



Republic of the Philippines
DEPARTMENT OF HEALTH
Center for Health Development Davao Region
SOUTHERN PHILIPPINES MEDICAL CENTER
J.P. Laurel Avenue, Davao City



BID BULLETIN NO. 1
(September 6, 2016)

**Consignment of Chemistry Machine and Immunology Machine and Delivery of
its Corresponding Reagents and Consumables**

Bid No: 2016-08-0336

This BID BULLETIN No. 1 is issued to modify or revise some requirements in the Checklist of Requirements and Bid Data Sheet, and some items in the Technical Specifications of the abovementioned project. This modification shall prevail and form an integral part of the Bidding Documents.

a. Checklist of Requirements and Bid Data Sheet:

- The required "*Certification that the equipment being offered must be of the latest version or model and has no record of violations such as electrical or mechanical malfunction since the time of its production*" maybe issued by the bidder in the absence or in lieu of the manufacturer/principal.
- The **Certificate of Product Registration (CPR)** is a mandatory requirement per **Republic Act 9711** and **DOH Department Order 2015-0005**. Therefore, the CPR must be part of the Technical Requirements and must be submitted during the scheduled submission of bids and opening of bids. *In case that the In vitro Diagnostic Kits does not require the said certificate, the bidder must attach a **Certificate of Exemption issued by FDA**.*

b. Clinical Chemistry Analyzer Technical Specifications:

- In page 2 under the column for the item, there were some corrections in the description, to wit:
 - Analyzers of Clinical Chemistry must be identical that is capable of performing and processing both arrays of tests.
 - Analyzers of Immunology must be identical that is capable of performing and processing both arrays of tests.
- In page 3, item 7, delete *additional tests other than* and the specification will become **"Must be equipped with built-in ISE system that is capable of processing electrolytes"**.
- The following requirements stated in the Other Terms and Conditions in page 5 item 1.1 to 1.2 and in page 6 item 6.1 to 6.2 will be submitted during **post-qualification** by the bidder declared as the Lowest Calculated and Responsive Bid:
 - 1.1 With Certificate of Date Manufactured and date released from the manufacturing company.
 - 1.2 Certificate of calibration and material safety data sheet.
 - 6.1 operating manual of each machine/analyzer;
 - 6.2 Product inserts for each test parameter;

- In page 7, some corrections were made for item 9 and the requirement becomes *"List of installations in two (2) tertiary hospital level 3 (within the past three (3) years) ISO certified or its equivalent with a satisfactory performance with supporting documents issued by the consignee/vendee in the form of Certificate of Acceptance reflecting the following:*

9.1 Machine performance (poor, good, excellent)

9.2 Availability of reagents and consumables (good, excellent)

9.3 After sales service (poor, good, excellent)

c. Immunoassay Analyzer Technical Specifications:

- Delete item 18 in page 19 *"Must be capable of close-capped tube sample run to prevent airborne/aerosol infection transmissions (e.g. merscov, ebola, other emerging and re-emerging diseases)"*.
- The following requirements stated in the Other Terms and Conditions in page 19 item 1.1 to 1.2 and in page 20 item 6.1 to 6.2 will be submitted during **post-qualification** by the bidder declared as the Lowest Calculated and Responsive Bid:
 - 1.1 With Certificate of Date Manufactured and date released from the manufacturing company.
 - 1.2 Certificate of calibration and material safety data sheet.
 - 6.1 operating manual of each machine/analyzer;
 - 6.2 Product inserts for each test parameter;

All other provision of the bidding documents which are not affected shall remain in effect.

For guidance and information of all concerned.


RICARDO S.D. JUSTOL, MARE, MPA
Chairperson, Bids and Awards Committee