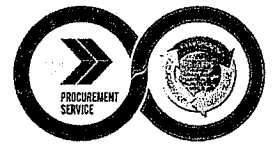




Republic of the Philippines
 Department of Budget and Management
PROCUREMENT SERVICE
BIDS AND AWARDS COMMITTEE



Bid Bulletin No. 2

November 19, 2018

Public Bidding 18-360-3

**Supply and Delivery of Cartridge-Based Nucleic Acid Amplification Test
 for the Department of Health (DOH)**

Issued pursuant to Sec. 22.5 of the IRR of R.A. 9184 to clarify, include and/or amend certain provisions in the Bidding Documents issued for this project.

Please be informed that the activities for the above-mentioned project shall be adjusted as follows:

A. AMENDMENT

REFERENCE	BASES FOR CLARIFICATION/ AMENDMENT/INCLUSION						
SECTION III. BID DATA SHEET							
Page 44 BDS Clause 29.2 (9) xxx <table border="1" style="width: 100%;"> <tr> <td style="width: 10%;">29.2</td> <td>xxx</td> </tr> <tr> <td></td> <td>9. Copy of the World Health Organization (WHO) prequalification listing in the website. <u>Supplier must provide a copy of World Health Organization (WHO) Guidelines that product offered is in the Implementation Manual as a diagnostic test.</u></td> </tr> <tr> <td></td> <td>xxx</td> </tr> </table> xxx	29.2	xxx		9. Copy of the World Health Organization (WHO) prequalification listing in the website. <u>Supplier must provide a copy of World Health Organization (WHO) Guidelines that product offered is in the Implementation Manual as a diagnostic test.</u>		xxx	To amend the additional requirement to be submitted during Pos-Qualification in the Bid Data Sheet.
29.2	xxx						
	9. Copy of the World Health Organization (WHO) prequalification listing in the website. <u>Supplier must provide a copy of World Health Organization (WHO) Guidelines that product offered is in the Implementation Manual as a diagnostic test.</u>						
	xxx						
SECTION VII. TECHNICAL SPECIFICATIONS							

For the purpose of this Bulletin and for better understanding of its contents, the following rules shall apply: (a) ~~double-strikeout~~ – denotes deletion; (b) Underline – denotes inclusion or new item/requirement; and “xxx” – denotes separation of phase/s being amended from the rest of the main text.

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Supply and Delivery of Cartridge Based Nucleic Acid Amplification Test for the Department of Health (DOH)



xxx

TECHNICAL SPECIFICATIONS

LOT NO.1 : Cartridge-Based Nucleic Acid Amplification Test

QUANTITY : 506 kits

AGENCY SPECIFICATIONS	BIDDER'S STATEMENT OF COMPLIANCE
CARTRIDGE-BASED NUCLEIC ACID AMPLIFICATION TEST	Brand:
xxx	xxx
<ul style="list-style-type: none"> World Health Organization (WHO) prequalification listing in the website: Copy of World Health Organization (WHO) Guidelines that product offered is in the Implementation Manual as a diagnostic test. 	
xxx	xxx

xxx

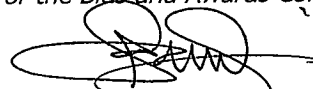
To amend the requirement in the Technical Specifications

Please refer to revised Technical Specifications labeled as Appendix "A".

All other related provisions in the Bidding Documents affected by these amendments are likewise deemed amended to conform to this Bid Bulletin.

Amendments/inclusions/clarifications made herein shall be considered an integral part of the Bidding Documents.

For the Bids and Awards Committee III:



ENGR. EDWARD R. SADDI

Chairperson
Bids and Awards Committee III

Technical Specifications

LOT NO.1 : Cartridge-Based Nucleic Acid Amplification Test
 QUANTITY : 506 kits

AGENCY SPECIFICATIONS	BIDDER'S STATEMENT OF COMPLIANCE
CARTRIDGE-BASED NUCLEIC ACID AMPLIFICATION TEST	Brand:
<ul style="list-style-type: none"> • Nucleic acid amplification cartridge-based test that can rapidly detects Myobacterium Tuberculosis (MTB) and Rifampicin resistance in less than 2 hours (50s in kit) • Cartridge should be compatible with existing machine at health facilities that can detect Myobacterium Tuberculosis (MTB) and Rifampicin resistance in less than 80 minutes and/or not more than 2 hours • A copy of World Health Organization (WHO) Guidelines that product offered is in the Implementation Manual as diagnostic test. 	
SHELF LIFE	
<ul style="list-style-type: none"> • Supplies must be fresh commercial stock with a total shelf life of at least fifteen (15) months from the date of delivery 	
PACKAGING INSTRUCTIONS	
<ul style="list-style-type: none"> • With one (1) sample reagent bottle per cartridge, with one (1) piece of 2mL pipette per cartridge, with one (1) CD-ROM per kit. • Fifty (50) cartridges in a kit. Number of kit per carton will follow standard packaging of Manufacturer. 	
REQUIRED DOCUMENTS	
<ul style="list-style-type: none"> • License to Operate (LTO) issued by the PFDA • Certification from the Manufacturer that the Bidder is Exclusive/ Authorized Distributor or Dealer of the products, or a Distributorship/ Dealership Agreement from the Manufacturer or Manufacturer's Agent or Main Distributor or Dealer in the Philippines. If the Bidder is the Manufacturer, Certificate that the Bidder manufactures the products/items. • Declaration of conformity with the appropriate ISO/IEC/PNS standard issued to the Manufacturer by an ISO certifying body or any appropriate agency or body. • Original brochure or downloaded from the internet or product insert. 	

LABELLING INSTRUCTIONS <ul style="list-style-type: none"> Each box and kits should be imprinted with: "Philippine Government Property – Department of Health – NOT FOR SALE" 	
Manufacturing Date: _____ Expiration Date: _____ Batch / Lot No.: _____	
DELIVERY SCHEDULE <ul style="list-style-type: none"> 506 kits – Sixty (60) calendar days upon receipt of Notice to Proceed (NTP) 	
DELIVERY SITE <ul style="list-style-type: none"> DOH Different Warehouse/s or Service Provider/s within Metro Manila 1. QMMC, Project 4, Quezon City; 2. POPCOM Warehouse, Welfare Ville Compound, Mandaluyong City; 3. Non-Phareil Warehouse, Temperatue Control, Sun Valley, Paranaque City (aircon); 4. Veterans, Taguig City (Non-aircon); 5. Ximex Warehouse, Tanyag, Taguig City; 6. Philpost, Port Area, Manila; 7. DOH Central Office Bldg. 25, Tayuman , Sta. Cruz, Manila 	

I hereby certify that the statement of compliance to the foregoing technical specifications are true and correct, otherwise, if found to be false either during bid evaluation or post-qualification, the same shall give rise to automatic disqualification of our bid.

Name of Company	Signature Over Printed Name of Authorized Representative	Date
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