

Republic of the Philippines DEPARTMENT OF HEALTH PHILIPPINE CHILDREN'S MEDICAL CENTER Bids and Awards Committee Quezon Avenue, Quezon City 1100

website: <u>www.pcmc.gov.ph</u> email: <u>bac@pcmc.gov.ph</u> Trunkline: 8588-9900 local 361/355 Telefax No.: 8924-0870

## **SECTION II**

## **Instructions to Bidders**

## Supply and Delivery of Supplies/Reagents/Consumables thru Reagent Tie-ups / Machine Placement Agreements for Three (3) Years

IB-2022-003 to IB-2022-022

#### 1. Scope of Bid

The **Philippine Children's Medical Center (PCMC)** wishes to receive Bids for the following Project/s:

IB NUMBER	ITEM DESCRIPTION	TOTAL ABC (for the 1 <sup>st</sup> year)
One (1) Lot Su of the followin	apply and Delivery of supplies/consumables under Machine Placement Agreement for t g:	hree (3) years
IB-2022-003	Apheresis kits with anticoagulant	1,254,400.00
IB-2022-004	Blood bag, quintuple with filter	8,100,000.00
IB-2022-005	Hemoglobin Screening for blood donors	867,000.00
IB-2022-006	Sterile Connecting Device	533,610.00
One (1) Lot Su the following:	pply and Delivery of reagents/consumables under Reagent Tie-up Agreement for three	e (3) years of
IB-2022-007	Automated Immunohaematology Analyzer	6,308,698.00
IB-2022-008	Automated Immunology Analyzers for Transfusion-Transmissible Infection (TTIs)	7,422,812.00
IB-2022-009	Blood Gas Analyzer	2,932,672.00
IB-2022-010	Automated Clinical Chemistry Analyzer	7,032,210.00
IB-2022-011	Coagulation Machine	2,814,725.00
IB-2022-012	Flow Cytometry Machine	7,160,546.82
IB-2022-013	Automated Blood and other Sterile Body Fluid Culture Machine	1,192,250.00
IB-2022-014	Automated Hematology Analyzer	5,457,924.00
IB-2022-015	Automated Identification and Susceptibility for Multi-Drug Organism Test Machine	793,506.00
IB-2022-016	Automated Identification and Susceptibility for Microorganism Test Machine	2,113,317.00
IB-2022-017	Automated Immunohistochemical Antibody Staining Analyzer	2,800,000.00
IB-2022-018	Immunology Analyzer for Hepatitis B profile, Thyroid profile, Anti-HAV IgM, CMV IgM, Toxoplasma IgM, Rubella IgM, AFP, B-HCG, Serum Ferritin, Cortisol, Intact PTH and Total Vitamin D Assays	3,000,975.00
IB-2022-019	Immunoanalyzer for Procalcitonin	3,152,890.71
IB-2022-020	Automated Urine Microscopic and Urine Chemistry Analyzer	1,627,387.20
IB-2022-021	Immunology Analyzer for Dengue Antigen and HBsAg Confirmatory Testing	1,597,000.00
IB-2022-022	Automated Analyzer for Methotrexate determination	3,453,071.52

The above Procurement Projects, the details of which are described in Section VII (Technical Specifications).

#### 2. Funding Information

- 2.1. The GOP through the source of funding as indicated below for GAA CY 2022 in the amount of Sixty-Nine Million Six Hundred Fourteen Thousand Nine Hundred Ninety-Five Pesos and 25/100 (Php 69,614,995.25).
- 2.2. The source of funding is:
  - a. GOCC and GFIs, the Corporate Operating Budget.

#### **3. Bidding Requirements**

The Bidding for the Project shall be governed by all the provisions of RA No. 9184 and its 2016 revised IRR, including its Generic Procurement Manuals and associated policies, rules and regulations as the primary source thereof, while the herein clauses shall serve as the secondary source thereof.

Any amendments made to the IRR and other GPPB issuances shall be applicable only to the ongoing posting, advertisement, or **IB** by the BAC through the issuance of a supplemental or bid bulletin.

The Bidder, by the act of submitting its Bid, shall be deemed to have verified and accepted the general requirements of this Project, including other factors that may affect the cost, duration and execution or implementation of the contract, project, or work and examine all instructions, forms, terms, and project requirements in the Bidding Documents.

#### 4. Corrupt, Fraudulent, Collusive, and Coercive Practices

The Procuring Entity, as well as the Bidders and Suppliers, shall observe the highest standard of ethics during the procurement and execution of the contract. They or through an agent shall not engage in corrupt, fraudulent, collusive, coercive, and obstructive practices defined under Annex "I" of the 2016 revised IRR of RA No. 9184 or other integrity violations in competing for the Project.

#### 5. Eligible Bidders

5.1. Only Bids of Bidders found to be legally, technically, and financially capable will be evaluated.

5.2.

- a. Foreign ownership exceeding those allowed under the rules may participate pursuant to:
  - i. When a Treaty or International or Executive Agreement as provided in Section 4 of the RA No. 9184 and its 2016 revised IRR allow foreign bidders to participate;
  - ii. Citizens, corporations, or associations of a country, included in the list issued by the GPPB, the laws or regulations of which grant reciprocal rights or privileges to citizens, corporations, or associations of the Philippines;
  - iii. When the Goods sought to be procured are not available from local suppliers; or
  - iv. When there is a need to prevent situations that defeat competition or restrain trade.
- 5.3. Pursuant to Section 23.4.1.3 of the 2016 revised IRR of RA No.9184, the Bidder shall have an SLCC that is at least one (1) contract similar to the Project the value of which, adjusted to current prices using the PSA's CPI, must be at least equivalent to:
  - a. For the procurement of Expendable Supplies: The Bidder must have completed a single contract that is similar to this Project, equivalent to at least twenty-five percent (25%) of the ABC of the item/s joined
  - b. For procurement where the Procuring Entity has determined, after the conduct of market research, that imposition of either (a) or (b) will likely result to failure of bidding or monopoly that will defeat the purpose of public bidding: the Bidder should comply with the following requirements: [Select either failure or monopoly of bidding based on market research conducted]
    - i. Completed at least two (2) similar contracts, the aggregate amount of which should be equivalent to at least *fifty percent* (50%) *in the case of non-expendable supplies and services or twenty-five percent* (25%) *in the case of expendable supplies*] of the ABC for this Project; and
    - ii. The largest of these similar contracts must be equivalent to at least half of the percentage of the ABC as required above.
- 5.4. The Bidders shall comply with the eligibility criteria under Section 23.4.1 of the 2016 IRR of RA No. 9184.

#### 6. Origin of Goods

There is no restriction on the origin of goods other than those prohibited by a decision of the UN Security Council taken under Chapter VII of the Charter of the UN, subject to Domestic Preference requirements under **ITB** Clause 18.

#### 7. Subcontracts

7.1. The Bidder may subcontract portions of the Project to the extent allowed by the Procuring Entity as stated herein, but in no case more than twenty percent (20%) of the Project.

The Procuring Entity has prescribed that:

a. Subcontracting is not allowed.

#### 8. Pre-Bid Conference

The Philippine Children's Medical Center will hold a Pre-Bid Conference on October 27, 2021 at 10:00A.M. through video conferencing via google meet which shall be open to prospective bidders, as indicated in paragraph 6 of the **IB**.

#### 9. Clarification and Amendment of Bidding Documents

Prospective bidders may request for clarification on and/or interpretation of any part of the Bidding Documents. Such requests must be in writing and received by the Procuring Entity, either at its given address or through electronic mail indicated in the **IB**, at least ten (10) calendar days before the deadline set for the submission and receipt of Bids.

#### **10.** Documents comprising the Bid: Eligibility and Technical Components

- 10.1. The first envelope shall contain the eligibility and technical documents of the Bid as specified in Section VIII (Checklist of Technical and Financial Documents).
- 10.2. The Bidder's SLCC as indicated in **ITB** Clause 5.3 should have been completed within **the past two (2) years** prior to the deadline for the submission and receipt of bids.
- 10.3. If the eligibility requirements or statements, the bids, and all other documents for submission to the BAC are in foreign language other than English, it must be accompanied by a translation in English, which shall be authenticated by the appropriate Philippine foreign service establishment, post, or the equivalent office having jurisdiction over the foreign bidder's affairs in the Philippines. Similar to the required authentication above, for Contracting Parties to the Apostille Convention, only the translated documents shall be authenticated through an apostille pursuant to GPPB Resolution No. 13-2019 dated 23 May 2019. The English translation shall govern, for purposes of interpretation of the bid.

#### 11. Documents comprising the Bid: Financial Component

- 11.1. The second bid envelope shall contain the financial documents for the Bid as specified in Section VIII (Checklist of Technical and Financial Documents).
- 11.2. If the Bidder claims preference as a Domestic Bidder or Domestic Entity, a certification issued by DTI shall be provided by the Bidder in accordance with Section 43.1.3 of the 2016 revised IRR of RA No. 9184.
- 11.3. Any bid exceeding the ABC indicated in paragraph 1 of the **IB** shall not be accepted.
- 11.4. For Foreign-funded Procurement, a ceiling may be applied to bid prices provided the conditions are met under Section 31.2 of the 2016 revised IRR of RA No. 9184.

#### 12. Bid Prices

- 12.1. Prices indicated on the Price Schedule shall be entered separately in the following manner:
  - a. For Goods offered from within the Procuring Entity's country:
    - i. The price of the Goods quoted EXW (ex-works, ex-factory, ex-warehouse, ex-showroom, or off-the-shelf, as applicable);
    - ii. The cost of all customs duties and sales and other taxes already paid or payable;
    - iii. The cost of transportation, insurance, and other costs incidental to delivery of the Goods to their final destination; and
    - iv. The price of other (incidental) services, if any, listed in e.
  - b. For Goods offered from abroad:
    - i. Unless otherwise stated in the **BDS**, the price of the Goods shall be quoted delivered duty paid (DDP) with the place of destination in the Philippines as specified in the **BDS**. In quoting the price, the Bidder shall be free to use transportation through carriers registered in any eligible country. Similarly, the Bidder may obtain insurance services from any eligible source country.

ii. The price of other (incidental) services, if any, as listed in Section VII (Technical Specifications).

#### 13. Bid and Payment Currencies

- 13.1. For Goods that the Bidder will supply from outside the Philippines, the bid prices may be quoted in the local currency or tradeable currency accepted by the BSP at the discretion of the Bidder. However, for purposes of bid evaluation, Bids denominated in foreign currencies, shall be converted to Philippine currency based on the exchange rate as published in the BSP reference rate bulletin on the day of the bid opening.
- 13.2. Payment of the contract price shall be made in:
  - a. Philippine Pesos.

#### 14. Bid Security

- 14.1. The Bidder shall submit a Bid Securing Declaration<sup>1</sup> or any form of Bid Security in the amount indicated in the **BDS**, which shall be not less than the percentage of the ABC in accordance with the schedule in the **BDS**.
- 14.2. The Bid and bid security shall be valid until *120 calendar days*. Any Bid not accompanied by an acceptable bid security shall be rejected by the Procuring Entity as non-responsive.

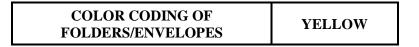
#### 15. Sealing and Marking of Bids

Use of indelible ink <u>color blue</u> shall be used by the authorized signatory in signing the required forms. *Strictly NO using of staple wire and thick materials for tab* 

The **First**  $(1^{st})$  **Envelope**, shall contain the following <u>Technical Component</u> accomplished in two (2) sets, each set filed in a folder

The Second  $(2^{nd})$  Envelope shall contain the <u>Financial Component</u> accomplished in three (3) sets, each set filed in a folder

#### All copies should be certified as true copy



#### LABEL ON THE ENVELOPE/S:

Name of PROCURING ENTITY Name of CONTRACT TO BE BID IB Number DATE of Bid Opening Name of the Bidder Company Address of the Bidder Company

#### **IDENTIFY THE ENVELOPES:**

- as: > Technical Component Requirements (original copy and copy 1) > Financial Component Requirement
  - Financial Component Requirement (original, copy 1 and copy 2)

#### 16. Deadline for Submission of Bids

16.1. The Bidders shall submit on the specified date and time and either at its physical address or through online submission as indicated in paragraph 7 of the **IB**.

#### **17.** Opening and Preliminary Examination of Bids

17.1. The BAC shall open the Bids in public at the time, on the date, and at the place specified in paragraph 9 of the **IB**. The Bidders' representatives who are present shall sign a register evidencing their attendance. In case videoconferencing, webcasting or other similar technologies will be used, attendance of participants shall likewise be recorded by the BAC Secretariat.

In case the Bids cannot be opened as scheduled due to justifiable reasons, the rescheduling requirements under Section 29 of the 2016 revised IRR of RA No. 9184 shall prevail.

17.2. The preliminary examination of bids shall be governed by Section 30 of the 2016 revised IRR of RA No. 9184.

#### **18.** Domestic Preference

18.1. The Procuring Entity will grant a margin of preference for the purpose of comparison of Bids in accordance with Section 43.1.2 of the 2016 revised IRR of RA No. 9184.

#### **19. Detailed Evaluation and Comparison of Bids**

- 19.1. The Procuring BAC shall immediately conduct a detailed evaluation of all Bids rated "*passed*," using non-discretionary pass/fail criteria. The BAC shall consider the conditions in the evaluation of Bids under Section 32.2 of the 2016 revised IRR of RA No. 9184.
- 19.2. If the Project allows partial bids, bidders may submit a proposal on any of the lots or items and evaluation will be undertaken on a per lot or item basis, as the case maybe. In this case, the Bid Security as required by **ITB** Clause 15 shall be submitted for each lot or item separately.
- 19.3. The descriptions of the lots or items shall be indicated in **Section VII** (**Technical Specifications**), although the ABCs of these lots or items are indicated in the **BDS** for purposes of the NFCC computation pursuant to Section 23.4.2.6 of the 2016 revised IRR of RA No. 9184. The NFCC must be sufficient for the total of the ABCs for all the lots or items participated in by the prospective Bidder.
- 19.4. The Project shall be awarded as follows:

Option 3 - One Project having several items, which shall be awarded as separate contracts per item.

19.5. Except for bidders submitting a committed Line of Credit from a Universal or Commercial Bank in lieu of its NFCC computation, all Bids must include the NFCC computation pursuant to Section 23.4.1.4 of the 2016 revised IRR of RA No. 9184, which must be sufficient for the total of the ABCs for all the lots or items participated in by the prospective Bidder. For bidders submitting the committed Line of Credit, it must be at least equal to ten percent (10%) of the ABCs for all the lots or items participated in by the prospective Bidder.

#### 20. Post-Qualification

20.2. Within a non-extendible period of five (5) calendar days from receipt by the Bidder of the notice from the BAC that it submitted the Lowest Calculated Bid, the Bidder shall submit its latest income and business tax returns filed and paid through the BIR Electronic Filing and Payment System (eFPS) and other appropriate licenses and permits required by law and stated in the **BDS**.

#### 21. Signing of the Contract

21.1. The documents required in Section 37.2 of the 2016 revised IRR of RA No. 9184 shall form part of the Contract. Additional Contract documents are indicated in the **BDS**.

#### **CONFORME:**

Authorized Signatory Signature over printed name Contact No:

Name of Company/Firm

Company's Official Email Address (where notices will be sent)

Company's Official Contact No.



Republic of the Philippines DEPARTMENT OF HEALTH PHILIPPINE CHILDREN'S MEDICAL CENTER Bids and Awards Committee Quezon Avenue, Quezon City 1100

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## **SECTION III**

# **Bid Data Sheet**

## Supply and Delivery of Supplies/Reagents/Consumables thru Reagent Tieups / Machine Placement Agreements for Three (3) Years

IB-2022-003 to IB-2022-022

a. Supply and delivery of Various Medical and Laboratory Supplies.								
b. completed within <i>the last two (2) years</i> prior to the deadline for the submission and receipt of bids.								
quoted in Pl	hilippine							
quoted in 1	imppine							
of the follow	wing							
tem(s) joine								
r irrevocable	e letter of							
tem(s) joine	<b>ed</b> , if bid							
	TOTAL ABC (for the 1 <sup>st</sup> year)							
e Placement								
	1.254.400.00							
	1,254,400.00							
	8,100,000.00							
	867,000.00							
nt Tie-up Agr	533,610.00							
u Tie-up Agr	eement jor							
	6,308,698.00							
smissible								
	7,422,812.00							
	2,932,672.00							
	7,032,210.00							
	2,814,725.00							
lachine	7,160,546.82 1,192,250.00							
	5,457,924.00							
ıg	793,506.00							
	775,500.00							
anism	2,113,317.00							
lyzer	2,800,000.00							
	3,000,975.00							
	3,152,890.71							
yzer	1,627,387.20							
	1,597,000.00							
	3,453,071.52							
	IgM, AFP, Vitamin D alyzer							

### **Bid Data Sheet**

1	. Latest Income and Business Tax Returns filed and paid through the BIR Electronic Filing (EFPS) within the last three (3) months
2	. Valid and current Certificate of PhilGEPS Registration.
3	. Manufacturer's Certification or if the Bidder is not a manufacturer, authenticated copy of certification from the manufacturer as authorized or exclusive distributor or dealer of the products / items.
4	. Valid and current Certificate of Product Registration issued Food and Drugs Administration (FDA).
5	. With manufacturer and/or products certification by an independent 3rd party Certifying body (ISO 14020, 14021, 14024, 14025 or its equivalent), is preferred.
6	. Consumer guidelines regarding disposal of the supplies (Information about how and where the used/decommissioned products/ packaging/parts can be returned for recycling and/or disposal e.g. buy-back program)
7	. Other appropriate licenses and permits required by law and stated in the Bidding Documents
agair	re of the Bidder declared as LCB to duly submit the requirements stated above or a finding st the veracity of such shall be ground for forfeiture of the bid security and disqualify the er for award.
	Requirement Nos. 3 to 6 for items JOINED must be accomplished and submitted using ummary Sheet to be provided by PCMC.
21.2 No a	dditional contract documents relevant to the Project

#### **CONFORME:**

Authorized Signatory Signature over printed name

Contact No:

Name of Company/Firm

Company's Official Email Address (where notices will be sent) Company's Official Contact No.



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## **SECTION V**

## Special Conditions of Contract

## Supply and Delivery of Supplies/Reagents/Consumables thru Reagent Tie-ups / Machine Placement Agreements for Three (3) Years

IB-2022-003 to IB-2022-022

### **Special Conditions of Contract**

GCC Clause							
1	For Goods supplied from within the Philippines:						
	Upon delivery of the Goods to the Project Site, the Supplier shall notify the Procuring Entity and present the following documents to the Procuring Entity:						
	<ul> <li>(i) Original and four copies of the Supplier's invoice/delivery receipt showing Goods' description, quantity, unit price, and total amount;</li> </ul>						
	(ii) Four copies of Material Safety Data Sheet for a specified product upon initial delivery						
	Packaging –						
	The Supplier shall provide such packaging of the Goods as is required to prevent their damage or deterioration during transit to their final destination, as indicated in this Contract. The packaging shall be sufficient to withstand, without limitation, rough handling during transit and exposure to extreme temperatures, salt and precipitation during transit, and open storage. Packaging case size and weights shall take into consideration, where appropriate, the remoteness of the Goods' final destination and the absence of heavy handling facilities at all points in transit						
	The packaging, marking, and documentation within and outside the packages shall comply strictly with such special requirements as shall be expressly provided for in the Contract, including additional requirements, if any, specified below, and in any subsequent instructions ordered by the Procuring Entity.						
	<ol> <li>The Supplier shall have an established disposal and retrieval program or take back system for their products (items with container):</li> <li>The Supplier shall submit MSDS for a specified product.</li> <li>The product label shall bear the following information:         <ul> <li>a. Product specifications and ingredients</li> <li>b. Manufacturing and Expiration Dates</li> <li>c. Precautions</li> </ul> </li> </ol>						
	<ul> <li>d. Instructions for proper use and disposition</li> <li>e. Hazardous items shall be properly labeled as a hazardous product (e.g. flammable cytotoxic, radioactive, poison, etc.)</li> <li>4. The product shall not contain halogenated plastics and PVCs.</li> <li>5. The product shall be packed in suitable packaging materials which are reusable and recyclable.</li> </ul>						
	Transportation –						
	Where the Supplier is required under Contract to deliver the Goods CIF, CIP, or DDP, transport of the Goods to the port of destination or such other named place of destination in the Philippines, as shall be specified in this Contract, shall be arranged and paid for by the Supplier, and the cost thereof shall be included in the Contract Price.						
	Where the Supplier is required under this Contract to transport the Goods to a specified place of destination within the Philippines, defined as the Project Site, transport to such place of destination in the Philippines, including insurance and storage, as shall be specified in this Contract, shall be arranged by the Supplier, and related costs shall be included in the contract price.						

#### **CONFORME:**

Authorized Signatory Signature over printed name Contact No:

Name of Company/Firm

Company's Official Email Address (where notices will be sent) Company's Official Contact No.



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**SECTION VI** 

# Schedule of Requirements

Supply and Delivery of Supplies/Reagents/Consumables thru Reagent Tie-ups / Machine Placement Agreements for Three (3) Years

IB-2022-003 to IB-2022-022

The delivery schedule expressed as weeks/months stipulates hereafter a delivery date which is the date of delivery to the project site.

IB NUMBER	IB NUMBER ITEM DESCRIPTION					
One (1) Lot Su	pply and Delivery of supplies/consumables under Machine Placement Agreement for t	hree (3) years				
of the followin						
IB-2022-003	Apheresis kits with anticoagulant	1,254,400.00				
IB-2022-004	Blood bag, quintuple with filter	8,100,000.00				
IB-2022-005	Hemoglobin Screening for blood donors	867,000.00				
IB-2022-006	Sterile Connecting Device	533,610.00				
One (1) Lot Su	pply and Delivery of reagents/consumables under Reagent Tie-up Agreement for three	e (3) years of				
the following:						
IB-2022-007	Automated Immunohaematology Analyzer	6,308,698.00				
IB-2022-008	Automated Immunology Analyzers for Transfusion-Transmissible Infection (TTIs)	7,422,812.00				
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IB-2022-012	Flow Cytometry Machine	7,160,546.82				
IB-2022-013	Automated Blood and other Sterile Body Fluid Culture Machine	1,192,250.00				
IB-2022-014	Automated Hematology Analyzer	5,457,924.00				
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IB-2022-019	Immunoanalyzer for Procalcitonin	3,152,890.71				
IB-2022-020	Automated Urine Microscopic and Urine Chemistry Analyzer	1,627,387.20				
IB-2022-021	Immunology Analyzer for Dengue Antigen and HBsAg Confirmatory Testing	1,597,000.00				
IB-2022-022	Automated Analyzer for Methotrexate determination	3,453,071.52				

Delivery Site	PCMC Requirement DELIVERY PERIOD			
Materials Management Division, G/F PCMC, Quezon Avenue, corner Agham Road Quezon City	Within seven (7) working days from receipt of DELIVERY ORDER SLIP			

#### DELIVERY AND ACCEPTANCE

- Staggered delivery and staggered payment
- Supplies to be delivered should have an expiration at least one (1) year and longer or as expressed/required by the end-user.
- > The Supplier should submit Materials Safety Data Sheet upon initial delivery, if applicable.
- The supplier should deliver the goods called for in the Purchase Order (PO) within seven (7) working days or as stated on Delivery Period upon receipt of approved Delivery Order Slip, faxed or personally received during office hours at the Materials Management Division (MMD).
- All goods delivered pursuant to the Purchase Order (PO) with Delivery Order Slip shall be subject to acceptance and inspection by the end-user as well as by the House Inspector and of the Resident Auditor or their representatives. Goods delivered not in conformity with specifications shall be rejected and the contractor held in default.

#### **CONFORME:**

Authorized Signatory (Signature over printed name)

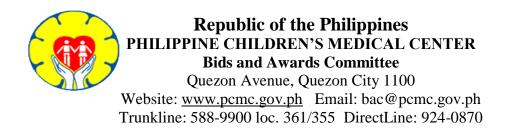
Contact No:

Name of Company/Firm

Company's Official Email Address

Company's Official Contact No.

(where notices will be sent)



### **SECTION VII**

# **Technical Specifications**

## PURCHASE OF REAGENTS UNDER MACHINE PLACEMENT AGREEMENT FOR TWO (2) YEARS

**IB Nos. IB-2020-014 to IB-2020-016** 

#### PHILIPPINE CHILDREN'S MEDICAL CENTER

#### Quezon Avenue, Quezon City

#### Invitation to Bid No. IB-2022-007

Instruction: Accomplish this form by presenting a clear statement of your offer. It should be specific, precise and complete which demonstrates how your statement complies with the required specifications. Please do not write "comply" of the page numbers of the brochure/data sheet, etc.

#### **TECHNICAL SPECIFICATIONS**

ITEM			PCMC's REQUIREMENT	BIDDER'SOFFER			
NO	QTY	UNIT	Item Description	MACHINE SPECIFICATIONS/COMPLIANCE			
		reagen	) Lot Supply and Delivery of ts/consumables under Reagent Tie-up for three rs of Automated Immunohaematology Analyzer				
2	unit	Latest Analyz	Model of Automated Immunohaematology zer				
		1.	An automated machine that can operate, process and analyze samples simultaneously from putting the specimen in the machine up to the releasing of results without human interruption for the following IH procedures such as:				
			a. Blood Grouping (newborn and adults)				
			b. Antibody Screen and Identification (patients and blood units)				
			c. Crossmatch				
			d. Other Blood Banking procedures				
		2.	Random access and capable to prioritize emergency samples with capacity to:				
			a. Detect both IgM and IgG including anti-Mi(a+) for the 3 cell antibody screen				
			b. Detect weak D and partial D				
			c. Run pediatric tubes.samples (500ul)				
		3.	Throughput should not be less than:				
			a. 48 samples for blood typing (forward & reverse) per hour				
			b. 37 samples for antibody screening (3 cells-for patients) per hour				
			c. 42 samples for antibody screening (pooled-for units) per hour				
			d. 42 samples for crossmatch per hour				
		4.	Single piercing of cards or cassettes				
		5.	With daily internal quality control system				
		6.	With back-up unit (modular machine), in which testing can be done, semi-automated with the same format. The format should be capable to resolve discrepancies either by increasing the antibody concentration or incubating in different temperatures. The system should also be able to detect with accuracy, clinically significant antibodies				

#### PHILIPPINE CHILDREN'S MEDICAL CENTER

#### Quezon Avenue, Quezon City

#### Invitation to Bid No. IB-2022-007

Instruction: Accomplish this form by presenting a clear statement of your offer. It should be specific, precise and complete which demonstrates how your statement complies with the required specifications. Please do not write "comply" of the page numbers of the brochure/data sheet, etc.

#### **TECHNICAL SPECIFICATIONS**

ITEM		1	PCMC's REQUIREMENT	BIDDER'SOFFER				
NO	QTY	UNIT	Item Description	MACHINE SPECIFICATIONS/COMPLIANCE				
		7.	Minimum volume sample required: 500uL centrifuged whole blood (in microtainer tube, without plasma separation) Dead volume sample: 400ul centrifuged whole blood (in microtainer tube, without plasma separation)					
			(in microtainer tube, without plasma separation) Dead volume of reagent: 400uL					
		8.	On-board stability of reagents: 3 to 5 days					
		9.	Automated identification of samples and reagents					
		10.	With software for antibody identification					
		11.	Capable of the following:					
			• full positive identification of lot numbers.					
			• liquid detection, sample clot detection and low level notification					
		12.	Automatic cross-checking of previous results					
		13.	Blood grouping reagents with certificate of product registration (CPR) from BFAD and evaluation result from the National Reference Laboratory for Immunohematology (NKTI)					
		14.	At least one (1) local installation of the proposed brand and model in a tertiary hospital on Metro Manila					
			Total Number of procedures/test that should be covered by the proposal	<b>Reagents/Consumables needed to complete the</b> <b>required number of tests (Use Annex ''A'')</b> (List down per Test and indicate Brand, Packing, Number of test per pack Specifications, etc.)				
	1,200	test	Blood typing, for newborn					
	24,400	test	Blood typing, for adults					
	12,000	test	Antibody screening, for patients (3 cells)					
	9,600	test	Antibody screening, for units (pooled)					
	200	test	Antibody identification					
	14,400	test	Crossmatch					
	Coomb's	Test:	1					
	12,000	test	Direct Coomb's Test					
	800	test	Indirect Coomb's Test					

#### **Additional Requirements**

> Machine to be offered should have passed end-user's evaluation

#### PHILIPPINE CHILDREN'S MEDICAL CENTER

#### Quezon Avenue, Quezon City

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#### **TECHNICAL SPECIFICATIONS**

ITEM			PCMC's REQUIREMENT	<b>BIDDER'SOFFER</b>
	QTY	UNIT	Item Description	MACHINE SPECIFICATIONS/COMPLIANCE

PCMC has the right to reject any or all bids without offering any reason, waive any required formality and award the contract to any bidder whose proposals as evaluated by PCMC is the most advantageous to the government.

#### **AUTHORIZED REPRESENTATIVE'S INFOS:**

Signature over Printed Name

**COMPANY'S DETAILS:** 

NAME OF COMPANY

ADDRESS

COMPANY'S OFFICIAL EMAIL ADDRESS

**Designation/Position** 

TEL. NO. / FAX NO.



Republic of the Philippines DEPARTMENT OF HEALTH PHILIPPINE CHILDREN'S MEDICAL CENTER Bids and Awards Committee Quezon Avenue, Quezon City 1100 website: www.pcmc.gov.ph\_\_email: pcmcbac@gmail.com

website: <u>www.pcmc.gov.ph</u> email: pcmcbac@gmail.com Trunkline: 8588-9900 local 361/355 Telefax No.: 8924-0870

## **SECTION VIII**

# Checklist of Technical and Financial Documents

Supply and Delivery of Supplies/Reagents/Consumables thru Reagent Tie-ups / Machine Placement Agreements for Three (3) Years

**IB-2022-003 to IB-2022-022** 

### **Checklist of Technical and Financial Documents**

The Bidder shall submit the following <u>TECHNICAL COMPONENT ENVELOPE</u> (ARRANGED, <u>NUMBERED AND TABBED</u>) [Strictly NO using of staple wire and thick materials for tabs] as enumerated below:

#### I. TECHNICAL COMPONENT ENVELOPE

#### Class "A" Documents

#### Legal Documents

1. Valid PhilGEPS Registration Certification (Platinum Membership) and its Annex A.

<u>or</u>

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,

and

- 3. Mayor's/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. and
- 4. Valid Tax Clearance per Executive Order 398, series of 2005, as finally reviewed and approved by the BIR.

#### Note:

In the event the bidder opted to submit only requirement nos. 2 to 4 Legal Documents, the Certificate of PhilGEPS Registration (Platinum Membership) shall remain a post-qualification requirement to be submitted in accordance with Section 34.2 of the 2016 Revised IRR of RA 9184 (Pursuant to GPPB Circular 07-2017 dated 31 July 2017)

#### Technical Documents

- 5. 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. (Use of Form No. DOBA-PCMC-SCF3b is required)
- 6. Statement of the bidder's Single Largest Completed Contract (SLCC) similar to the contract to be bid (*Refer to BDS Clause 5.3*), except under conditions provided for in Sections 23.4.1.3 and 23.4.4.4 of the 2016 Revised IRR of RA 9184, within **two (2) years** prior to bid opening (use of Form No. DOBA-PCMC-SCF3a is required).
- 7. Bid Security (**Refer to BDS Clause 14.1**)
- 8. Duly accomplished and signed Production/ Delivery Schedule using the form as provided for in Section VI
- 9. Duly accomplished and signed Technical Specification using the form as provided for in Section VII
- 10. Omnibus Sworn Statement (Use of the Form provided is required)

- 11. Certificate of Performance in letterhead of their clients indicating the contact numbers and email addresses signed by the authorized head of the Department from three (3) clients of the bidder issued within the last six (6) months prior to bid opening.
  - *Note:* Certification issued by PCMC Materials Management Division (MMD) must be included if bidder had done business with us. Certification of which should be of same category (e.g. equipment/supplies) of project being bided.
- 12. Valid and current License to Operate (LTO) issued by Food and Drug Administration (FDA).

#### Financial Documents

- 1. The prospective bidder's Audited Financial Statements, showing, among others, the prospective bidder's total current assets and liabilities, stamped "received" by the BIR or its duly accredited and authorized institutions, for the preceding calendar year which should not be earlier than two (2) years from the date of submission.
- 2. The prospective bidder's computation of the Net Financial Contracting Capacity (NFCC) that must be at least equal to the ABC to be bid (*Use of Form No. DOBA–PCMC–NFF4 is required*); OR.

a committed Line of Credit from a Universal or Commercial Bank, in lieu of its NFCC computation, it must be at least equal to 10% of the ABC.

#### Class "B" Documents

1. For Goods, valid joint venture agreement (JVA), in case the joint venture is already in existence. In the absence of a JVA, duly notarized statements from all the potential joint venture partners should be included in the bid, stating that they will enter into and abide by the provisions of the JVA in the event that the bid is successful. Failure to enter into a joint venture in the event of a contract award shall be ground for the forfeiture of the bid security. (*Use of Form No. DOBA-PCMC-JVF6 is required*).

Each partner of the joint venture shall submit their legal eligibility documents or Certificate of PhilGEPS Registration (Platinum Membership). The submission of technical and financial eligibility documents by any of the joint venture partners shall constitute compliance: Provided, That the partner responsible to submit the NFCC shall likewise submit the Statement of all its ongoing contracts and Audited Financial Statements.

The Bidder shall submit the following **FINANCIAL COMPONENT ENVELOPE** (ARRANGED, **NUMBERED AND TABBED**) [Strictly NO using of staple wire and thick materials for tabs] as enumerated below:

#### II. FINANCIAL COMPONENT ENVELOPE

- 1. Duly accomplished and signed Bid Form
- 2. Duly accomplished and signed **Price Schedule** using the form as provided

#### *Note:* Bidder <u>shall return</u> to PCMC the issued <u>USB Flash Drive</u> <u>containing</u> the <u>soft copy</u> of their <u>accomplished</u> <u>TECHNICAL</u> <u>SPECIFICATIONS</u> <u>AND</u> <u>PRICE</u> <u>SCHEDULE</u> (in <u>EXCEL FORMAT</u>)

- 3. With manufacturer and/or products certification by an independent 3rd party Certifying body (ISO 14020, 14021, 14024, 14025 or its equivalent)
- 4. Duly notarized Certificate from Principal Manufacturer/ Distributor that the machine being offered has been/ currently installed to at least one (1) tertiary hospital in Metro Manila (attach supporting documents)

- 5. Section II. Instructions to Bidders with signature (conforme) on all pages
- 6. Section III. Bid Data Sheet with signature (conforme) on all pages
- 7. Section IV. General Conditions of the Contract with signature (conforme) on all pages
- 8. Section V. Special Conditions of the Contract with signature (conforme) on all pages
- 9. Signed Conforme on the Terms of Reference, if applicable
- 10. Certification for Assurance of Stocks Availability [use of Form No. DOBA-PCMC-CAF10 is required]
- 11. Return Policy [use of Form No. DOBA PCMC CRF34 is required]
- 12. Brochure for the machine to be provided under the agreement

#### **CONFORME:**

Authorized Signatory Signature over printed name Contact No:

Name of Company/Firm

Company's Official Email Address (where notices will be sent) Company's Official Contact No.

#### PHILIPPINE CHILDREN'S MEDICAL CENTER Quezon Avenue, Quezon City

#### **TERMS OF REFERENCE**

#### 1. DESCRIPTION OF THE PROJECT

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One (1) Lot Reagent Tie-up Agreement for Three (3) Years on Supply and Delivery of Reagents and Consumables with Installation, Commissioning and free use of one (1) brand new latest model, and one (1) back- up with same specifications (can be refurbished or brand new) Automated Immunohaematology Analyzer with one (1) unit of semi automated machine as back-up.

#### 2. OBLIGATIONS OF THE SUPPLIER

#### 2.1 Installation and Commissioning of the Machine and Peripherals

- 2.1.1 Install, test and commission two (2) latest model of Automated Immunohaematology Analyzer with one (1) unit of semi automated machine as back-up conforming to the required technical specifications including electrical safety of the machine such as uninterruptible compatible power supply at Philippine Children's Medical Center (PCMC) - Pathology Division within the required period.
- 2.1.2 Responsible for every installation step required to set up the machines including manpower, supplies, and materials for all civil works necessary to complete the installation at its own expense. The PCMC will only provide the tapping points for electricity and waste disposal as needed by the system.
- 2.1.3 Pay all related expenses and requirements on LIS connectivity.
- 2.1.4 Provide (2) high-end printers including consumables which can produce an average volume print-outs of 10,000 pages per month for 24/7 operation.
- 2.1.5 Upgrade the software as necessary free of charge within the contract period.
- 2.1.6 Specify the conditions of delivery, such as packing and necessary environmental requirements (such as data loggers/temperature monitoring) upon delivery.
- 2.1.7 Be responsible for any loss or damage to the machine not caused by PCMC's negligence.

#### 2.2 Supply and Delivery of Reagents and Consumables

- 2.2.1 Supply and deliver the reagents and consumables within seven (7) calendar days after receipt of approved Delivery Order Slip (DOS) with a shelf life of at least one (1) year from date of delivery. Quantity may increase or decrease depending on patients' census.
- 2.2.2 Guarantee that the quantity to be provided is sufficient to cover the required number of tests. In case not sufficient, additional quantity shall be provided at no cost to PCMC.

Projected number of IH procedures/tests f	or 2022:
Blood typing, for newborn	1,200
Blood typing, for adults	24,400
Antibody screening, for patients (3 cells)	12,000
Antibody screening, for units (pooled)	9,600
Antibody identification	200
Crossmatch	14,400
Coomb's Test:	
Direct Coomb's Test	12,000
Indirect Coomb's Test	800

- 2.2.3 Guarantee that there will be no price adjustment during the duration of the contract.
- 2.2.4 Replace defective product and other consumables including arising from errors or malfunction of the machine and its accessories free of charge within seven (7) calendar days from receipt of notice which may be through email or SMS.
- 2.2.5 Ensure availability of the reagents and consumables equivalent to three (3)-month inventory stored off-site to be used in conjunction with immunohaematology analyzer with no minimum quantity per month. Quantity may increase or decrease depending on patient census.
- 2.2.6 Provide phosphate buffer solution for performing immunohaematology tests.
- 2.2.7 Provide additional antibody identification panel cells.
- 2.2.8 Ensure immunohaematology results adhere to currently accepted standards at all times following strict standard operating procedures.

#### 2.3 Warranty/Service and Maintenance

2.3.1 Guarantee the serviceable lifespan of the automated immunohaematology analyzer is at least three (3) years after the acceptance test.

- 2.3.2 Bind itself to service and maintain the equipment in good working condition, free of charge, during the duration of the contract. This warranty includes availability of spare parts and quarterly calibration/preventive maintenance with corresponding certificate/service report and sticker.
- 2.3.3 Guarantee that, in the event of machine malfunction, the service response time via phone call is within thirty (30) minutes and the Field Service Engineer renders service promptly within three (3) hours after notification twenty four (24) hours a day seven (7) days a week, including holidays.
- 2.3.4 Provide back-up unit of the same model within twenty four (24) hours if both machines malfunction for a maximum period of eight (8) hours.

#### 2.4 Training and Technical Service Support

- 2.4.1 Provide End-user's orientation and training on the proper operation, minor troubleshooting and maintenance of the equipment by the Principal Manufacturer /Distributor's certified trainer within seven (7) days upon installation until the End-users can operate the equipment confidently and shall issue a Certificate of Training to all participants.
- 2.4.2 Provide training updates and quality assurance-related lectures, training, and workshop to personnel when necessary free of charge.

#### 2.5 Green Procurement

2.5.1 Observe and comply with PCMC's policy on green procurement while contract is enforced.

#### 3. DURATION OF THE CONTRACT

- 3.1 This Agreement shall be valid for three (3) years or until December 31, 2024 whichever comes earlier.
- 3.2 Staggered delivery and staggered payment, subject to government accounting and auditing, rules and regulations.
- 3.3 Both parties shall not assign or transfer any portion of this Agreement nor any item of the machine without prior written consent of the other.

#### 4. OTHER PROVISIONS

- 4.1 PCMC shall not be liable for any failure to perform under this machine placement agreement due to *force majeure*, strikes (legal or illegal), lockouts, fires, flood or water damage, riots, governmental acts or orders, interruption of transportation, inability to obtain materials upon reasonable prices or terms, or any other cause beyond its control.
- 4.2 In the event that either party fails to perform under, or commits, or allows to be committed, a breach of any to covenants and conditions of this Agreement and other provisions contained in the bidding documents, the other party shall notify such party in writing of such failure or default. The breaching party shall then have the right to remedy such failure or default within thirty (30) days from receipt of such notice, otherwise, the other party may terminate this Agreement immediately upon notice of its failure to remedy such breach.
- 4.3 Should the parties be constrained to resort to court action, the losing party shall pay the prevailing party an amount equivalent to twenty percent (20%) of the total amount claimed, and as by way of attorney's fees but not less than Twenty Thousand Pesos (20,000.00). Venue of such shall be laid exclusively in Quezon City, Metro Manila.

#### **CONFORME:**

Authorized Signatory Signature over printed name

Name of Company/Firm

### PRICE SCHEDULE

PROCURING ENTITY : PHILIPPINE CHILDREN'S MEDICAL CENTER						NAME OF BII	DDER :			
NAME OF PROJECT : One (1) Lot Supply and Delivery of reagents/consumables under Reagent Tie-up for three (3) years of Automated Immunohaematology Analyzer			INVITATION TO BID NO. IB 2022-007							
PCMC REQUIREMENT						<b>BIDDER'S OFFE</b>	BIDDER'S OFFER			
1	2	3	4	5	6	7	8	9	10	11
ITEM DESCRIPTION	Brand/Packaging of Reagents / Consumables to be provided	Qty	Manufacturer	Country of Origin	UNIT PRICE EXW	COST OF LOCAL LABOR, RAW MATERIAL AND COMPONENT	TOTAL PRICE EXW	UNIT PRICES FINAL DESTINATION AND UNIT PRICE OF OTHER INCIDENTAL SERVICES	SALES AND OTHER TAXES PAYABLE PER ITEM IF CONTRACT IS AWARDED	TOTAL PRICE DELIVERED FINAL DESTINATION [(col. 9 + 10) x 2]
One (1) Lot Supply and Delivery of reagents/consumables under Reagent Tie-up for three (3) years of Automated Immunohaematology Analyzer ** Total ABC = Php 6,308,698.00 (for the first year)										
** Use Annex''A'' for the det	ailed offer **									
Additional Requirements										
• Conforme to the Terms of Refe	erence.									
<ul> <li>Supplies to be delivered should have expiration of at least one (1) year and longer or as expressed/ required by Pathology.</li> <li>The price of the bided item(s) shall be valid for three (3) years or within the contract duration The quantities specified are estimated requirements during the period and may be decreased depending upon th</li> <li>actual need of PCMC. It is understood therefore that PCMC is not bound to order / purchase all the items / quantities called for on this bidding</li> </ul>					pending upon the	TERMS OF (For discounts b	PAYMENT eing offered, if there	e's any. Otherwise	e, state "NONE")	:
• The supplier should submit Ma	• The supplier should submit Materials Safety Data Sheet upon delivery, if applicable.									
• PCMC has the right to reject any or all bids without offering any reason, waive any required formality and award the contract to any bidder whose proposals as evaluated by PCMC is the most advantageous to the government.						NAM	E AND SIGNATUR	E OF AUTHORIZ	LED REPRESEN	TATIVE