



Running Contract Details	
Equipment Name	Non Invasive BIPAP Ventilator
Running Contract Valid Till	07-06-2025
Tender Ref No	KMSCL/EP/T430(R)/136B/2021
Tendered Quantity	250
Supplier Name	M/s ResMed India Pvt Ltd
GST No	07AADCR5533K1Z7
Installation & Delivery Period	8 Week(s)
Up-time / PM vist	95% & 4 Visits per year
Warranty period	3 Years

Supplier`s Details		
Address	Contact Details	
1st Floor Worldmark 1 Asset Area 11 Aerocity Hospitality District IGI Airport Delhi-110037	Contact Person	Mr. Vinay Rattan
	Phone	
	Mobile No	9560299926
	Email	enquiries@resmed.co.in,ajay.chambyal@resmed.co.in

Item-wise Price Details				
#	Item Details	Unit Rate (Incl.all taxes & charges)	Service Charges (Through KMSCL)	Grand Total
1	Non Invasive BIPAP Ventilator <i>Model & Make : Stellar 150</i>	233968 Incl.GST :12%	17255.14	251223.14
		233968	17255.14	251223.14

Annual / Comprehensive Maintenance Charges (Exl.Tax)							
Rate	4 th Year	5 th Year	6 th Year	7 th Year	8 th Year	9 th Year	10 th Year
Non Invasive BIPAP Ventilator							
Labour	4,178.00	4,178.00	4,178.00	4,178.00	4,178.00	4,178.00	4,178.00
Comprehensive	12,534.00	12,534.00	12,534.00	12,534.00	12,534.00	12,534.00	12,534.00

Other terms & conditions

1. The supplier shall execute an agreement with the purchaser as per tender conditions (agreement format is given in the tender

document).

2. The supplier shall submit performance security amounting to 5.00% of the value of the supply order.
3. The labour & comprehensive charges of equipment after the completion of warranty period is finalized by KMSCL as mentioned above.
4. Since discount rate is not applicable for equipment under Running Contract of KMSCL, purchase/supply order can be issued directly to supplier at the given rate with tax & other charges (exclusive of KMSCL service charges).
5. If purchase/supply order is issued directly to the supplier, KMSCL service charge need not be paid. But the copy of the said order may be forwarded to KMSCL for information.

Technical Specification

Equipment :Non Invasive BIPAP Ventilator

ItemName:Non Invasive BIPAP Ventilator

1. NIV for adults and pediatrics.
2. Light weight, small, user friendly and quiet device.
3. Should have the following modes. S -T (spontaneous - timed), CPAP (Spontaneous), T (Timed), PAC (Pressure Assisted Control)/ PC (Pressure Control), Volume Assured Pressure Support (VAPS).
4. Should incorporate latest algorithms for leak compensation and synchronization.
5. Should have color screen at least 3.5 inch for real time monitoring of minute volume/ tidal volume, respiratory rate, percentage of leak, I:E ratio, Delivered IPAP and EPAP.
6. Should be able to display real time flow and pressure curves / values simultaneously and the Ti bar graph.
7. Should include user adjustable alarms and essential nonadjustable fixed alarms for patient safety.
8. Should include alarms for leak, power supply failure, apnea, patient circuit disconnection, occlusion, low internal battery etc. and should have adjustable alarms for minute volume, high/low pressure, RR, apnea.
9. Should have oxygen port to accept flow up to 15 l/min of oxygen to achieve a high FiO₂.
10. Should provide and maintain optimal humidification at patient desired temperature regardless of ambient humidity changes throughout night.
11. Pressure range: IPAP- 4/ 2-40 cm H₂O, EPAP-2/4-20cm H₂O.
12. Pressure support 0-30cmH₂O.
13. Respiratory rate 5-40bpm or more.
14. Rise time upto 600msec.
15. Inspiratory time upto 3sec or more.
16. Flow/ auto trigger and cycle settings.
17. Air outlet should be 22mm taper compatible with ISO 5356-1:2004.
18. Machine should be fitted with electrostatic fibre mesh air filter.
19. Should have built in internal battery for minimum 2 hrs of back up and should have capability to add optional external battery
20. NIV ventilator to be supplied with patient ckt 2nos, air inlet filters, power supply pack, reusable face mask standard 3 sizes (Small, medium and Large) 2 pieces each, Oxygen connector, Fio₂ Monitoring accessories.
21. Power supply input 100-240v ac.
22. Should have safety certificate from a competent authority CE issued by a notified body registered in European commission / FDA (US) / STQC CB certificate / STQC S certificate or valid detailed electrical and functional safety test report from ERTL. Copy of the certificate / test report shall be produced along with the technical bid.