

Bid Number: GEM/2020/B/710802

Dated: 15-07-2020

Bid Document

Bid bocument				
Bid Details				
Bid End Date/Time 18-07-2020 17:00:00				
Bid Opening Date/Time	18-07-2020 17:30:00			
Bid Life Cycle (From Publish Date)	90 (Days)			
Bid Offer Validity (From End Date)	30 (Days)			
Ministry/State Name	Ministry Of Steel			
Department Name	Na			
Organisation Name	N/a			
Office Name	Kiocl Limited/ Bangalore			
Total Quantity	2			
Item Category	ICU Ventilator			
Experience Criteria	1 Year (s)			
MSE Exemption for Years of Experience and Turnover	No			
Startup Exemption for Years of Experience and Turnover	No			
Document required from seller	Experience Criteria,OEM Authorization Certificate *In case any bidder is seeking exemption from Experience / Turnover Criteria, the supporting documents to prove his eligibility for exemption must be uploaded for evaluation by the buyer			
Bid to RA enabled	Yes			
Inspection Required	No			

EMD Detail

Required	No
Nequired	110

ePBG Detail

Required	No
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Splitting

Bid splitting not applied.

1. Experience Criteria: In respect of the filter applied for experience criteria, the Bidder or its OEM {themselves or through reseller(s)} should have regularly, manufactured and supplied same or similar Category Products to any Central / State Govt Organization / PSU / Public Listed Company for number of years as indicated in the bid document before the bid opening date. Copies of relevant contracts to be submitted along with bid in support of having supplied some quantity during each of the year. In case of bunch bids, the category of primary product having highest value should meet this criterion.

ICU Ventilator (2 pieces)

Technical Specifications

* As per GeM Category Specification

Specification	Specification Name	Values	Bid Requirement (Allowed Values)
Standards	Certifications for the product	-	*
	Certification number and date	-	*
	Confirmity to quality management standards	ISO 9001 & ISO 13485	*
	Confirmity to electrical safety standards	IEC 60601-1 or ISO 80601-2-12 or BIS equivalent	*
Performance	UMDNS Code	17-429	*
Parameters	Scope of item	Intensive care ventilators are defined as mechanical ventilators that can be configured to provide invasive ventilation (e.g., with an endotracheal tube or tracheostomy tube) or noninvasive ventilation (eg, with a face mask)	*
	Clinical application	Mechanical ventilators are life support devices that move gas (e.g., air and/or oxygen) to and from a patient's lungs. These devices may provide temporary or permanent respiration for patients who cannot breathe on their own, or who require assistance maintaining	*

		adequate ventilation	
	Patient Type	Adult & pediatric	Adult & pediatric, Adult, pediatric and neonatal
	Type of technology	Compressor	Turbine, Compressor
	Tidal volume in ml	50 ml or less - 1500 ml or more	50 ml or less - 1500 ml or more, 2 ml or less to 1500 ml or more, 100 ml or less-1500 ml or more
	Respiration rate, breaths/min	up to 80 or more	upto 150 or more, up t 80 or more, Up to 120 or more, up to 50 or more
	Trigger mechanism	Pressure /flow	Pressure /flow, Both pressure and flow
	FiO2%	21-100	*
	Inspiratory flow rate, L/min	upto 120 or more	upto 120 or more, upto 30 or more, up to 150 or more
	Inspiratory pressure, cm H2O	upto 80 or more	upto 50 or more, upto 80 or more, up to 150 or more
	IE ratio	1:10 to 4:1	1:10 to 4:1, 1;6 to 3:1
	Sigh breath function	Yes	Yes, No
	PEEP/CPAP, cm H2O	upto 40 or more	*
	Pressure support, cm H2O	0-40 or more	*
	Leak compensation	Yes	*
	Auto 100%/Increase O2 button	Yes	*
	Control panel lock	Yes	Yes, No
	Facility for double lung ventilation	Yes	Yes, No
Patient Assessment Tools	Maximum wave forms displayed	3 (Pressure and time, volume and time , flow and time)	*
	Number of loops	2 loops (P-V, F-V) with facility of saving of 2 Loops for reference	2 loops (P-V, F-V) with facility of saving of 2 Loops for reference, 2 loops (P-V, F-V)
	Maximum trending time in hrs	≥24 hr	*
	Lung recruitment tools (PV loops) / OLT (Open lung tools)	Yes	Yes, No

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Lung mechanics visualization tool	No	Yes, No
Capnography/CO2 monitoring	No	Yes, No
Esophageal / transpulmonary pressure monitoring	No	Yes, No
Stress index	No	Yes, No
Modes of ventilation	Volume controlled, Pressure Controlled, Pressure Support, SIMV with Pressure support ,SIMV with volume control with pressure support, ,CPAP/PEEP,Inverse Ratio Ventilation,Non invasisve ventilation,Apnea /back up ventilation,SIMV (VC) with Pressure support; SIMV (PC) with Pressure Support; SIMV (PRVC) with Pressure Support	*
ASV(Adaptive support ventilation) or ALT	No	Yes, No
APRV/Bivent/Bi level	Yes	Yes, No
MMV +PSV /ASV or ALPV or Automode	No	Yes, No
PRVC/Auto flow	Yes	Yes, No
High flow oxygen therapy	No	Yes, No
Automatic weaning system provided	Yes	Yes, No
Volume Support	Yes	Yes, No
Ventilation monitoring facility	The Ventillator is a Microprocessor Controlled ventilator with integrated facility for Ventilation monitoring	*
Monitored / Displayed parameters	Peak inspiratory pressure , Mean airway pressure, PEEP pressure, Tidal volume, Minute volume, Spontaneous minute volume, FiO2 (analyzed	*

Availability of Status indicator for Ventilator	%), Respiratory rate , Inspiratory time, Expiratory time, IE ratio, Plateau Pressure Yes	*
mode, Battery life, patient data, alarm settings, clock etc on display		
Patient alarms	Low/high FiO2, Low/high minute volume , Low inspiratory pressure , High pressure, Low PEEP , High PEEP , Apnea, Continuous high pressure/occlusion, Inverse IE, High respiratory rate, Breathing circuit disconnect	*
Equipment alarms	Gas-supply failure , Power failure, Vent inoperative, Low battery, Self-diagnostic	*
Non-forced Slow Vital Capacity	No	Yes, No
Physiologic Dead space	Yes	Yes, No
RSBI	Yes	Yes, No
Imposed work of Breathing (WOBi)	Yes	Yes, No
Expiratory Time constant (Tcexp)	Yes	Yes, No
Compliance static and dynamic	Yes	Yes, No
Inspired and expired resistance	Yes	Yes, No
Occlusion pressure	Yes	Yes, No
Inspiratory and expiratory hold	Yes	Yes, No
Spontaneous frequency	Yes	Yes, No
Total peep, intrinsic peep, extrinsic peep	Yes	Yes, No
Expiratory block is autoclavable and no routine calibration	Yes	Yes, No

required]
Auto compensation for ET Tube	No	Yes, No
Facility for automatic compliance & Leakage compensation for circuit available	Yes	Yes, No
RS 232 out put port	Yes	Yes, No
USB/Memory card	No	Yes, No
Availability of Remote alarm/display port	Yes	*
Facility to Report (vent alarms and patient status)	Yes	*
HLT Compliant out put	Yes	Yes, No
Type of coloured display	TFT	LED, LCD, TFT
Display should be touch screen	Yes	Yes, No
NIV (Non Invasive Ventilation) to be possible in all modes of ventilation available	Yes	Yes, No, NIV available in pressure targeted modes only
Size of display (in inches)	15	12, 15, 17, 18
Graphic display have automatic scaling facility for waves	Yes	Yes, No
Power supply	220-240 V , 50 Hz AC single phase	*
Provision of UPS	No	Yes, No
Backup time in hrs	1	1, 2, NA if not provided
Internal backup battery	Yes	*
Backup time for internal battery in mins	45 mins or more	*
Built in air source	Air compressor (external)	Air compressor (external), Turbine (In built)
Availability of stand alone compressor	No	Yes, No
Compressor certification	European CE	US FDA, European CE, Both European CE and US FDA, BIS

	Compressor shall provide an oil free Medical air, with Peak output flow should be minimum 150 LPM and Air quality complying with ISO compressed air purity class	Yes	Yes, NA (for turbine)
	Medical Air Compressor should automatically activate in the event of wall air supply loss and replacement of internal filters should be performed without removing the compressor, have washable air filter	Yes	*
	Seller shall ensure compatibility of compressor with ventilator	Yes	Yes, NA for Turbine
	Inbuilt nebulizer with particle size less than 3 micron	Yes	Yes, No
	Comprehensive Warranty in years	5	5
Additional Accessories	Reusable silicon breathing circuits for Adult,pediatric and Neonatal	2 each	*
	NV mask(Re usable)-Small,medium and large	2 Each	*
	Length of Power cord in m	>3 m	*
	Type of flow sensor	Disposable	Reusable, Disposable, NA
	No of reusable flow sensor	2 no.	*
	No of disposable flow sensor	50	50, 100, NA
	Autoclavable exhalation valve/expiratory cassette / filter	2	*
	Humidifier: Servo controlled with digital monitoring of inspired gas	1	1, 2, NA

	temperature with heating wire		
	Trolley for ventilator with circuit holding arm	Yes	*
Miscellaneous Parameters	Number of installations in Central /State/PSU Govt Hospitals (Hint: Seller should supply a performance certificate of the device to the buyer incase demanded after placement of order)	0	*
	OEM/Reseller (if supplied by reseller shall ensure uninterrupted availability of all spares for 10 years)	Yes	*
	Availability of toll free facility for technical support maintened by OEM or authorized agencies	Yes	*
	During the warranty period it shall be ensured that all the break down calls are attended with in 24 hrs and the complete details of service agents / contact details should be furnished to buyer and consignee at time of supplies	Yes	*
	User/Technical/Mainten ance manuals to be supplied in English in hard and soft copy	Yes	*
	Details of equipments and procedures required for local calibration and routine maintenance to be supplied and advanced maintenance task documentation also to be furnished	Yes	*
	List of important spares and accessories, with their part numbers and price lists to be supplied to the buyer	Yes	*

	time of supplying uipment		
equipn to be p comple	nstration of nent and training provided after eting supplies acceptance	Yes	*
have d Presen	acturer must	Yes	*
certific furnish deman supplie	of reports and ations to be ed to buyer on d at time of	Yes	*

^{*} Specifications highlighted in bold are the Golden Parameters.

Additional Specification Documents

Consignees/Reporting Officer and Quantity

S.No.	Consignee/Reporti ng Officer	Address	Quantity	Delivery Days
1	Kumaravelu Kathaiyan	575010,KIOCL LIMITED, PELLET PLANT UNIT, PANAMBUR, MANGALORE, PIN - 575 010	2	30

Special terms and conditions for category ICU Ventilator

1. Comprehensive warrantyComprehensive warranty shall include preventive maintenance including calibration as per technical/ service /operational manual of the manufacturer, service charges and spares,. During the warranty period commencing from date of the successful completion of warranty period. Service personnel shall visit each consignee site as recommended in the manufacturer's technical/ service /operational manual, at least once in six months, warranty shall not be including the consumables .Further there will be 98% uptime warranty during warranty period on 24 (hrs) X 7 (days) X 365 (days) basis, with penalty, to extend warranty period by double the downtime period. Service centres Details of Service outlets in India to render services for equipment to be furnished to buyer/consignees with complete address ,telephone numbers, e mails etc at time of making the supplies .lt shall be the responsibility of seller to ensure that authorized service centres are available to cater to the areas where supplies are made within reasonable distance from where the service calls can be handled .Details of toll free numbers for service call and online registration of service requests also to be provided buyer/consignee at the time of supplies. Source of supplyIt shall be responsibility of seller to provide Documents regarding source of equipments such as copy of Performa invoice or any other documents to establish that the products supplied are manufactured by OEM indicated and sourced from

^{*} Bidders may note that In respect of non-golden Parameters, the specifications 'Values' chosen by Buyer will generally be preferred over 'Bid requirement (allowed Values) by the Buyer.

them. Packing and Marking Medical equipments being very delicate and sensitive packing for the goods should be strong and durable enough to withstand transit including transhipment (if any), rough handling, open storage etc. without any damage, deterioration etc. .The size, weights and volumes of the packing cases, remoteness of the final destination of the goods, availability or otherwise of transport and handling facilities at all points during transit up to final destination,. Quality of packing, the manner of marking within & outside the packages and provision of accompanying documentation shall take in to consideration the type of medical equipments being supplied. The accessories shall be suitably labelled and packed .Each of the package shall be marked on three sides with indelible paint of proper quality: indicating contract number and date, brief description of goods including quantity,. Packing list reference number, country of origin of goods and any other relevant details. Spare PartsSeller shall provide materials, information etc. pertaining to spare parts manufactured and supplied by the OEM. It shall be ensured that the required spares are available for purchase at least for 10 years from date of supplies. In case due to any reasons the production of the spare parts is discontinued sufficient advance notice should be given to the buyer/consignee before such discontinuation to provide adequate time to purchase the required spare parts etc. Further, OEM and their service centres/dealers shall carry sufficient inventories to assure ex-stock supply of consumables and spares for the equipments so that the same are available.OEM or reseller shall always accord most favoured client status to the buyer/consignee and shall give the most competitive price for spares and consumables of its machines/equipments supplied.Installation, Training, ManualsSeller shall be responsible to carry out Installation & commissioning, Supervision and Demonstration of the goods. They shall provide required jigs and tools for assembly, minor civil works for the completion of the installation and Training of Consignee's representatives for operating and maintaining the equipment and Supplying required number of operation & maintenance manual for the goods. In case the category parameters are specifying any requirements regarding the installations, training and manuals the same shall also be applicable. Electrical safety checkingSellers are required to make sure that they furnish the list of equipments for carrying out routine and preventive maintenance to buyer/consignee. They should make sure to periodically check the electrical safety aspects as per BIS Safety Standards or equivalent .In case they do not have required equipment for such testing should ensure that the equipments checked for electrical safety compliance through labs with facilities for such checking during every preventive maintenance call. Software All software updates should be provided free of cost during warranty period.

Bid Specific Additional Terms and Conditions

1. Scope of supply (Bid price to include all cost components): Supply Installation Testing Commissioning of Goods and Training of operators and providing Statutory Clearances required (if any)

This Bid is also governed by the General Terms and Conditions

---Thank You---