

Technical Specifications

Note: Bidders must state in the Statement of Compliance either “*Comply*” or “*Not Comply*” against each of the individual parameters of each Specification stating the corresponding performance parameter of the equipment offered. Statements of “Comply” or “Not Comply” must be supported by evidence in a Bidders Bid and cross-referenced to that evidence. Unless otherwise categorically stated hereunder, evidence shall be in the form of manufacturer’s un-amended sales literature, unconditional statements of specification and compliance issued by the manufacturer, samples, independent test data etc., as appropriate. A statement that is not supported by evidence or is subsequently found to be contradicted by the evidence presented will render the Bid under evaluation liable for rejection. A statement either in the Bidders statement of compliance or the supporting evidence that is found to be false either during Bid evaluation, post-qualification or the execution of the Contract may be regarded as fraudulent.

A. General Specifications

No.	General Specifications	Statement of Compliance
1	Supply, Delivery, Testing and Commissioning of Brand New One (1) Unit of Nasal High Flow	

Detailed Technical Specifications

ITEM NO.	ITEM	SPECIFICATIONS	STATEMENT OF COMPLIANCE
1	Nasal High Flow		
	<u>Clinical Features:</u>		
	Temperature Settings	31 - 37 C	
	Max Temperature	43 C	
	Humidity Requirement	> 12 - > 33 mg/L (10 - 95% Relative Humidity)	
	Flow Range	10 - 60 Lpm (minimum range)	
	Oxygen Output	60Lpm	
	FiO2 Monitoring Range	21 - 100%	
	FiO2 Monitoring Accuracy	+ / - 4%	
	Warm - up Time	at least 10 - 30 mins.	
		(at temp range 31 - 37 C)	
	Alarms	< / = 45 dbA	
	Alarm Silence	At least 1 min.	
	<u>Physical</u>		
	Weight	less than 10 lbs.	
	Mobility	with stand and at least 4 wheels	
	<u>Electrical</u>		
	Frequency	50 - 60Hz	
	Current/Voltage Requirement	Autovolt (110 -240)	

	UPS (Uninterrupted Power Supply)	(at least 30 mins to 1 hour)	
	Accessories		
	Breathing Tubes	Disposable	
	Humidification Chamber	Disposable	
	HFNC Interface	Small, Medium, Large	
		Documentary Requirements:	
		1. Certificate of availability of stocks and can deliver within the prescribed delivery schedule.	
		2) Valid and current License to Operate (LTO) as medical device distributor issued by the Philippine Food and Drug Administration. In Case of expired LTO, the following must be submitted: i) Copy of expired LTP, ii) Application for renewal, iii) Official Receipt as proof of payment for the renewal of LTO	
		Requirements if awarded the Contract:	
		1. Delivery and Completion Schedule: Ten (10) working days with no extension upon receipt of approved NOA and NTP. Failure to comply with the delivery schedule shall mean forfeiture of the contract	
		2. Testing: Prior to acceptance, the COVID Task Force / 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 an orientation/ training on the proper use and maintenance of the equipment to the end-users.	
		4. Warranty: Warranty certificate for One (1) year on parts and 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 testing and commissioning.	
		5. Manuals: The supplier must provide Manual in English Language (2 hard copies and 1 soft copy of Operator's/User's Manual and 2 hard copies and 1 soft copy of Service/Technical and Maintenance Manual) upon delivery	

		6. Product brochure or technical data sheet(s) of the equipment showing the technical specifications in English Language.	
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<i>Conforme:</i>
<i>Company/Bidder's Name:</i>
<i>Name and Signature of Authorized Representative:</i>
<i>Date:</i>