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

SUPPLEMENTAL/ BID BULLETIN NO. 1

IB 2022 – 059
PROCUREMENT OF VARIOUS HOSPITAL SUPPLIES (LOT BIDDING)

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

Revision and clarification to provisions/specifications in the Bidding Documents:	
Item No. 1 PROCUREMENT OF 9,165 BOXES OF DISPOSABLE INJECTION NEEDLE GAUGE 23 Compatible with Luer slip tip 1 cc syringe	
FROM	**TO**
Item No. 2 PROCUREMENT OF 5,000 BOXES TUBERCULIN (1 CC) SYRINGE Disposable-Single use only	Disposable
Quantity: 9,164 boxes (100 pcs per box)	Quantity: 5,000 boxes (100 pcs per box)
Item No. 3 PROCUREMENT OF 920 BOXES THREE (3) CC SYRINGE Disposable-Single use only	Disposable
Item No. 4 PROCUREMENT OF 633 BOXES 0.5 ML AD SYRINGE 0.5 ml AD Syringe with needle for intramuscular injection	0.5 ml AD Syringe with needle for intramuscular injection or Needle retractable safety syringe
Disposable-Single use only	Disposable
Item No. 5 PROCUREMENT OF 22,450 PCS OF 0.3 ML AD SYRINGE Quantity: 22,450 BOXES	Quantity: 22,450 BOXES
0.3 ml AD Syringe with needle for intramuscular injection	0.3 ml AD Syringe with needle for intramuscular injection or Needle retractable safety syringe
Disposable-Single use only	Disposable

Bidders are advised to use the following attached forms and submit together with all required documents for the submission of bids on 25th day of April 2022, 9:00 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 16th day of April 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

Technical Specifications

Republic of the Philippines Department of Health Metro Manila Center for Health Development TECHNICAL SPECIFICATIONS			
Item No. 1	**PROCUREMENT OF 9,165 BOXES OF DISPOSABLE INJECTION NEEDLE GAUGE 23**	Qty./Unit	9,165 BOXES
Name of Manufacturer:		Country of Origin	
Brand:		Model: (if applicable)	
ABC: **P 1,374,750.00**			
PURCHASER'S SPECIFICATION		STATEMENT OF COMPLIANCE	
A. Technical Specifications: Gauge: 23 Length: 1 inch Disposable-Single use only Compatible with Luer slip tip 1 cc syringe Quantity: 9,165 boxes (100 pcs per box)			
B. Upon delivery the following must be complied: **Shelf life:** Must be fresh commercial stock, with a total shelf life 24 months from the date of manufacture but not less than 18 months from the date of delivery **Packaging Instructions:** 1. Primary packaging: 100 pcs per box 2. Standard packaging of the manufactures as approved by the Philippine Food Drug and Authority **Labelling Instructions:** Standard packaging of the manufacturers 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 of removed b. On each small and bigger box/carton, the following should			

be legible imprinted or stickered with non-removable or permanent sticker or label that is binding and will residue and ripping if removed Philippine Government Property-Department of Health **NOT FOR SALE** Date Manufacture: _____ Date of Expiry: _____ Batch/Lot No._____ **Delivery Period:** 30 Calendar days **Place of Delivery:** DOH-MMCHD Pasig Warehouse	
C. <u>Additional Requirement to be submitted by the Single/Lowest Calculated Bidder (SCB/LCB) as part of the 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 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.	
D. <u>Additional Requirements to be attached with this form arranged, numbered and tabbed as enumerated below:</u> 1. Valid Certificate of Product Registration (CPR) issued by Philippine Food and Drug Administration (PFDA) or valid extension The CPR must be valid for the entire period of the award. If the CPR is about to expire, the supplier must have submitted a copy of an application of renewal to the FDA at least 3 months before the expiry date (**a copy of the expiring CPR which is stamped with an "Extension of Validity" shall be submitted as proof**); [2. Valid and current License to Operate (LTO) as Medical Device Importer/ Wholesaler issued by PFDA. Provided that in case of expired LTO, the application for renewal was made timely as per PFDA Circular No. 2011-004. **In case of expired LTO, the following copies may be submitted:** a. Expired LTO; b. Application for renewal; and c. Official Receipt as proof of payment of renewal of LTO	

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Republic of the Philippines Department of Health Metro Manila Center for Health Development TECHNICAL SPECIFICATIONS			
Item No. 2	**PROCUREMENT OF 5,000 BOXES TUBERCULIN (1 CC) SYRINGE**	Qty./Unit	5,000 BOXES
Name of Manufacturer:		Country of Origin	
Brand:		Model: (if applicable)	
ABC: **P 1,500,000.00**			
PURCHASER'S SPECIFICATION		STATEMENT OF COMPLIANCE	
A. <u>Technical Specifications:</u> Syringe Volume: 1 cc/1 ml Graduation Scale: 0.01 ml Syringe Tip Type: Luer Slip Tip With plunger molded to the luer cone (Low Dead Volume) Detachable Needle (23 to 27 x 5/8 inch / 1 inch / 1 ½ inch) Disposable Quantity: 5,000 boxes (100 pcs per box)			
A. <u>Upon delivery the following must be complied:</u> **Shelf life:** Must be fresh commercial stock, with a total shelf life 24 months from the date of manufacture but not less than 18 months from the date of delivery **Packaging Instructions:** 1.Primary packaging: 100 pcs per box 2.Standard packaging of the manufactures as approved by the Philippine Food Drug and Authority **Labelling Instructions:** Standard packaging of the manufacturers 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 of removed b. On each small and bigger box/carton, the following should be legible imprinted or stickered with non-removable or permanent sticker or label that is binding and will residue and ripping if removed Philippine Government Property-Department of Health **NOT FOR SALE** Date Manufacture: _____ Date of Expiry: _____ Batch/Lot No._____ **Delivery Period:** 30 Calendar days **Place of Delivery:** DOH-MMCHD Pasig Warehouse	
B. <u>Additional Requirement to be submitted by the Single/Lowest Calculated Bidder (SCB/LCB) as part of the 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 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.	
D. <u>Additional Requirements to be attached with this form arranged, numbered and tabbed as enumerated below:</u> 1. Valid Certificate of Product Registration (CPR) issued by Philippine Food and Drug Administration (PFDA) or valid extension The CPR must be valid for the entire period of the award. If the CPR is about to expire, the supplier must have submitted a copy of an application of renewal to the FDA at least 3 months before the expiry date (**a copy of the expiring CPR which is stamped with an "Extension of Validity" shall be submitted as proof**); [3. Valid and current License to Operate (LTO) as Medical Device Importer/ Wholesaler issued by PFDA. Provided that in case of expired LTO, the application for renewal was made timely as per PFDA Circular No. 2011-004. **In case of expired LTO, the following copies may be submitted:** a. Expired LTO; b. Application for renewal; and c. Official Receipt as proof of payment of renewal of LTO	

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Republic of the Philippines Department of Health Metro Manila Center for Health Development TECHNICAL SPECIFICATIONS			
Item No. 3	**PROCUREMENT OF 920 BOXES THREE (3) CC SYRINGE**	Qty./Unit	920 BOXES
Name of Manufacturer:		Country of Origin	
Brand:		Model: (if applicable)	
ABC: **P 276,000.00**			
PURCHASER'S SPECIFICATION		STATEMENT OF COMPLIANCE	
A. <u>Technical Specifications:</u> Syringe Volume: 3 cc/3 ml Graduation Scale: 0.1 ml Syringe Tip Type: Luer Lock Tip Detachable Needle (23 G x 1 inch) Disposable Quantity: 920 boxes (100 pcs per box)			
B. <u>Upon delivery the following must be complied:</u> **Shelf life:** Must be fresh commercial stock, with a total shelf life 24 months from the date of manufacture but not less than 18 months from the date of delivery **Packaging Instructions:** 1. Primary packaging: 100 pcs per box 2. Standard packaging of the manufactures as approved by the Philippine Food Drug and Authority **Labelling Instructions:** Standard packaging of the manufacturers 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 of removed			

b. On each small and bigger box/carton, the following should be legible imprinted or stickered with non-removable or permanent sticker or label that is binding and will residue and ripping if removed Philippine Government Property-Department of Health **NOT FOR SALE** Date Manufacture: _____ Date of Expiry: _____ Batch/Lot No._____ **Delivery Period:** 30 Calendar days **Place of Delivery:** DOH-MMCHD Pasig Warehouse	
C. <u>Additional Requirement to be submitted by the Single/Lowest Calculated Bidder (SCB/LCB) as part of the 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 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.	
D. <u>Additional Requirements to be attached with this form arranged, numbered and tabbed as enumerated below:</u> 1. Valid Certificate of Product Registration (CPR) issued by Philippine Food and Drug Administration (PFDA) or valid extension The CPR must be valid for the entire period of the award. If the CPR is about to expire, the supplier must have submitted a copy of an application of renewal to the FDA at least 3 months before the expiry date (**a copy of the expiring CPR which is stamped with an "Extension of Validity" shall be submitted as proof**); [3. Valid and current License to Operate (LTO) as Medical Device Importer/ Wholesaler issued by PFDA. Provided that in case of expired LTO, the application for renewal was made timely as per PFDA Circular No. 2011-004. **In case of expired LTO, the following copies may be submitted:** a. Expired LTO; b. Application for renewal; and c. Official Receipt as proof of payment of renewal of LTO	

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Item No. 4	**PROCUREMENT OF 633 BOXES 0.5 ML AD SYRINGE**	Qty./Unit	633 BOXES
Name of Manufacturer:		Country of Origin	
Brand:		Model: (if applicable)	
ABC: **P 126,600.00**			
PURCHASER'S SPECIFICATION		STATEMENT OF COMPLIANCE	

A. Technical Specifications:

0.5 ml AD Syringe with needle for intramuscular injection or Needle retractable safety syringe

Gauge 23, 1"in length with protective cap

Polyprophylene syringe material

Quantity: 100 pcs per box

Disposable

B. Upon delivery the following must be complied:

Shelf life: 36 months to not less than 24 months from the date of manufacture

Each small box should be legibly imprinted or stickered with non-removable or permanent sticker/label that is binding and will leave residue and ripping of removed:

Philippine Government Property-Department of Health

NOT FOR SALE

Date Manufacture: _____

Date of Expiry: _____

Batch/Lot No. _____

Delivery Period: 30 - 45 Calendar days

Place of Delivery: DOH-MMCHD Pasig Warehouse

C. <u>Additional Requirement to be submitted by the Single/Lowest Calculated Bidder (SCB/LCB) as part of the 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 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.	
A. <u>Additional Requirements to be attached with this form arranged, numbered and tabbed as enumerated below:</u> 1. Valid Certificate of Product Registration (CPR) issued by Philippine Food and Drug Administration (PFDA) or valid extension The CPR must be valid for the entire period of the award. If the CPR is about to expire, the supplier must have submitted a copy of an application of renewal to the FDA at least 3 months before the expiry date (**a copy of the expiring CPR which is stamped with an "Extension of Validity" shall be submitted as proof**); [2. Valid and current License to Operate (LTO) as Medical Device Importer/ Wholesaler issued by PFDA. Provided that in case of expired LTO, the application for renewal was made timely as per PFDA Circular No. 2011-004. **In case of expired LTO, the following copies may be submitted:** d. Expired LTO; e. Application for renewal; and Official Receipt as proof of payment of renewal of LTO	

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Item No. 5	**PROCUREMENT OF 22,450 PCS OF 0.3 ML AD SYRINGE**	Qty./Unit	22,450 PIECES
Name of Manufacturer:		Country of Origin	
Brand:		Model: (if applicable)	
ABC: **P 224,500.00**			
PURCHASER'S SPECIFICATION		STATEMENT OF COMPLIANCE	
A. <u>Technical Specifications:</u> 0.3 ml AD Syringe with needle for intramuscular injection or Needle retractable safety syringe Gauge 23, 1"in length with protective cap Polyprophylene syringe material Quantity: 100 pcs per box Disposable			
B. <u>Upon delivery the following must be complied:</u> **Shelf life:** 36 months to not less than 24 months from the date of manufacture Each small box should be legibly imprinted or stickered with non-removable or permanent sticker/label that is binding and will leave residue and ripping of removed: Philippine Government Property-Department of Health **NOT FOR SALE** Date Manufacture: _____ Date of Expiry: _____ Batch/Lot No._____ **Delivery Period:** 30 - 45 Calendar days **Place of Delivery:** DOH-MMCHD Pasig Warehouse			

C. <u>Additional Requirement to be submitted by the Single/Lowest Calculated Bidder (SCB/LCB) as part of the 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 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.	
D. <u>Additional Requirements to be attached with this form arranged, numbered and tabbed as enumerated below:</u> 3. Valid Certificate of Product Registration (CPR) issued by Philippine Food and Drug Administration (PFDA) or valid extension The CPR must be valid for the entire period of the award. If the CPR is about to expire, the supplier must have submitted a copy of an application of renewal to the FDA at least 3 months before the expiry date (**a copy of the expiring CPR which is stamped with an "Extension of Validity" shall be submitted as proof**); [4. Valid and current License to Operate (LTO) as Medical Device Importer/ Wholesaler issued by PFDA. Provided that in case of expired LTO, the application for renewal was made timely as per PFDA Circular No. 2011-004. **In case of expired LTO, the following copies may be submitted:** f. Expired LTO; g. Application for renewal; and Official Receipt as proof of payment of renewal of LTO	