The Health Sciences Center

BIDS & AWARDS COMMITTEE 1 (BAC 1)

 Proj. Ref. No.:
 PUR21-10-0976
 Opening of Bids: 17 June 2022

 End-User:
 Dept. of Anesthesia
 ABC: Php12,000,000.00

Project: SUPPLY, DELIVERY AND TESTING OF FOUR (4) UNITS OF

ANESTHESIA MACHINE WITH MULTIPARAMETER

PATIENT MONITOR

Contract: Single Bid

Item No.	Qty.	UOM	Item Description	Unit Price		ations included)
NO.					in figures	in words
1	4	Unit	Anesthesia Machine with Multiparameter Patient Monitor	3,000,000.00		
			Components:			
			A. Anesthesia Machine with Vaporizers B. Multiparameter Cardiac Monitor C. Transport Monitor			
			Anesthesia Machine Technical Specifications			
			 Must have Three Gas System (02, Med. Air and N2O) Must have dual tubes (Macro and Micro) for each gas; Min. oxygen flow for micro must be 50ml or below With separate auxiliary outlet of oxygen with own flow meter for nasal cannula/face mask use Must have auxiliary common gas outlet for non-rebreathing system (NRBS) Can provide nominal 21% concentration of oxygen in O2/N2O mixture (hypoxia guard proportioning system) Must have at least two (2) Vaporizer Mounts: One (1) Isoflurane and One (1) Sevoflurane vaporizer compatible with the machine. Must be equipped with standard pin index yoke for gases (for oxygen only); May have yoke for N2O also. Must have reusable breathing circuit natural latex-free and autoclavable at 134° for up to 10mins. or settings prescribed by manufacturer Breathing system must be fully integrated in the workstation 			

	Approved by:
	Dean BIENVENIDO S. BALOTRO, RPh, DBA, MS Chairperson
(Signature over Printed Name of President / Gen. Manager)	onun porson
(Name & Address of Company)	

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	10. One step bag-vent switch turns ventilator on/off 11. Adjustable pressure limiting valve range 1 to 75 cmH2O and with tactile indicator 12. Circuit volume of 2.6 to 2.7L maximum including canister capable of low-flow anesthesia 13. No tools needed for assembly/disassembly of breathing system 14. Quick-change CO2 absorber with water tap (CO2 cannister, 1500G or lower) 15. Must have active gas scavenging system 16. Fresh Gas Decoupling or equivalent feature. 17. One bellows for all patient range (neonate to adult) 18. Must be equipped with gas pressure gauges (pipeline & cylinder) 19. Must be equipped with oxygen flush valve 20. Re-usable breathing head corrugated tubings must have universal adaptors/ coupling 21. High-pressure tubings /adapter/		
	feature. 17. One bellows for all patient range (neonate to adult) 18. Must be equipped with gas pressure gauges (pipeline & cylinder) 19. Must be equipped with oxygen flush valve 20. Re-usable breathing head corrugated tubings must have universal adaptors/ coupling		
	connector/ coupling for pipeline gases: Machine side: DISS; Gas pipeline outlet side: Medstar/ Oxequip TM type or DISS 22. Medical grade electrical outlets with circuit breaker fuse in anesthesia machine base unit 23. Anesthesia Machine Base Unitmanufacturer's Standard for equipment model but must include the following: a. With Lockable Caster Wheels b. With drawers c. With Monitor Mount		

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Chairperson

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d.	Pneumatics		
24. Venti	ator Specifications		
a.	Suitable for Adult, Pediatric and		
	Neonates		
b.	Operating Modes:		
	i. Volume controlled ventilation		
	ii. Pressure Controlled Ventilation		
	iii. Pressure Support		
	iv. Synchronized Intermittent		
	Mandatory Ventilation		
	v. Manual Ventilation		
	vi. Spontaneous Breathing		
C.	Monitored Parameters:		
	i. Expired Volume		
	ii. Expired Flow		
	iii. Respiratory Rate		
	iv. Airway Pressure with Pressure		
	waveform display		
	v. Allows Alarm Management		
d.	Control Input Ranges:		
	i. Breathing Frequency (rate) 4		
	to 60 bpm (1/min)		
	ii. Positive End Expiratory		
	Pressure (PEEP) 0 to 20		
	cmH20 or OFF, 3 to 20 cmH20.		
	Up to 30cm H2O PEEP is		
	acceptable.		
	iii. Inspiration/Expiration Ratio		
	(Ti:Te) 4:1 to 1:4 2:1-1:8		
	iv. Pressure Limiting (Pmax) 15 to		
	70 cmH20 (hPa). Up to 100 cm		
	H20 is acceptable		
	v. Tidal Volume (Vt) 20 to 1500		
	mL in Volume Control		
	vi. Compliance Compensation on		
	Delivered TV		
36 1.1	vii. Low-flow compensation		
_	eter Cardiac Monitor		
Specification			
	least 19-inch color LCD TFT		
-	pacitive touchscreen with at least		
170	degree viewing angle		

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Suitable for monitoring the following patients: adult, pediatric, and neonate Able to display 10 to 12 real-time 3. waveforms Full capacitive touchscreen with multi-touch operation. Units with navigation wheel, hard keys in combination with capacitive touchscreen are acceptable. With Transport Module a. With at least 5.5-inch monitor b. Able to display 10 to 12 waveforms c. Parameters: 3/5 lead ECG, SPO2 (Nellcor or equivalent), Respiration, NIBP (Variable cycle), Sidestream or Mainstream CO2: **Invasive Blood Pressure** d. Battery: Lithium Ion with at least 3-hours capacity Parameters: 6. a. 12-Lead ECG b. Impedance respiration c. SpO2 d. Non-invasive blood pressure e. Dual temperature End Tidal CO2-Sidestream or mainstream **Dual Invasive Blood Pressure** With arrhythmia analysis including atrial fibrillation. With motion tolerant SpO2 signal processing and perfusion indicator to determine signal quality NBP measurement modes: a. Manual b. Automatic Stat. d. Venous puncture mode.

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10. With the following Screen layout

a. 10 to 12 waveform layout

choices:

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	b. Short trend layout
	c. Large number layout
	d. Oxy-Cardiorespirogram
	e. Cascading ECG Layout
	f. 7-12 ECG layout
	11. Can derive a multi-axis portrait from
	the ST analysis to help detect changes
	in ST values.
	12. Able to create a record of events or
	screen image capture capability and
	transfer PNG or PDF data to USB
	13. Can be connected to a central station
	using standard wired connection. Can
	be upgraded to connect using wireless
	connection.
	14. Back-up Battery: Lithium-ion battery
	capable of at least 2 hours operation
	15. With alarm reminder: Audio and
	visual reminder
	16. Can store up to at least 1000 events
	including parameter alarms,
	arrhythmia events and technical
	alarms. and manual events for the
	current patient
	17. Trend data can store >120 hours of
	trend data for a single patient. Full
	disclosure that can store up to 48
	hours for all parameters and
	waveforms.
	18. With at least 3 default profiles
	(General ward, ICU, Operating Room)
	and at least 3 customized profile.
	19. With "demo" feature for staff training
	in lieu of a demo mode, the supplier
	may provide a laptop or a simulator
	for training purposes. 20. Capable of external slave display
	connection
	21. HL7 Compliant output
	22. With USB port and speakers
	23. With hight mode to automatically
	reduce screen brightness, alarm
	reduce screen drightness, alarm

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volume and HR/pulse tone volume to minimize noise and promote anesthesia induction 24. Capable for dash board connectivity that allows hospital biomedical engineers to quickly view the status of the monitor, upgrade monitor software, and update configuration settings from a remote server 25. With Integrated handle for portability. 26. Capable of connection to a global rack and support the following modules: BIS, dual IBP, microstream or mainstream, and Gas 27. Power Supply: Autovolt 100 -240 Volts AC, 50 to 60Hz 28. Resistant to AC and high-frequency electro surgical interference from devices (e.g., cautery, defibrillators, 29. The following module must be provided/unit: a. One (1) Neuromuscular Transmission (with adult and pediatric mechanosensors for blockade monitoring modes: single twitch, TOF, DBS, tetanus, PTC; nerve localization mode with electrosensor). Stand-alone NMT Module is also acceptable. 30. Each unit must be delivered with the following: Two (2) sets of the following NBP cuffs with interconnect tubing: Big/Large Adult NBP cuffs ii. Adult NBP cuffs iii. Small Adult NBP cuffs iv. Pediatrics cuffs v. Infant cuffs vi. Neonate cuffs b. One (1) pc Reusable SpO2 sensor

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(Adult)/unit

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		with grabber cable or clip type/unit One (1) pc Power cord/unit. Power cord must be compatible with existing outlet. One (1) Manual/Unit One (1) Unit AVR appropriate for the machine (Third Party)		
Approved Budget for the Contract (ABC) =>			PHP12,000,000.00	

TERMS & CONDITIONS:

A. Requirement/s if declared as Lowest/Single Calculated Bids:

1. Presentation of Technical data sheet and/or presentation of a prototype equipment within seven (7) calendar days after receipt of Notice of Lowest / Single Calculated Bid. Presentation may be an online demonstration.

B. Requirement/s if awarded the contract:

- 1. Delivery Period: Within Ninety (90) Calendar days after receipt of Notice to Proceed (NTP).
- 2. Delivery Place: Equipment Section, Property & Supply Division, Philippine General Hospital, Taft Avenue, Manila.
- 3. Warranty Period / Coverage of Warranty: One (1) year on parts and Two (2) years on service. Free quarterly preventive maintenance during the warranty period. Warranty Period shall commence from the date of acceptance by the end user after installation, testing and commissioning.
- 4. Signed service level agreement with the Philippine General Hospital.
- 5. Original hard copy (not photocopy) or soft copy of operators and service manuals in English Language.
- 6. Product Orientation for end users and troubleshooting training for at least two (2) biomedical engineers.
- 7. Quotation of the Annual Preventive Maintenance Cost after the warranty period expires.
- 8. Acceptance Procedures and Parameters: Visual and functional testing.

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C. Documents required of the bidder to submit during Post-Qualification:

- 1. Brochures/Technical data Sheet.
- 2. SEC registration to prove that the supplier is in the business of importing and supplying medical equipment.
- 3. Certified true copy of the Certificate of Distributorship for the last five (5) years. The principal and the local distributor must have been in business partnership for at least five (5) years.
- Certification that the Brand has been in the local/international market for at least five (5) years. Proof required: Invoices or Purchase Orders.
 Certification from supplied hospitals is also acceptable proof of brand presence.
- 5. Certification by the supplier that at least one service engineer is available locally to provide quick on-site support.
- 6. List of local Service Center/s
- 7. Certificate of Performance Evaluation from the Single Largest Contract.
- 8. License to Operate (LTO) from the Philippine FDA.

D. Requirements Required of the Manufacturer to be submitted during Post-Qualification:

- 1. Certification that the manufacturer has been in the business of manufacturing hospital equipment for at least 10 years.
- 2. Guarantee Letter from the manufacturer to ensure availability of supplies, parts and accessories for at least five (5) years after expiration of the warranty period.
- 3. Certification by the principal that service engineers are factory trained on service and repair.
- 4. ISO/IEC compliance document of the manufacturer.
- 5. List of the manufacturer's office and contact details in the following territories: Western Europe, US/Canada and Japan.

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