medinCNO®mini

Technical specifications



The medinCNO[®]mini CPAP driver is used in combination with the Medijet[®] nCPAP generator to administer CPAP therapy or with a suitable Nuflow[®] nasal cannula to administer High Flow therapy to premature infants, and newborns. The medinCNO[®]mini must be used under the supervision of expert, specially trained staff in a clinical setting, and the patient's oxygen saturation must be monitored at the same time.

Specification

| Dimensions (L x W x H) | 29 x 23.5 x 18.5 cm | |
|--------------------------------|------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|--|
| Weight | 4.75 kg | |
| Gas feed system | Computer-controlled electronic gas mixer with integrated oxygen sensor; flow-controlled Setting range: 0 l/min to 15 l/min Working range: 4 l/min to 10 l/min Accuracy: ±1 l/min (in working range) ±2 l/min (outside of the working range) | |
| Gas feed | Air supply: 300 to 600 kPa (= 3.0 to 6.0 bar) Oxygen: 300 to 600 kPa (= 3.0 to 6.0 bar) | |
| Gas connection | Connector standard: DISS or NIST (as preferred) | |
| Patient flow outlet | Dimensions M22 (OD) or F15 (ID) | |
| CPAP pressure meter connection | Luer-type – 4.3 mm ID | |
| Power supply | 1x internal battery, 14.4 V DC, approx. 3 hours run time, rechargeable External power supply 100 to 240 V AC / 50 to 60 Hz | |
| Display | 7.0" – color, 800 x 480 pixel | |
| Data | Patient pressure (diagram and measurement) Trend: CPAP, flow, FiO ₂ , RR, push frequency (among other things, up to 28 days) | |
| СРАР | Measurement range 0 to 18 mbar (in 0.1 mbar increments) Verification: Redundant measurement by two sensors Accuracy: ± 1.3 mbar | |
| Push (inspiration support) | Setting range Additional flow during the inspiration push: Min: 0 I/min Max: 17.5 I/min (Basic flow + push flow) Duration 200 ms to 2 seconds Manual and automatic triggering of pushes Accuracy ± 1 I/min (if the total flow = basic flow + additional inspiration flow - is within the flow working range) ± 2 I/min (if the total flow = basic flow + additional inspiration flow - is outside of the flow working range) | |
| Leak-Assist | Leakage compensation (±2 L/min to maintain the target pressure) in CPAP mode | |
| RR (respiration rate) | Display of the measured respiratory rate, no breaths up to 120 breaths per minute, in CPAP and Apnea CPAP mode | |
| Apnea time | 2 to 30 s (in 0.1 s increments) | |
| T _{insp} . | 0.2 to 2.0 s (in 0.1 s increments) | |





| Triggering | Pressure trigger, based on the CPAP pressure sensitivity, ± 0.2 to ± 2.0 mbar (in 0.1 mbar increments) | |
|------------------|---------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|--|
| FiO ₂ | Oxygen concentrationSetting range:21 to 100% (in 1% increments, in the flow working range)Measurement range:21 to 100%Alarm settings: ± 2 -5% difference O_2 flush (level: +5, +10, +15% (vol.) above set target oxygen concentration) for 60 sAccuracy: $\pm 3\%$ (vol.) | |
| Alarms | Visual (LED & display) and acoustic (adjustable push alarm settings) Alarm countdown Interface to external alarm system | |
| Safety | Mechanical overpressure valve (opening pressure 4 kPa (= 40 mbar)) Electronic shut-off valve (in the event of an error, interrupts the flow supply to the patient and opens the tubing system to the atmosphere) | |
| External data | USB/RS232 port, export of live and trend data | |
| Operating time | The medinCNO®mini can be used for continuous, long-term operation up to 4 weeks without a restart in the interim. | |

Environmental Conditions

| Operation | Temperature: | 15 to 35°C |
|-----------|-----------------------|---------------------------------------------------------------------------------------------------|
| | Relative humidity: | 20 to 80% (not condensing) |
| Storage | Short-term/transport: | Temperature: -20 to +50°C Relative humidity: 20 to 80% (not condensing) |
| | Long-term: | Temperature: Room temperature (about 20°C) Relative humidity: 20 to 80% (not condensing) |
| | ••••• | |

Description of modes

| CPAP | Standard CPAP; additional function: Leak-Assist |
|------------|-------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| Apnea CPAP | CPAP with apnea detection (automatically generated pushes); additional function: Backup (backup rate: 5 to 120 1/min as a function of $T_{insp.}$ (minimum 0.2 s) |
| NIPPV | Bi-level CPAP with measurement of the I:E ratio as a function of the set rate and inspiration time (minimal I:E ratio 1:1.5) |
| High Flow | Flow emission for Nuflow [®] nasal cannulas in flow range of 0 to 12 l/min, adjustable $\rm O_2$ flush |
| Standby | No flow, deactivated alarms, settings made are saved, keeps medinCNO®mini ready for use |

Software settings

| Languages | EN, DE, FR, IT, NL, ES, EL, CS, PL, NO, DA, SV, RU, LT, ZH, TR, JA, RO |
|----------------|------------------------------------------------------------------------|
| Pressure units | mbar or cmH ₂ O |
| Pressure scale | Three settings: 0 to 10, 0 to 15 and 0 to 20 mbar |

Standards and certifications

| Declaration | The medinCNO®mini is manufactured within an EN ISO 13485 and EN ISO 9001, Council Directive 93/42/ EEC, Annex II excluding Section 4 certified quality management system. The nCPAP device meets the Essential Requirements of Council Directive 93/42/EEC, Annex I. |
|---------------------------------------------|----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| Classification to EC directive 93/42/EEC | Class llb |
| Certifications of IEC | 60601-1, 60601-1-2, 60601-1-8 |
| Class of protection | Class II equipment |
| IP class | IP20 |

Warranty

Period

24 months

Humidification (recommended)

Hamilton Medical

HAMILTON-H900 humidifier

Accessories and configurations

Suitable accessories and possible configurations of the device can be found in the corresponding configuration sheet.



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