Technical Specifications of Ventilator

S.N.	Purchaser's Specifications of Ventilator	Bidder's Compliance Sheet			
	Ventilator (Infant to Adult)	Yes/No	Page no. in Catalogue	Remarks	
	Manufacturer				
	Brand				
	Type / Model				
	Country of Origin				
1	Description of Function				
1.1	Advanced automated adaptive assisted wide-range turbine/blower based ventilation system.				
2	Operational Requirements				
2.1	Electronic microprocessor controlled ventilator				
	with integrated facility controlled by volume and				
	pressure to fit for all types of patients from				
	neonate to Adults.				
3	Technical Specifications				
3.1	Should have TFT Touch Screen of minimum 15 inch				
3.2	Should be able to easily switch between a NIV				
	mode and IV modes by UI operation only				
3.3	. It Should have following Ventilation Mode:				
	A Investive Vantilation Made				
	A .Invasive Ventilation Mode:				
	• Assist control (Volume Control Ventilation)				
	Assist control (Pressure Control Ventilation)				
	VSIMV (Volume Synchronized Intermittent				
	Mandatory Ventilation)				
	PSIMV (Pressure Synchronized Intermittent				
	Mandatory Ventilation)				
	CPAP/PSV (Continuous Positive Airway Description (Continuous Positive Airway				
	Pressure/Pressure Support Ventilation)				
	• PRVC (Pressure Regulated Volume Control)				
	• BiPAP (Bi-level Positive Airway Pressure)				
	 APRV (Airway Pressure Release Ventilation) 				
	 Aprea Ventilation 				
	B. Non-invasive Ventilation Mode:				
	D. Ron-myasive ventilation wode.				
	• PC-ACV(Pressure Controlled – Assist				
	Control Ventilation)				
	PVC-ACV(Pressure Volume Controlled-				
	Assist Control Ventilation)				
	 PC-SIMV(Pressure Controlled – 				
	Synchronized Intermittent Mandatory				
	Ventilation)				
	 PVC-SIMV(Pressure Volume Controlled – 				

	Synchronized Intermittent Mandatory			
	Ventilation)			
	PC-Dual PAP(Pressure Controlled-Duo			
	Positive Airway Pressure)			
	• PC-APRV(Pressure Controlled – Airway			
	Pressure Release Ventilation)			
	CPAP/PSV			
	CPAP-VG(Continuous positive airway			
2.1	pressure-Volume guarantee)			
3.4	It should have following Measured and displayed			
	patient parameters:			
	• Tidal volume: Neonates:5 to 200mL			
	Pediatric:20 to 400mL			
	Adult:100 to 2000mL			
	• Rate: 1 to 100bpm			
	• Texp : 0.3 to 12.0 s			
	• Tslope : 0.00 to 2.00 s			
	• PEEP/CPAP: 0 to 50 cmH2O			
	 Inspired Oxygen Concentration (FiO2 			
):21 to 100%			
	• I:E ratio: 1:10 to 4:1			
	• Inspiratory time (TI): 0.2 to 5 s			
	• Flow trigger: 0.2 to 15 l/min			
	• Pressure trigger: -0.5 to -20cmH2O			
	• Pressure control: 5 to 90 cmH2O			
	• Pressure support: 0 to 90 cmH2O, added			
	to PEEP/CPAP			
	• Pressure ramp: 0 to 2s			
	• Apnea absence: 10 to 30s			
3.5	Should have O2 therapy with controlled flow of			
	2-50L/min and controlled flow accuracy of \pm			
	(2L/min+10% of Setting) (BTPS) (body			
	temperature, pressure, water vapor saturated) and O2			
	concentration between 21 to 100% with accuracy of			
	\pm (3 Vol.% +1 % of setting).			
3.6	Must have real time graphical representation of			
	 Pressure-time Waveform Flow-time Waveform			
	Volume-time Waveform			
	 Measurement of volumetric capnography 			
	of CO 2			
	Pressure-volume Loop			
	Flow-time Loop			
	Pressure-flow Loop			
3.7	It should have operator-adjustable alarm system			
	with :			
	• Low/high minute volume			
L		I	1	1

	• low/high pressure : 1 - 90 cmH2O		
	• low/high tidal volume : OFF, 10 - 2000 ml		
	• low/high respiratory rate: OFF, 1 - 90 BPM		
	• apnea time : 5 - 60 sec		
	 low/high oxygen 		
	• Optional: low/high CO2, low/high SPO2		
	• Special alarms:		
	• Oxygen concentration,		
	Circuit disconnection,		
	loss of PEEP		
	• exhalation obstruction,		
	 flow sensor power supply betteries 		
	 power supply batteries, gas supply failure 		
3.8	should have Internal and swappable Li-ion batteries		
	with battery running time of 3 hours and 2 Backup		
	battery batteries of 5200mAh each		
3.9	Should have apnea ventilation facilities with		
	volume-controlled apnea ventilation.		
3.10	Must have integrated self-check system with leak		
	check and system tightness to secure the		
	ventilation result		
3.11	Should have synchronized Tube Resistance		
	Compliance with:		
	• ET Tube, Trach Tube		
3.13	Should have sigh switch that can be turned		
	ON/OFF with interval of 30s-3 hrs except for		
0.1.1	CPAP/PSV, DuoLevel, and APRV.		
3.14	Should have communication Interface Nurse call,		
2.15	USB, NET, RS232, VGA		
3.15	Should have logging capability that can store and		
216	display of up minimum log of minimum 1000		
3.16	Should have operating temperature of 5-40°C		
	,Relative Humidity of 10-95% for both operating		
	,storage and transport and Barometric Pressure		
2 17	of 62-106kPa		
3.17	Must have value added functions of nebulizer,		
	PV Tool, P 0.1, Inspiratory Hold, Expiratory		
4	Hold, NIF and leakage compensation		
4 4.1	Accessories, spares and consumables Accessories:		
4.1	Trolley 1/Pedestal rotatable		
	trolley with castors brake		
	Neonatal Humidifier chamber		
	Resuable pediatric patient		
	circuit/W Heating		

	Support arm		
	 Disposable bacterial filter 		
	Reusable Neonate patient		
	circuit /W Heating		
	circuit / w Heating		
4.2	All standard accessories, consumables and parts		
	required to operate the ventilator to be included in		
	the offer. Bidders must specify the quantity of every		
	item included in their offer (including items not		
	specified above).		
5	Operating Environment		
5.1	The product offered shall be designed to be stored		
	and to operatenormally under the conditions of the		
	purchaser's country. The conditions include Climate,		
	Temperature, Humidity, etc.		
5.2	Power supply: 220-240v, 50/60Hz fitted with		
	appropriate plug. The power cable must be at least 3		
6	meter in length. Standards and Safety Requirements		
6.1	Must submit ISO13485:2003/AC:2007 AND		
6.2	Must submit European CE certificate		
7	User Training		
7.1	Must provide user training (including how to use and		
	maintain the equipment).		
8	Warranty		
8.1	Warranty for 2year after acceptance.		
9	Maintenance Service During Warranty Period		
9.1	During the warranty period supplier must		
	ensure planned preventive maintenance (PPM) along		
	with corrective/breakdownmaintenance whenever		
10	required. Installation and Commissioning		
10.1	The bidder must arrange for the equipment to be		
10.1	installed and commissioned by certified or qualified		
	personnel; any prerequisites for installation to be		
	communicated to the purchaser in advance, in detail.		
11	Documentation		
11.1	User (Operating) and Service (Technical /		
	Maintenance) manualin English.		
11.2	The bidder should submit the original brochure or e-		
11.0	copy.		
11.3	The bidder should submit a valid authorization		
11 7	letter from the manufacturer		
11.5	The bidder should compulsorily fill the technical		
	Specification tender form and clearly mention		
	theManufacturer, Brand, Model and Country of		
	Origin.		