



Republic of the Philippines  
Department of Health

**METRO MANILA CENTER FOR HEALTH DEVELOPMENT**

**SUPPLEMENTAL/ BID BULLETIN NO. 1**

**IB 2023 – 008**

**PROCUREMENT OF SYPHILLIS RAPID TEST (30 TESTS/KIT) (SHORT OF AWARD)**

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

<b>Revision and clarification to provisions/specifications in the Bidding Documents:</b>	
<b>FROM</b>	<b>TO</b>
With DOH/FDA CPR and NRL/SACCL completed evaluation	With DOH/FDA CPR and NRL-SACCL completed evaluation
<b>Packaging Instructions:</b> 1.Primary packaging: 100 pcs. per box	<b>Packaging Instructions:</b> 1.Primary packaging: 100 kits. per box

Bidders are advised to use the following attached forms and submit together with all required documents for the submission of bids on 28<sup>th</sup> day of November 2022, 9:30 AM:

This Supplemental/Bid Bulletin No. 1 shall form an integral part of the Bidding Documents. All other provisions indicated in the bidding documents which are not affected by this Supplemental/Bid Bulletin No. 1 shall remain in effect.

For guidance and information of all concerned.

Issued this 18<sup>th</sup> day of November 2022 in MMCHD

Approved by:

**ALELI ANNIE GRACE P. SUDIACAL, MD, MPH**  
Director III / BAC Chairperson



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

Republic of the Philippines Department of Health Metro Manila Center for Health Development			
<b>TECHNICAL SPECIFICATIONS</b>			
Item No. 1	<b>PROCUREMENT OF SYPHILLIS RAPID TEST (30 TESTS/KIT) (SHORT OF AWARD)</b>	Qty./Unit	2,434 KITS
Name of Manufacturer:		Country of Origin	
Brand:		Model: (if applicable)	
<b>ABC: P 6,085,000.00</b>			
PURCHASER'S SPECIFICATION		STATEMENT OF COMPLIANCE	
<p><b><u>SPECIFICATIONS:</u></b></p> <p>Principle solid based Immunochromatographic Assay            Detect all isotypes (Ig, IgM and IgA)            Specimen – Serum (10 uL) Plasma (10 uL), Whole blood (20 uL)            Sensitivity: 99.5% vs TPHA back up with documents            Specificity: 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 &amp; 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)</p> <p><b>Shelf Life:</b></p> <p>Must be fresh commercial stock, with a total shelf life of 24 months from the date of manufacture but not less than 18</p>			



## METRO MANILA CENTER FOR HEALTH DEVELOPMENT

months from the date of delivery

**Packaging Instructions:**

1. Primary packaging: 100 kits per box
2. 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 Manufacture: \_\_\_\_\_

Date of Expiry: \_\_\_\_\_

Batch/Lot No.: \_\_\_\_\_

Delivery period: 30 Calendar days

Area of delivery: DOH-MMCHD Pasig Warehouse

**ADDITIONAL REQUIREMENT TO BE SUBMITTED  
BY THE SINGLE/LOWEST CALCULATED BIDDER  
(SCB/LCB) AS PART OF POST QUALIFICATION:**



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## METRO MANILA CENTER FOR HEALTH DEVELOPMENT

<p>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 award of contract. Prototype of the labelling instruction must be part of the sample submitted however, the technical specifications of the labelling instruction of the product must be complied upon delivery.</p>	
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