Republic of the Philippines
DEPARTMENT OF HEALTH
PHILIPPINE CHILDREN'S MEDICAL CENTER
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BID BULLETIN NO. BB-2025-013

Date : **November 13, 2024**

TO : **ALL PROSPECTIVE BIDDERS**

FROM : **BIDS AND AWARDS COMMITTEE**

SUBJECT : **AMENDMENT/ADDENDUM/CLARIFICATION AS STATED**

Name of Project : **One (1) Lot Supply and Delivery of reagents/consumables under Reagent Tie-up Agreement for three (3) years for Flow Cytometry**
Invitation to Bid No. IB-2025-034
Total ABC = Php 7,160,319.30

This Bid Bulletin No. **BB-2025-013** outlines the amendments and additional requirements in the bidding documents for above project as follows:

I. AMENDMENTS

A. Under Section VII. Technical Specifications

ITEM DESCRIPTION	
FROM	**TO**
One (1) unit Flow cytometry system, fully automated machine	One (1) unit Flow cytometry system, brand new automated machine
1. With three (3) multi-colored (blue, red, violet)	1. With three (3) multi-colored lasers (blue, red, violet)
5. Panels must follow the Euroflow Consortium guidelines and protocols for leukemia and lymphoma immunophenotyping	5. Panels must follow the Euroflow Consortium guidelines and protocols for MRD, leukemia and lymphoma immunophenotyping. Panels must be validated and readily available.
7. Instrument should be capable of full automation of tests (end to end automation)	7. Instrument should be capable of sample acquisition and analysis with predefined templates and reports

B. Under Terms of Reference (TOR)

ITEM DESCRIPTION	
FROM	**TO**
2.1 Installation and Commissioning of the machine peripherals	**2.1 Installation and Commissioning of the machine peripherals**
2.1.3 Install the Flow Cytometer and all its accessories as well as provide PCMC with a start up kit of reagents and calibrate all the necessary tests to be run by the laboratory FREE OF CHARGE	2.1.3 At least 20-30 samples/tests should be provided for the start up kit of reagents to be provided by the supplier

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II. ADDITIONAL REQUIREMENT

Under Terms of Reference (TOR)

2.2 Supply and Delivery of Reagents and Consumables

2.2.9 The winning bidder shall provide all necessary reagents and supplies required for the equipment validation, which will be conducted by the PCMC Pathology Division upon delivery of the machine. The validation studies must include 20 tests for each of the following criteria:

> Linearity
> Precision (within run and between run)
> Accuracy (within run and between run)
> Carryover
> Limit of blank
> Lower and higher limit of quantification
> Contamination
> Matrix of interferences
> Comparison bias
> Reference range

For information and guidance of all concerned.

FRANCIS S. DELA CUESTA, RN, MAN
Chairperson, Bids and Awards Committee