Republic of the Philippines
DEPARTMENT OF HEALTH
PHILIPPINE CHILDREN'S MEDICAL CENTER
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BID BULLETIN NO. BB-2025-012

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 Hema Analyzer** **Invitation to Bid No. IB-2025-017** **Total ABC = Php 2,295,540.00**

This Bid Bulletin No. **BB-2025-012** outlines the amendments and additional requirements in the bidding documents for above project as follows:

I. AMENDMENTS

Under Section VII. Technical Specifications

ITEM DESCRIPTION	
FROM	**TO**
1. With the capability to be connected to an automated slide maker with a throughput of at least 25 slides per hour with where can be personalized based on patients hematocrit count.	1. With the capability to be connected to an automated slide maker with a throughput of at least 25 slides per hour
3. With sample aspiration volume of not more than 100 uL for whole blood sample and not more than 80 uL for pre-diluted samples to allow samples to be re-analyzed several times.	3. With sample aspiration volume of not more than 165 uL for whole blood sample and pre-diluted samples
4. Must be capable of accomodating primary tubes with sample collected as low as 350 uL for both whole body and body fluis, and as low as 200 uL collected in a micro collection tube without causing errors relating to insufficient blood samples to prevent the need for recollection for low volume samples.	4. Must be capable of accomodating primary tubes with sample collected as low as 200 uL for both whole body and body fluids
7. Capable of analyzing the Cerebrospinal fluid (CSF), Peritoneal Fluid, Pleural Fluid and Synovial Fluid. Must be able to provide proof of US FDA clearance for the bofy fluids analysis	7. Must be able to provide FDA clearance for the body fluids analysis

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II. ADDITIONAL REQUIREMENT

Under Terms of Reference (TOR)

2.2 Supply and Delivery of Reagents and Consumables

2.2.8 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