

Rizal Avenue, Sta. Cruz, Manila

BAC SECRETARIAT

SEALED BID

SEALED BID NO.: HBAC-A2-2020-06-0702 Date: 01 June 2020

Gentlemen:

Please quote your price for item/s indicated below. Price quoted is final, and without condition and we of all taxes. Bid Proposal should be submitted by inclusive of all taxes. Bid Proposal should be submitted in sealed envelope, properly labelled, indicating the date and Sealed Rid No. not be submitted in sealed envelope. date and Sealed Bid No., not later than 2:00pm of 04 June 2020

BAC Secretariat Office and thereafter publicly opened. Any change/erasure must be duly initialed otherwise the bid of the public opened. otherwise, the bid shall be disqualified. Attendance of authorized representative is not mandatory on bid opening. In accordance with Section 52.2 of the opening. In accordance with Section 53.2 of the Revised Implementing Rules and Regulation of the Republic Act 9184. Republic Act 9184.

THE COMMITTEE RESERVES THE RIGHT TO REJECT ANY AND ALL BIDS, DECLARE A FAILURE OF BIDDING AT ANY TIME PRIOR TO THE CONTRACT AWARD, OR NOT TO AWARD THE CONTRACT, WITHOUT THERERY INCURRING ANY LIABRITUDE. WITHOUT THEREBY INCURRING ANY LIABILITY, AND MAKE NO ASSURANCE THAT A CONTRACT SHALL BE ENTERED INTO AS A RESULT OF THE BIDDING. EMELITO O. VALDEZ-TAN, MD

Chairman, HBAC-A2

PR No.: 2020-05-2923 (HEMC)

PLEASE INDICATE BRAND / MODEL IF APPLICABLE

D. J. M /D Marriana			ABC	Unit Price	Total Price
Qty.	Unit	Description/Particulars CARDIAC MONITOR	1100		
25	Units	BRANDED, BRAND NEW			
		Display Monitor At least 12 inches size			
		> Color TFT touch screen			
		Capable of displaying 4 waveforms	to 6		
		ECG/Cardiac Monitoring ECG waveform display (at lead II, Lead II, III) and with corresponding sound on each QRS waveform.	Lead beep		
		> Heart rate display			
		With arrhythmia Analysis Calculation and Pace Analysis	S, ST 200,000.00		
		> Lead selection switch			
		> Sensitivity switch: 2.5 t mm/mV	0 20		
		> Filter switch for interference	from		
		Mains power frequer	icy		
		Low and high pass si	gnal		
		Common Mode Rejection (more than +100dB	CMR):		
		ECG signal measurement ran mV to +2 mV.	ge: -2		
		Frequency range: At least 0 150 Hz or wider range	.67 to		
		Input impedance: 2.5 MΩ at 1	0 Hz		

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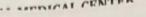
- Frequency response: -3dB at 0.05 Hz to 100 Hz
- Automatic internal data storing for at least 40 ECG records
- ECG leads connector for at least 3lead patient cable (protection from interference)
- Patient cable with at least 3 leads with electrical screening
- ECG recorder/printer capable of printing at least 4 waveforms simultaneously, Arrhythmia Analysis, ST Calculation and Pace Analysis and heart rate, etc.

Pulse Oximeter

- Hinge finger probe or rubber finger probe and ear sensors for adult, pediatric and infant use. The connection of the probes to the main unit must have locking mechanism.
- Oxygen saturation (Sp02): 70 to 99% with minimum graduation of 1%.
- Pulse rate in beats per minute (bpm). Pulse rate range at least 30 to 240 bpm, with minimum gradation of 1 bpm.
- Pulse waveform or indicator that illustrates the strength of pulse being detected.
- Sp02 limit alarm activation settings.
- Pulse rate limit alarm activation settings
- Alarm sound level adjustment and alarm override and temporary silence control
- Accuracy of SpO2 measurement: ± 3 %
- Accuracy of pulse rate measurement: ± 5 bpm

Temperature Measurement

- Digital thermometer temperature probe
- Body temperature measured at degrees Celsius with measurement





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range of 32 - 43 °C

- Measurement accuracy: ± 0.1 °C between 35°C to 41°C
- Non-Invasive Blood Pressure Monitor (oscillometric method)
 - Inflatable rubber cuff surrounded by durable and flexible cover and with Velcro strips
 - Rubber tubes with at least 30cm in length
 - Systolic and diastolic blood pressure measurement with a maximum pressure reading of 300mmHg
 - Reading accuracy: ± 5 mmHg or better
 - Measures blood pressure at least every 10 minutes
- Respiratory Monitoring
 - Thoracic impedance measurement via ECG leads
 - Breaths per minute with measurement range of 0-120 BPM
 - > Respiratory waveform display
 - Measurement accuracy: ± 3 BPM or better
- Safety Features
 - Protection against defibrillation and electrosurgical equipment
 - Equipment compatible with patients with pacemakers
 - Degree of protection against electrical shock: Type CF
 - Alarms (the equipment must have an alarm setting switch and sound adjustment switch)
 - 4 Arrhythmia
 - Ventricular fibrillation
 - Tachycardia
 - Bradycardia
 - Electrode and/or sensor disconnection
 - High and low Sp02
 - High and low pulse rate
 - 4 Sensor failure
 - Apnea Alarm

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Low battery

Power Supply

- Autovolt at 100 240V AC, 60 Hz or 220V, 60 Hz with external Automatic Voltage Regualtor (AVR) with at least 1 kVA capacity
- With internal re-chargeable backup battery that can allow the equipment to operate up to 3 hours

Mobility

Mounted onpole stand or cabinet cart with 4 anti-static and rust-free swivel wheels with two locking brakes cart with brakes.

Accessories

- Protective case
- > Two (2) sets of ECG electrodes
- Non-removable embossed DOH letters on the visible part of the equipment

IF CENTRAL MONITORING IS REQUIRED BY THE PROCURING ENTITY

- The Patient Monitor must have a provision for telemetry data transmission for central monitoring
- Central Station Monitoring System
 - LED touch screen with resolution of at least 1920 x 1080 pixels and at least 32 inches size
 - Data reception must be through telemetry
 - With licensed Operating System software
 - Capable of receiving and displaying data simultaneously from at least 16 patient monitors
 - Capable of trend review per patient
 - Capable of alarm history review
 - Capable of freezing data



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for further review and analysis

- Capable of graphic and tabular data trend presentation
- CPU: at least 2 processors with minimum of 4 cores and 4 threads, minimum of 2.6 GHz
- ♣ RAM: at least 16 GB
- Video Card: at least 4 GB video RAM
- Hard drive/storage: at least 2 TB
- **USB** ports
- Licensed Operating
 System (OS)
- 4 220V, 60 Hz
- Accessories: Keyboard, mouse, external speaker, Automatic Voltage Regulator, laser printer

Documentary Requirements:

- Mayor's Permit
- Omnibus Sworn Statement
- Income / Business Tax Return
- Product brochure or technical data sheets of the equipment showing the technical specifications in English Language
- Valid and current Certificate of Compliance
 of Manufacturer of the Equipment with the
 latest version of ISO 13485: Quality
 Management System Requirements for
 Regulatory Purposes in the name of the
 manufacturer. The Certificates must be
 issued by an independent Certifying Body /
 Agency
- Valid Marketing Authorization, Registration Approval or Free Sale Certificate for the equipment issued by the Health Authority in the Country of Origin.
- Valid Certificate of Distributorship (as first Tier Distributor) issued by the Manufacturer of the Equipment Authorizing the bidder to sell / distribute

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the offered equipment

- List and address of the equipment manufacturer's branch office, sales office and / or distributor's office in any of the following:
 - > Western Europe;
 - > USA or Canada or;
 - > Japan
- Proof (such as sales invoice) that the Brand of the equipment has been sold to other health facilities in the Philippines.
- Notarized Certificate from the Bidder:
 - That the brand of the equipment has been in the local and / or international market for at least ten (10) years.
 - That the equipment and its accessories are brand new, unused, not-discontinued models and were not subjected to any product recall
- Bidder's valid and current License to Operate (LTO) as a medical device distributor issued by the Philippine Food and Drug Administration. In case of expired LTO, the following must be submitted:
 - Copy of expired LTO
 - > Application for renewal
 - > Official Receipt as proof of payment for the renewal of LTO

Requirements if Awarded the Contract (Please attach a Certification that the following will be provided / will be followed upon delivery of the item)

- 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 <u>Seven (7) Calendar Days</u> upon receipt of the Notice to Proceed.
- 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.
- Training: The supplier shall provide a training on the proper use and maintenance of the equipment to the endusers and to the hospital.
- Warranty: Warranty certificate for two



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(2) years on Parts and on Services. 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	
Notarized undertaking that the supplier shall conduct the necessary corrective maintenance within five (5) calendar days upon notification of 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.	
Manuals: The supplier must provide the end-user one (1) hard and one (1) soft copy of the following: > Service Manual in English Language > Operations manual in English Language	

Check Mode of Payment:1. Terms/Deferred Payment	Name of Company
	Address/TIN No.
THE THE PERSON OF THE PERSON O	Authorized signature over printed name/ Telephone no. quotation, is conforming to our payment terms you check. ; WITHIN 7 WORKING DAYS UPON RECEIPT OF NTP AWARDS COMMITTEE FOR SUPPLIES, MATERIALS & EQUIPMENT
Chairman	
Vice-Chairman	
Member	
Member	
Member	
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