Republic of the Philippines
Department of Health
METRO MANILA CENTER FOR HEALTH DEVELOPMENT

SUPPLEMENTAL/ BID BULLETIN NO. 1

IB NO. 2024-004
Procurement of Syphilis Rapid Test (30/tests/kits) (SHORT OF AWARD)

This Supplemental/Bid Bulletin No. 1 is being issued to revise provisions/specifications in the Bidding Documents for a fore cited project:

Revision and clarification to provisions/specifications in the Bidding Documents:	
ORIGINALTECHNICAL SPECIFICATIONS	**AMENDED**
Sensitivity: 99.5% vs TPHA back up with documents	Sensitivity: At least 99.5% vs TPHA back up with documents
Specificity: 99.5% vs TPHA back up with documents	Specificity: At least 99.5% vs TPHA back up with documents
Packaging Instructions: Primary packaging: 100 pcs. per box	Primary packaging: 100 pcs per box or Manufacturer Standard

Bidders are advised to use the following attached forms and submit together with all required documents for the submission of bids on November 8, 2023, 9:00 AM

This Supplemental/Bid Bulletin No. 1 shall form part of the Bidding Documents. Any provisions in the Bidding Documents inconsistent herewith is hereby amended, modified and superseded accordingly.

For guidance and information of all concerned.

Issued this 28th day of October, 2023 in MMCHD.

SGD.
PRETCHELL P. TOLENTINO, MD, MCHM
Director III / BAC Chairperson

Section VII. Technical Specifications

<table>
<tr><td colspan="4">Republic of the Philippines
Department of Health
Metro Manila Center for Health Development

TECHNICAL SPECIFICATIONS</td></tr>
<tr><td>Item No.</td><td>Procurement of Syphilis Rapid Test (30/tests/kits) (SHORT OF AWARD)</td><td>Qty./Unit</td><td>3,386 kits</td></tr>
<tr><td colspan="3">Name of Manufacturer:</td><td>Country of Origin</td></tr>
<tr><td colspan="3">Brand:</td><td>Model: (if applicable)</td></tr>
<tr><td colspan="3">ABC: 8,465,000.00</td><td></td></tr>
<tr><td colspan="3">PURCHASER’S SPECIFICATION</td><td>STATEMENT OF COMPLIANCE</td></tr>
<tr><td colspan="3">SPECIFICATIONS:
Principle solid based Immunochromatographic Assay
Detect all isotypes (Ig, IgM and IgA)
Specimen – Serum (10 uL) Plasma (10 uL), Whole blood (20 uL)
Sensitivity: At least 99.5% vs TPHA back up with documents
Specificity: At least 99.5% vs TPHA back up with documents
Materials provided: Capillary pipettes, alcohol swabs and lancets
Result time: 5-20 minutes
Can be stored at room temperature
With DOH/FDA CPR and NRL/SACCL completed evaluation
With 5 Clinical Studies from 5 independent bodies concerning
Sensitivity & specificity of the brand test kits
With comparative study of the brand to be offered versus RPR
from independent body
Provide quick guide usage (at least 50 pcs) and CD training
Guide (20 pcs)</td><td></td></tr>
</table>

Shelf Life: Must be fresh commercial stock, with a total shelf life of 24 months from the date of manufacture but not less than 18 months from the date of delivery **Packaging Instructions:** Primary packaging: 100 pcs. per box or Manufacturer Standard Standard packaging of the manufacturers as approved by the Philippine Food Drug and Authority **Labelling Instructions:** Standard labelling instruction as approved by FDA pursuant to Administrative Order No. 2016-0008 In addition to the labelling requirements of the PFDA: a. On each blister pack/foil strip and box, the following should be legibly imprinted or stickered using a permanent non-removable sticker/label that is binding and will leave residue and ripping if removed. b. On each small and bigger box/carton, the following should be legibly imprinted or stickered with non-removable or permanent sticker or label that is binding and will leave residue and ripping if removed Philippine Government Property- Department of Health NOT FOR SALE Date of Manufacturer: Date of Expiry: Batch/Lot No. Delivery period: 30 Calendar days Area of delivery: DOH-MMCHD Pasig Warehouse	
<u>ADDITIONAL REQUIREMENTS TO BE SUBMITTED BY THE SINGLE/LOWEST CALCULATED BIDDER (SCB/LCB) AS PART OF POST QUALIFICATION:</u> 1. One (1) original sample of manufacturer's product to be submitted and returned after evaluation. The sample submitted and approved during the evaluation shall be the same item to be delivered upon contract award. The prototype of the labeling instruction must be part of the sample submitted; however, the technical specifications of the	

labeling instruction of the product must be complied with upon delivery.	