

# Republic of the Philippines DEPARTMENT OF HEALTH Metro Manila Center for Health Development



# SUPPLEMENTAL/ BID BULLETIN NO. 1 <u>IB2024 - 066E</u> <u>PROCUREMENT OF INFUSION PUMP ADAPTIVE KVO FUNCTION</u>

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

<b>Technical Specifications</b>	Query	Response of the End User Unit	
- 0.10-30 ml/h (minimum increment: 0.01 ml/h)	0.10-3 ml/h or better (minimum increment: 0.01 ml/h)	0.10-3 ml/h or better (minimum increment: 0.01 ml/h) GRANTED	
Supports maximum of 30 profiles and 5000 drugs	Supports maximum of 30 profiles (patient) and 5000 drugs – for clarification with end user	Supports maximum of 30 profiles (patient) and 5000 drugs	
Programmable drug library information and pump configuration	Programmable drug library information and pump configuration – for clarification with end user	Programmable drug library information and pump configuration – Retained the original specs	

Bidders are advised to use the following attached forms and submit them together with all required documents for the submission of bids on the 7<sup>th</sup> day of August 2024, 9:00 AM:

This Supplemental/Bid Bulletin No. 1 shall be integral to the Bidding Documents. All other provisions indicated in the bidding documents not affected by this Supplemental/Bid Bulletin No. 1 shall remain in effect.

For guidance and information of all concerned.

Issued this 31st day of July 2024 in MMCHD

Approved by:

JEREMIAS FRANCIS Y. CHAN, MD Licensing Officer V / BAC Chairperson



# Republic of the Philippines DEPARTMENT OF HEALTH Metro Manila Center for Health Development



## SUPPLEMENTAL/ BID BULLETIN NO. 1 <u>IB2024 – 066E</u> PROCUREMENT OF INFUSION PUMP ADAPTIVE KVO FUNCTION

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

1. Query during Pre-bidding Conference:					
Query	Response of the End User Unit				
0.10-3 ml/h or better (minimum increment: 0.01 ml/h)	0.10-3 ml/h or better (minimum increment: 0.01 ml/h) GRANTED				
Supports maximum of 30 profiles (patient) and 5000 drugs – for clarification with end user	Supports maximum of 30 profiles (patient) and 5000 drugs				
Programmable drug library information and pump configuration – for clarification with end user	Programmable drug library information and pump configuration – Retained the original specs				
	Query  0.10-3 ml/h or better (minimum increment: 0.01 ml/h)  Supports maximum of 30 profiles (patient) and 5000 drugs – for clarification with end user  Programmable drug library information and pump configuration – for clarification				

Bidders are advised to use the following attached forms and submit them together with all required documents for the submission of bids on the  $7^{th}$  day of August 2024, 9:00 AM:

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### Republic of the Philippines Department of Health Metro Manila Center for Health Development

### **TECHNICAL SPECIFICATIONS**

	TECHNICAL SPECIFICAT	IONS	
Item No. 1	INFUSION PUMP WITH ADAPTIVE KVO FUNCTION	Qty./Unit	26 Units
Name of Manufacturer:			Country of Origin
Brand:			Model: (if applicable)
ABC: 2,600,00	00.00		
PURCHASER'S SPECIFICATION			STATEMENT OF COMPLIANCE
<b>Technical Spe</b>			
	Manufacturer's Standard		
	touchscreen with adjustable brightness		
	ode: Rate mode, Time mode, Weight mode, Seq		
	e, Micro mode, Loading dose mode and Drip mo	de	
	e range: 0.1-2000.0 ml/h (20d/ml IV set)		
	nl/h (60d/ml IV set)		
	crement of infusion rate:		
	nl/h (minimum increment: 0.01ml/h)		
	ml/h (minimum increment: 0.1ml/h)		
	00 ml/h (minimum increment: 1ml/h)		
• VTBI range	9ml (minimum increment: 0.01ml)		
• Infusion inac			
• KVO rate:	curacy. ±370		
	nl/h (minimum increment: 0.01 ml/h)		
	KVO included, can adjust KVO rate according to		
rate	, ,		
	or better (minimum increment: 0.01 ml/h)		
	occlusion: 150~975 mmHg, 12 levels are availa		
• Fluid side od	cclusion: Fluid side occlusion alarm is supported		
<ul> <li>Single Bubbl</li> </ul>			
	le alarm accuracy: ±15μl or ±20% (whichever is		
	evel: 25, 50, 100, 200, 300, 500, and 800 (µl)		
	& acoustic alarm	- C : ( : + l .	
	sfusion certified, can be used for blood trans	siusion (with	
transfusion se			
	recent therapies are recorded and can be used e can be changed anytime during infusion		
	infusion among multiple pumps		
	g library with dose error reduction system		
	aximum of 30 profiles (patient) and 5000 drugs		
	ble drug library information and pump configur	ation	
0	unction available, unintended bolus ≤ 0.2ml		
	nds again in 2 minutes if there is still alarm after	being muted	
	000 events can be stored for review		
•	er is cut off, the infusion automatically switch	h to internal	
battery power	r		

- Power: 220V, 60Hz
- Battery duration: 10 hours or better
- Charging time: 4 hours or less

### **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 60 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 a 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:

- a) 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 workmanship under normal use. The warranty period shall commence from the date of acceptance by the end-user after testing and commissioning.
- b) Preventive maintenance at least every six (6) months or according to the manufacturer's recommendations;
- c) Corrective maintenance within five (5) calendar days upon notification from the end-user regarding equipment breakdown/defects.
- d) The number of days where the equipment is unusable due to equipment defects/faults shall be added to the warranty period.
- e) The supplier shall specify post-warranty comprehensive preventive maintenance costs including list and prices of major spare parts of the equipment for three (3) years after the warranty period.
- 5. **Notarized undertaking** that the supplier shall conduct the necessary corrective maintenance, replacements and repair 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.

Source of Fund: SAA 2024-02-001002 (HFEP 2024)

Recipient: Taguig-Pateros District Hospital

### Republic of the Philippines Department of Health Metro Manila Center for Health Development

TECHNICAL SPECIFICATIONS					
Item No. 1	INFUSION PUMP WITH ADAPTIVE KVO FUNCTION	Qty./Unit	26 Units		
Name of Manufacturer:			Country of Origin		
Brand:			Model: (if applicable)		
ABC: <b>2,600,000</b>					
	PURCHASER'S SPECIFICATION		STATEMENT OF COMPLIANCE		
<ul> <li>Display: LCD to</li> <li>Infusion Mode</li> <li>Trapezia mode,</li> <li>Infusion rate r  - 0.1-667ml/</li> <li>Minimum incr  - 0.10-99 ml  - 1000-2000</li> <li>VTBI range  - 0.1~9999n</li> <li>Infusion inacco</li> <li>KVO rate:  - 0.10-30 ml  - Adaptive K</li> </ul>	ifications: Idanufacturer's Standard Souchscreen with adjustable brightness E: Rate mode, Time mode, Weight mode, Sec Micro mode, Loading dose mode and Drip mo ange: 0.1-2000.0 ml/h (20d/ml IV set) I/h (60d/ml IV set) I/h (minimum increment: 0.01ml/h) I/h (minimum increment: 0.1ml/h) I/h (minimum increment: 1ml/h) I/h (minimum increment: 1ml/h) I/h (minimum increment: 0.01ml)	de			
<ul> <li>Fluid side occl</li> <li>Single Bubble <ul> <li>Air bubble</li> <li>Bubble Lev</li> </ul> </li> <li>With visual &amp;</li> <li>Blood transfettransfusion sets</li> <li>At least 20 rec</li> <li>Infusion rate of</li> <li>Continuous in</li> <li>Built-in drug l</li> <li>Supports maxif</li> <li>Programmabl</li> <li>Anti-Bolus fur</li> <li>Alarms sound</li> <li>Maximum 300</li> </ul>	alarm accuracy: ±15μl or ±20% (whichever is rel: 25, 50, 100, 200, 300, 500, and 800 (μl) acoustic alarm asion certified, can be used for blood tran	greater) sfusion (with ation being muted			

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- d) The number of days where the equipment is unusable due to equipment defects/faults shall be added to the warranty period.
- e) The supplier shall specify post-warranty comprehensive preventive maintenance costs including list and prices of major spare parts of the equipment for three (3) years after the warranty period.
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