



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

(December 14, 2016)

Consignment of One (1) Brand New Unit Fully Automated Immunohematology Analyzer
including Delivery & Supply of its Reagents, & Consumables
Bid No: 2016-11-0699

This BID BULLETIN No. 1 is issued to modify or revise some Eligibility and Technical Requirements indicated in the *Checklist of Requirements, Bid Data Sheet (BDS)* 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"* must be issued by the principal.
- The 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 shall be supported by the **Purchase Order or Contract Agreement or NOA** stating the amount of the project;

b. Technical Specifications:

- In page 2, under the General Description, delete ID and it must be "Fully Automated Immuno Haematology Analyser that employs the **Microtyping System or Micro Column Systems or Micro Column Agglutination Systems** for Cross Matching, Blood Grouping, Antibody Screening.
- In page 3, item 7 letter c, delete gel and the item becomes **"Cards/cassettes"**.
- In page 4, under the Other Terms and Conditions for Acceptability item 2, the required *"With Certificate of Date Manufactured and date released from the manufacturing company"* will be submitted during the installation of the machine.
- In page 6 item 9, delete the word **gel**, and the requirement becomes *"A monthly report of the number of test run will be made by the end user and shall be noted by the supplier base on the number of cassettes/cards / consumed for blood grouping and tubes/columns for compatibility testing as reflected on the daily worksheet per shift"*.

- In page 8 item 16, the *Operating Manual of the machine/analyzer in English* (letter a) and *Product inserts for each test parameter* (letter b) will be submitted during the installation of the machine. While the letter c *"Valid Certificate of Product Registration or Certificate of Exemption for the reagents issued by BFAD to the specific supplier/bidder"* still to be submitted during the submission of Eligibility and Technical Requirements.
- In page 10 item 23, include the Biomedical Technician and the requirement becomes *"The end-user, Biomedical Technician and Department Chairman or his representative must have an actual inspection and or witness an actual installation and commissioning of the machine being offered"*.
- In page 10, item 23, letter b (End User shall evaluate the functionality of the machine/analyzer's advantages), this requirement will be conducted during the evaluation demonstration process;

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

For guidance and information of all concerned.


RICARDO SD. JUSTOL, MARE, MPA
Chairperson, Bids and Awards Committee