO certification or quality certification of similar nature.21. Products carried are covered by current Certificate of Product Registration or in the absence of the former, proof of renewal or updated Certificate of Exemption issued by the Philippine FDA;22. The supplier shall have installations of the same equipment and its accessories and using the same reagents in the Philippines. A certification from at least one institution having such installation shall be provided.23. The supplier shall submit a certificate of distributorship from the principal manufacturer of the equipment and reagents;24. Shall allow delivery on staggered or as per need basis;25. Supplier /distributor is preferably, but not necessarily, ISO 9001:2015 certified;26. Should there be persistence of erroneous, invalid, or inconsistent results that may jeopardize the safety of the patients or the quality of services provided despite attempts to repair, the contract shall be terminated at the discretion of the department. The supplier shall refund whatever is paid equivalent to the value of unused reagents or wasted reagents due to repeat analyses or repairs; or if not yet paid, shall not be paid for the said reagents. The supplier shall be compelled to				

ITEM NO.	UOM	ITEM DESCRIPTION	QTY.	ABC	TOTAL AMOUNT	AMENDMENT
		immediately remove the equipment from the section.				
103	test	Anti-HBc	700	356.33	249,431.00	
104	test	HBc IgM	700	436.50	305,550.00	
105	test	Anti-Hbe	500	410.00	205,000.00	
106	test	Hbe Ag	800	417.33	333,864.00	
107	test	HAV IgM	200	295.50	59,100.00	
108	test	Anti-HAV IgG	200	325.00	65,000.00	
109	test	Anti-Hbs	1,000	480.00	480,000.00	
110	test	T3	200	223.50	44,700.00	
111	test	T4	200	243.50	48,700.00	
112	test	TSH	2,400	257.67	618,408.00	
113	test	FT3	2,400	234.33	562,392.00	
114	test	FT4	2,400	244.33	586,392.00	
115	test	PSA	900	393.00	353,700.00	
116	test	CA 125	400	614.50	245,800.00	
117	test	CEA	500	429.50	214,750.00	
118	test	Troponin I, High Sensitivity	3,000	775.50	2,326,500.00	
119	test	Beta-HCG	400	611.00	244,400.00	
120	test	Procalcitonin	500	1,119.50	559,750.00	
121	test	Ferritin	500	414.50	207,250.00	
122	test	NT-proBNP	200	650.00	130,000.00	
		RAPID TEST KITS LOT BID 123-130				**Lot bid for item nos. 123-124**
123	pc	Treponema Pallidum Specific Rapid Test Kit, cassette type SACCL Evaluated	2,010	100.00	201,000.00	
124	pc	HBsAg Rapid Test Kit SACCL Evaluated	2,500	70.00	175,000.00	
						LOT BID for item nos. 125-126
125	pc	Dengue NS1Ag, 20's per box, with positive and negative control	1,800	200.00	360,000.00	
126	pc	Dengue IgG, IgM Rapid Test Kit with positive and negative control	1,800	250.00	450,000.00	
127	kit	Anti-Streptolysin O, Rapid Test 100 tests	2	3,347.50	6,695.00	
128	kit	C-Reactive Protein, Rapid Test 100 tests	4	3,060.00	12,240.00	
129	kit	Rapid test for Leptospira IgM & or w/ IgG, device/ card with positive and negative control	150	280.00	42,000.00	
130	pc	Rapid Antigen Test Kit for COVID 19 (Nasal swab), sensitivity 94%, specificity 94%, storage temperature 30 degress celcius.	3,600	200.00	720,000.00	
		HISTOPATHOLOGY				
131	gal	Buffered Neutral Formalin, 10%, 1 gallon	15	805.00	12,075.00	
132	gal	Ethyl Alcohol, absolute, 3.8L-4L	10	3,180.00	31,800.00	
133	gal	Ethyl Alcohol, 95%, 1 gallon	10	1,330.00	13,300.00	

ITEM NO.	UOM	ITEM DESCRIPTION	QTY.	ABC	TOTAL AMOUNT	AMENDMENT
134	gal	Xylene,For processing and staining of histological specimens, colorless, with certificate of analysis, 3.8L - 4L	10	4,200.00	42,000.00	
135	box	High profile Disposable blades 50's /box	5	9,000.00	45,000.00	
136	box	Low profile Disposable blades 50's /box	1	7,000.00	7,000.00	
137	pack	Thin Cover Slip or Micro cover glass, 24mm x 56mm, 100/pack	50	198.00	9,900.00	
138	bot	EA 50, 1L	5	4,000.00	20,000.00	
139	bot	Orange G-6, 1L	5	2,180.00	10,900.00	
140	bot	Eosin Y Stain, 1 liter	5	4,150.00	20,750.00	
141	bot	Harris Hematoxylin Stain, 1 liter	5	5,150.00	25,750.00	
142	gal	Acid Alcohol, 1%	10	1,350.00	13,500.00	
143	bot	Mounting Medium, 473 ml	1	8,050.00	8,050.00	Mounting Medium, 473-500 ml
144	box	Tissue Cassette 250's /box (pink/blue)	10	2,000.00	20,000.00	
145	bot	Ammonia Water/Scott's Tap 378.5 ml	1	2,250.00	2,250.00	
146	can	Spray for Rapid Freezing of fresh or paraffin embedded tissue, Ozone sale; CFC free; non-flammable; leaves no residue	1	3,360.00	3,360.00	
147	bot	Clear Frozen Section Compound Mounting Medium, 118ml dipper bottle	2	1,350.00	2,700.00	
148	kg	Paraffin wax, - pellet, melts at 56-60 degrees Celcius, excellent tissue penetration,excellent ribbon continuity,no crumbling or cracking	20	763.00	15,260.00	
149	bot	Immersion Oil, 30 ml	5	1,440.00	7,200.00	
		Antigen decloaker: for free of use, bench top with pressure chamber unit, built in slide rack (able to hold at least 70 slides per run), able to retrieve antigen within 60minutes or faster, digital display, working temperature of 60C-110C (temperature increment of 5C or lower), 220-230V (50-60Hz, 1000W). LOT BID 150-157				
150	kit	Polymer Detection System (contain pre-diluted reagents: Peroxidase Block, Protein Block, Post Primary Block, Polymer, DAB Chromogen, DAB Substrate Buffer (Polymer) and Hematoxylin)50 test	5	19,767.05	98,835.25	
151	bot	Epitope Retrieval Solution pH6, 1L, 1600 tests	3	31,010.13	93,030.39	

ITEM NO.	UOM	ITEM DESCRIPTION	QTY.	ABC	TOTAL AMOUNT	AMENDMENT
152	bot	Epitope Retrieval Solution pH9, 1L, 1600 tests	3	31,010.13	93,030.39	
153	bot	Wash Solution 10x conc., 1000ml, 640 tests	3	37,080.00	111,240.00	
154	bot	Primary Antibodies for ER, 7ml, 46 tests	3	31,028.50	93,085.50	
155	bot	Primary Antibodies for PR, 7ml, 46 tests	3	31,028.50	93,085.50	
156	bot	Primary Antibodies for HER2, 6ml, 46 tests	3	31,495.82	94,487.46	
157	pc	Pen, light blue hydrophobic reagent soluble in commonly used clearing agents; designed to minimize wastage of reagents by allowing the user to ring the tissue(s) or cells to be stained thereby localizing the staining reagents, 1000 tests	1	25,000.00	25,000.00	
		MOLECULAR LABORATORY				
		Supply and Delivery of the following inclusive of consumables,controls, extraction kit, VTM, and swabs LOT BID # 158 (compatible with the existing machine SANSURE):				
158	test	COVID-19 RT PCR nucleic acid and Diagnostic Kit	3,759	350.00	1,315,650.00	
159	box	20ul Filtered Tips, DnAse/RnAse free, sterile 96tips x 10 racks	2	4,500.00	9,000.00	
160	pc	Respirator Mask (Specification: NIOSH Approved for at least 95 percent filtration efficiency; Cushioning nose foam, collapsed resistant cup shape, two strap design with welded dual point attachment, Fit tested to molecular laboratory staff, Medium Size)	436	75.00	32,700.00	
161	box	2.0 ml microcentrifuge tube	14	100.00	1,400.00	
162	pack	Resealable (double sealed/zipper) Small Plastic Transparent Bags 5cm x 7cm, 100pcs/pack	45	100.00	4,500.00	
		TOTAL			**75,109,077.83**	

Other Matters:

The following matters were discussed, to wit:

- All quantity should be served and no loose items will be cancelled even the packaging do not conform to the required quantity. All requests for cancellation will be reflected to the Performance Evaluation of the Supplier.
- All packaging is acceptable provided that they met the total quantity requirement (per pieces/ per box)
- Content higher than the requirement is acceptable if advantageous to the government

PRICE SCHEDULE

- 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. Please include your packaging for each item to be bid.**
  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)
- Bid Bulletin will be posted, if any.
- Initial Product Registration from FDA is not allowed.
- 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).
- **Bid Opening will be on December 2, 2025 at 9:00am (Face-to-face)**

The pre-bidding conference was adjourned at 3:00 pm.

Prepared by:

AILEEN S. CALI, MPA (SGD)
BAC Secretariat

Noted by:

SHIRLENE V. VIANZON (SGD)
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 Proje*ct.

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[1]]*** 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"