



METRO MANILA CENTER FOR HEALTH DEVELOPMENT

SUPPLEMENTAL/ BID BULLETIN NO. 1

IB 2022 – 127E

PROCUREMENT OF VARIOUS MEDICAL DEVICE (LOT BIDDING) (REBID)

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:

Additional Requirement 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 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.

FROM	TO
Submission of Bids: SEPTEMBER 22, 2022	Submission of Bids: SEPTEMBER 21, 2022
ITEM NO. 1: 10 UNITS INFUSION PUMP	
Display: At least 4 inches, LCD with big font	Display: At least 3.5 inches, LCD with big font
• Micro Infusion Speed: 0.1ml/h up to 1500 ml/h	• Micro Infusion Speed: 0.1ml/h up to 1500 ml/h - RETAIN
ITEM NO. 2: 9 UNITS SYRINGE PUMP	
Suitable for syringe with the maximum range	• Suitable for syringe with the maximum range (50-60 ml)
Capable of working independently or together in a stacking system	Capable of working independently or together in a stacking system - RETAIN
Mechanical clutch design	Mechanical clutch design - REMOVED


Bidders are advised to use the following attached forms and submit together with all required documents for the submission of bids on 21st day of September 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 14th day of September 2022 in MMCHD

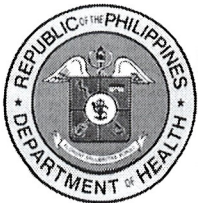
Approved by:


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 TECHNICAL SPECIFICATIONS			
Item No. 1	INFUSION PUMP	Qty./Unit	10 Units
Name of Manufacturer:			Country of Origin
Brand:			Model: (if applicable)
ABC: 850,000.00			
PURCHASER'S SPECIFICATION			STATEMENT OF COMPLIANCE
Technical Specifications: <ul style="list-style-type: none"> • Horizontal Door Clip • Display: At least 3.5 inches, LCD with big font • Ergonomics operation interface • Modes <ul style="list-style-type: none"> - Rate Mode - Drop Mode - Time Mode • Micro Infusion Speed: 0.1ml/h up to 1500 ml/h • Infusion Accuracy: $< \pm 3\%$ • With software adjusting function suitable for any brand of infusion sets • With high precision pressure sensor • With built-in drip sensor • Capable of storing up to 800 infusion records and configuration memory • Power Supply: Built-in battery (manufacturer's standard), can last up to 8 hours of operation if fully charged REQUIREMENTS IF AWARDED THE CONTRACT: 1. Completion Period: The delivery, installation, testing and commissioning of the equipment and its accessories, including the training of end-users and maintenance staff must be completed within 30 calendar days upon receipt of Notice to Proceed. 2. Testing: Prior to acceptance, the end user shall conduct a physical inspection and functionality test. The equipment must be functioning and must have no physical damage and defect.			



METRO MANILA CENTER FOR HEALTH DEVELOPMENT

3. **Training:** The supplier shall provide training on the proper use and maintenance of the equipment to the end-users and to the hospital maintenance staff within 3 days upon delivery of the equipment.

4. **Warranty:** Warranty certificate for two (2) years on parts and service. The supplier shall either repair or replace any item or part in the equipment that is found to be defective in material or in workmanship under normal use. The warranty period shall commence from the date of acceptance by the end-user after testing and commissioning.

5. Notarized undertaking that the supplier shall conduct the necessary corrective maintenance within five (5) calendar days upon notification of the equipment breakdown from the end-user. The undertaking shall include a statement that the number of days where the equipment is unusable due to defective material or workmanship, shall be added to the warranty period.

6. **Manuals:** The supplier must provide the end-user one (1) hard and one (1) soft copy of the following:

- a) Service manual in English language
- b) Operation manual in English language

7. 'With "**DOH-MMCHD HFEP**"(Government Property not for sale) sticker in each unit

[illegible]

Source of Fund: SAA 2021-0209 (HFEP 2021 CONAP)

Recipient: Novaliches District Hospital

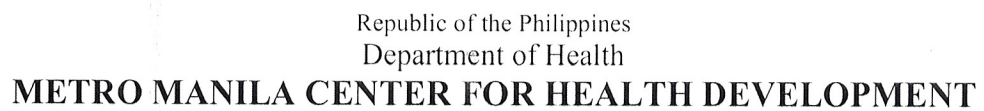
Additional Requirement to be submitted by the Single/Lowest Calculated Bidder (SCB/LCB) as part of post qualification:

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 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.



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 TECHNICAL SPECIFICATIONS			
Item No. 2	SYRINGE PUMP	Qty./Unit	9 Units
Name of Manufacturer:			Country of Origin
Brand:			Model: (if applicable)
ABC: 585,000.00			
PURCHASER'S SPECIFICATION			STATEMENT OF COMPLIANCE
TECHNICAL SPECIFICATIONS: <ul style="list-style-type: none"> Flow rate: 0.012-300 mL/min Suitable for syringe with the maximum range (50-60 ml) Capable of working independently or together in a stacking system Compatible with all types of syringes Hang clamp design 			
REQUIREMENTS, if awarded the contract <ol style="list-style-type: none"> 1. Completion Period: The delivery, installation, testing and commissioning of the equipment and its accessories, including the training of end-users and maintenance staff must be completed within 30 calendar days upon receipt of Notice to Proceed. 2. Testing: Prior to acceptance, the end user shall conduct a physical inspection and functionality test. The equipment must be functioning and must have no physical damage and defect. 3. Training: The supplier shall provide training on the proper use and maintenance of the equipment to the end-users and to the hospital maintenance staff within 3 days upon delivery of the equipment. 4. Warranty: Warranty certificate for two (2) years on parts and service. The supplier shall either repair or replace any item or part in the equipment that is found to be defective in material or in workmanship under normal use. The warranty period shall commence from the date of acceptance by the end-user after testing and commissioning. 5. Notarized undertaking that the supplier shall conduct the necessary corrective maintenance within five (5) calendar days upon notification of the equipment breakdown from the end-user. The undertaking shall include a statement that the number of days where the equipment is unusable due to defective material or workmanship, shall be added to the warranty period. 6. Manuals: The supplier must provide the end-user one (1) hard and one (1) soft copy 			



- a) Service manual in English language
- b) Operation manual in English language

[illegible]

Recipient: Novaliches District Hospital

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.