Republic of the Philippines
Department of Health

METRO MANILA CENTER FOR HEALTH DEVELOPMENT

SUPPLEMENTAL/ BID BULLETIN NO. 1

IB NO. 2024-007
Procurement of 900 kits Dengue NS1 AG + IgG/IgM kit (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**
SPECIFICATIONS: DENGUE NS1 AG + IgG/IgM RAPID DIAGNOSTIC TEST	
900 Kits	9,000 tests in 900 boxes
RDT kit for the qualitative detection of dengue non-structural protein 1 (NS1) and dengue Ns1 IgG and IgM antibodies from all four (4) Dengue serotypes	RDT kit for the qualitative detection of dengue non-structural protein 1 (NS1) and dengue NS1 IgG and IgM antibodies from all four (4) Dengue serotypes
3mL–5mL assay buffer	3 mL – 5 mL assay buffer

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 Any provisions in the Bidding Documents inconsistent herewith is hereby amended, modified and superseded accordingly.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 900 kits Dengue NS1 AG + IgG/IgM kit (SHORT OF AWARD)</td><td>Qty./Unit</td><td>900 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: 3,600,000.00</td><td></td></tr>
<tr><td colspan="3">PURCHASER’S SPECIFICATION</td><td>STATEMENT OF COMPLIANCE</td></tr>
<tr><td colspan="3">SPECIFICATIONS:
DENGUE NS1 AG + IgG/IgM RAPID DIAGNOSTIC TEST
9,000 tests in 900 boxes
Standard Features:
RDT kit for the qualitative detection of dengue non-structural protein 1 (NS1) and dengue NS1 IgG and IgM antibodies from all four (4) Dengue serotypes
*NS1 & IgG/IgM Combo in Immunochromatography in plastic cassette format
*3 mL – 5 mL assay buffer
*10 uL capillary pipette
*1 disposable dropper
*1 lancet, ultra thin, 28 gauge, consistent depth penetration, universal design firs almost all lancing devices</td><td></td></tr>
</table>

*assay time 15-20 min *Sensitivity: ≥ 92% and Specificity ≥ 98% Dengue NS1 Ag based on 3rd party evaluation or based on publication by national Reference Laboratory (NRL), Centers for Disease Control and Prevention (CDC) or World Health Organization (WHO) *Sensitivity ≥ 94% and Specificity Dengue IgG/IgM ≥ 96% Dengue IgG/IgM based on 3rd party evaluation or based on publication by national Reference Laboratory (NRL), Centers for Disease Control and Prevention (CDC) or World Health Organization (WHO) ***Packaging Instruction:** Manufacturer's standard/Distributor Packaging **Shelf life:** The product must have a minimum shelf life of eighteen (18) months remaining at the time of delivery. **Labelling Instructions:** On each box, the following should be legibly imprinted or stickered: Philippine Government Property – Department of Health NOT FOR SALE Manufacturing Date:_____ Expiration Date:_____ Batch/ Lot No.:_____ Delivery Period: Sixty (60) calendar days after receipt of notice to proceed Delivery site: DOH-MMCHD Tala Warehouse *Delivery Time: 60 days after receipt of PO Other Requirement: CERTIFICATE OF PRODUCT ANALYSIS (CPA) ISSUED BY THE MANUFACTURER	
ADDITIONAL REQUIREMENTS TO BE SUBMITTED BY THE SINGLE/LOWEST CALCULATED BIDDER (SCB/LCB) AS PART OF POST QUALIFICATION: 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.	