HAMILTON-C1

Technical specification for SW version 3.0.x

Ventilation modes

Standard: ✓ Option: O Not applicable: --

| Mode form | Mode name | Mode | Adult/Ped | Neonatal |
|-------------------------------------|------------------|--|-----------|----------|
| Volume-targeted | APVcmv / (S)CMV+ | Breaths are volume targeted and mandatory. | ✓ | ✓ |
| modes, adaptive pressure controlled | APVsimv / SIMV+ | Volume-targeted mandatory breaths can be alternated with pressure- supported spontaneous breaths. | ✓ | ✓ |
| | VS | Breaths are flow cycled and deliver a set tidal volume to support patient-initiated breaths. | ✓ | ✓ |
| Pressure-controlled modes | PCV+ | All breaths, whether triggered by the patient or the ventilator, are pressure-controlled and mandatory. | ✓ | ✓ |
| | PSIMV+ | Mandatory breaths are pressure controlled. Mandatory breaths can be alternated with pressure-supported spontaneous breaths. | ✓ | ✓ |
| | DuoPAP | Mandatory breaths are pressure controlled. Spontaneous breaths can be triggered at both pressure levels. | 0 | 0 |
| | APRV | Spontaneous breaths can be continuously triggered. The pressure release between the levels contributes to ventilation. | 0 | 0 |
| | SPONT | Every breath is spontaneous, with or without pressure-supported spontaneous breaths. | ✓ | ✓ |
| Intelligent ventilation | ASV | Operator sets %MinVol, PEEP, and Oxygen. Frequency, tidal volume, pressure, and I:E ratio are based on physiological input from the patient. | √ | |
| | INTELLIVENT-ASV | Ventilator management of CO2 elimination and oxygenation is based on clinician-defined target ranges and parameter limits, and physiological input from the patient. The underlying mode is ASV. | 0 | |
| Noninvasive modes | NIV | Every breath is spontaneous. | 0 | 0 |
| | NIV-ST | Every breath is spontaneous as long as the patient is breathing above the set Rate. A backup Rate can be set for mandatory breaths. | 0 | 0 |
| | nCPAP | Demand flow nasal continuous positive airway pressure. | | 0 |
| | nCPAP-PC | Breaths are pressure controlled and mandatory. | | 0 |
| | HiFlowO2 | High flow oxygen therapy. No supported breaths. | 0 | 0 |



Standard configuration and options (in alphabetical order)

Standard: ✓ Option: O Not applicable: --

| unctions | Adult/Ped | Neonatal |
|--|-----------|--|
| Capnography, mainstream (volumetric) and sidestream | 0 | 0 |
| ommunication board: | 0 | 0 |
| O2/Nurse Call/COM1, CO2/SpO2/COM1 ¹ , CO2/SpO2/Humidifier & COM1 ^{1, 2} | | |
| communication protocols. For details, see the Connectivity brochure | 0 | 0 |
| PR ventilation | ✓ | ✓ |
| lynamic Lung | ✓ | |
| vent log (up to 10,000 events with date and time stamp) | ✓ | ✓ |
| low trigger | ✓ | ✓ |
| lamilton Connect Module (connectivity) | 0 | 0 |
| IAMILTON-H900 humidifier integration | 0 | 0 |
| ntelliTrig (leak compensation) | ✓ | ✓ |
| anguages English, US English, Chinese, Croatian, Czech, Danish, Dutch, Finnish, French, German, Greek, Hungarian, ndonesian, Italian, Japanese, Korean, Norwegian, Polish, Portuguese, Romanian, Russian, Serbian, Slovak, panish, Swedish, Turkish, Ukrainian) | √ | ✓ |
| Manual breath/prolonged inspiration | ✓ | ✓ |
| lebulization, pneumatic | ✓ | |
|)2 enrichment | ✓ | ✓ |
| n-screen help | ✓ | ✓ |
| atient group HAMILTON-C1 neo | | ✓ |
| atient group HAMILTON-C1 | ✓ | 0 |
| rint screen | ✓ | ✓ |
| J-45 Ethernet port³ | ✓ | ✓ |
| creen lock | ✓ | ✓ |
| peak valve compatibility | 0 | |
| pO2 monitoring | 0 | 0 |
| tandby with timer | ✓ | ✓ |
| uctioning tool | ✓ | |
| rends/Loops | 0 | 0 |
| ISB port | ✓ | ✓ |
| /ent Status (visual representation of patient's ventilator dependence) | ·····✓ | ······································ |

¹ Applies only to devices with serial number > 6000 ² Only available with the HAMILTON-H900 Y-cable ³ Only available for use if the Hamilton Connect module is activated.

Technical performance

| Description | Specification |
|--------------------------------------|--|
| Automatic expiratory base flow | Adult/Ped: Fixed at 3 l/min |
| | Neonatal: Fixed at 4 I/min |
| Inspiratory pressure | 0 to 60 cmH2O |
| Maximum limited pressure | 60 cmH2O |
| Maximum working pressure | Adult/Ped: 60 cmH2O (total inspiratory pressure); ensured through pressure |
| | limiting |
| | Neonatal: 45 cmH2O (limitation depending on frequency) |
| Maximum inspiratory flow | 260 l/min (120 l/min with 100% O2) |
| Means of inspiratory triggering | Flow trigger control |
| Minimum expiratory time | 20% of cycle time; 0.2 to 0.8 seconds |
| Minute volume capability | Up to 60 l/min |
| Oxygen mixer accuracy | ± (volume fraction of 2.5% + 2.5% of actual reading) |
| Tidal volume | Adult/Ped: 20 to 2000 ml |
| | Neonatal: 2 to 300 ml |
| Preoperational checks | Leak test, flow sensor/circuit/O2 sensor calibration, CO2 sensor zero |
| | calibration ⁴ |
| Display device | Display of settings, alarms, and monitored data |
| | Type: Color TFT |
| | Size: 640 x 480 pixels, 8.4 in (214 mm) diagonal |
| Brightness setting for display | The range is 10% to 100% brightness. By default, Day = 80%; |
| | Night = 40%. |
| Alarm volume (loudness) ⁵ | The range is 1 to 10. The default setting is 5. |
| Sound power level ⁶ | 51 dB(A) ± 3dB(A) |
| Sound pressure level ⁶ | 43 dB(A) ± 3dB(A) |
| | |

⁴ CO2 option required
⁵ Volume at 1 meter distance from ventilator. A setting of 1 = 62 dB(A), 5 = 76 dB(A), and 10 = 85 dB(A), with accuracy of ±3 dB(A).
⁶ Per ISO 80601-2-12.

Standards and approvals

| Classification | Class Ilb, continuously operating according to EC directive 93/42/EEC |
|-------------------------------|--|
| Valid versions | IEC 60601-1:2005/A1:2012, ANSI/AAMI ES60601-1:2005/(R)2012, CAN/CSA-C22.2 No. 60601-1:14, IEC 60601-1-2:2014, ISO 80601-2-12:2011 + Cor.:2011, ISO 80601-2-55:2018, EN ISO 5356-1:2015, ISO 80601-2-61:2017, ISO 80601-2-49:2018 |
| Declaration | The HAMILTON-C1 was developed in accordance with pertinent international standards and FDA guidelines. The ventilator is manufactured within an EN ISO 13485 and EN ISO 9001, Council Directive 93/42/EEC, Annex II, Article 3 certified quality management system. The ventilator meets the Essential Requirements of Council Directive 93/42/EEC, Annex I. |
| Electromagnetic compatibility | According to IEC 60601-1-2:2014 |
| Safety class | Class II, Type B applied part (ventilator breathing system, VBS), type BF applied part CO2 sensor including CO2 module connector; SpO2 sensor including adapter, continuous operation according to IEC 60601-1 |

Pneumatic performance

| High-pressure oxygen inlet | Pressure: | 2.8 to 6 bar / 41 to 87 psi |
|--|----------------------------------|---|
| | Flow: | Maximum of 200 l/min |
| | Connector: | DISS (CGA 1240) or NIST |
| Low-pressure oxygen inlet | Pressure: | Maximum 6 bar / 87 psi |
| | Flow: | ≤ 15 l/min |
| | Connector: | Quick-coupling system, compatible with Colder Products Company (CPC) PMC series |
| Air supply | Integrated blower | |
| Gas mixing system | Delivered flow: | > 260 l/min ±10% against ambient pressure (at sea level) |
| | | • > 200 l/min with 100% oxygen |
| | Delivered pressure: | Adult/Ped: 0 to 60 cmH2O |
| | | Neonatal: 0 to 45 cmH2O |
| | Flow accuracy: | ±10% or ±300 ml/min (whichever is greater) |
| Inspiratory outlet (<i>To patient</i> port) | Connector: | ISO ID15/OD22 conical |
| Expiratory outlet (From patient port) | Connector (on expiratory valve): | ISO ID15/OD22 conical |
| | | |

Electrical specifications

| Input power | 100 to 240 VAC ±10%, 50/60 Hz | | | |
|---|----------------------------------|--|--|--|
| Power consumption 50 VA typical, 150 VA maximum | | | | |
| Battery | Hamilton Medical provides a high | Hamilton Medical provides a high-capacity battery ⁷ . | | |
| | Electrical specifications: | 6.7 Ah, 72 Wh, 50 W typical, 150 W maximum | | |
| | Туре: | Lithium-ion, supplied by Hamilton Medical only | | |
| | Recharge time: | While the ventilator is connected to primary power, approximately 3.25 h to fully recharge one battery. | | |
| | Storage: | -20°C to 60°C, \leq 85% relative humidity. The storage location should be free from vibration, dust, direct sunlight, moisture, and corrosive gases, and with a recommended temperature range $<$ 21°C. | | |
| | | Extended exposure to temperatures above 45°C can degrade battery performance and life. | | |
| | Normal operating time: | Operating times are measured with one fully charged battery, the blower in use, without communication board, and with the following settings: Mode = PCV+, Rate = 10 b/min, Δ Pcontrol = 10 cmH2O, I:E = 1:4, PEEP = 5 cmH2O, Flow trigger = 5 l/min, FiO2 = 40%. | | |
| | | Approximate operating times under these conditions are as follows: | | |
| | | • One battery, display brightness = 80%: 4 h | | |
| | | • One battery, display brightness = 20%: 4.5 h | | |
| | | This operating time applies to new, fully charged Li-ion batteries that have not been exposed to extreme temperatures. The actual operating time depends on battery age and on how the battery is used and recharged. | | |

Graphical patient data

| Graphic type/tab name | Options |
|--|---|
| Waveforms Pressure, Volume, Flow, PCO2 ⁸ , FCO2 ⁸ , Plethysmogram ⁹ , Capnogram ¹⁰ | |
| Intelligent panels | Dynamic Lung ¹¹ , Vent Status, ASV Graph ¹² , INTELLiVENT-ASV Oxygenation and CO2 elimination maps and horizons ¹⁰ |
| Trends | 1-, 6-, 12-, 24-, or 72-h trend data for a selected parameter or combination of parameters |
| Loops | Pressure/Volume, Pressure/Flow, Volume/Flow, Volume/PCO2 ⁸ , Volume/FCO2 ⁸ |

⁷ PN 369108, revision 4 and later. ⁸ CO2 option required ⁹ SpO2 option required ¹⁰ INTELLIVENT-ASV required ¹¹ Only for adult/pediatric patients ¹² Only in ASV mode

Alarms

| Priority | Alarm |
|-----------------|--|
| High priority | Apnea, Apnea time, ExpMinVol high/low, Oxygen high/low, Minute volume high/low, Pressure high/low, High Pressure during Sigh, Pressure not released Flow sensor calibration needed (during ventilation), Check flow sensor tubing, Check flow sensor, Check patient interface, External flow sensor failed, Replace O2 sensor, Oxygen supply failed, Buzzer defective, Loudspeaker defective Disconnection on patient/ventilator side, Exhalation obstructed, Obstruction Options not found, Self test failed, Blower fault, Device temperature high, Vent outlet temperature high Battery low, Battery power loss, Battery totally discharged, Battery temperature high, Battery communication error, Battery defective SpO2:13 SpO2 low |
| Medium priority | High Flow, fTotal high/low, Frequency high/low, Vt high/low, Inspiratory volume limitation, High PEEP, Loss of PEEP, Pulse high/low, Pressure limitation Wrong expiratory valve, Circuit calibration needed, Flow sensor calibration needed, Flip the flow sensor, Check flow sensor for water (Neonatal) Check for blockage, Fan failure, Function key not operational, Performance limited by high altitude, Real-time clock failure, Battery low CO2: ¹⁴ PetCO2 high/low INTELLIVENT-ASV: FiO2 set to 100% due to low SpO2, Oscillation %MinVol, Oscillation PEEP/CPAP, Oxygenation adjustment off, Oxygen control limit exceeded, Ventilation adjustment off SpO2: ¹³ SpO2: Adapter missing, SpO2: Light interference, SpO2: Low perfusion index, SpO2: Poor signal, SpO2: Probe missing, SpO2: Patient disconnected, SpO2: Sensor error, Pl low/high, PVI low/high, Pulse low/high, SpO2 low |
| Low priority | Check Plimit, ASV: Cannot meet the target, Maximum leak compensation, Pressure limit has changed, CPR ON, SpeakValve ON/OFF, Suctioning maneuver, Apnea ventilation/Apnea ventilation ended Flow sensor calibration needed, Preventive maintenance required, Replace HEPA filter, Blower service required, Loss of external power, IRV (inverse ratio ventilation), Release valve defective, Touch not functional, Check settings Battery calibration required, Battery replacement required, Wrong battery, Battery low O2 sensor calibration needed, O2 sensor defective, O2 sensor missing, O2 sensor not system compatible External connections disabled ¹⁵ , JTAG not working, Invalid communication board CO2: ¹⁴ CO2 calibration needed, CO2 sensor defect, CO2 sensor disconnected, CO2 sensor over temperature, CO2 sensor warmup, Check CO2 sampling line, Check CO2 airway adapter, CO2: Poor signal INTELLIVENT-ASV: ¹⁶ Oxygen controller at limit, PetCO2 target range changed, Ventilation controller at limit SpO2: ¹³ SpO2 high |

 ¹³ If the SpO2 option is installed and enabled.
 ¹⁴ If the CO2 option is installed and enabled.
 ¹⁵ If the Hamilton Connect module is installed and enabled.
 ¹⁶ If INTELLIVENT-ASV is installed.

Control settings and ranges

| %MinVol (%) ¹⁸ Apnea backup ETS (%) Flow (l/min) ¹⁹ | 25 to 350 On, Off 5 to 80 2 to 100 ²⁰ | On, Off 5 to 80 |
|---|--|--|
| Apnea backup ETS (%) | 5 to 80 | |
| ETS (%) | | 5 to 80 |
| | 2 to 100 ²⁰ | |
| 11000 (1/111111) | | 2 to 30 |
| l:E ²¹ | 1:9 to 4:1 | 1:9 to 4:1 |
| IBW (kg) <i>(calculated)</i> | 3 to 139 | |
| Oxygen (%) | 21 to 100 | 21 to 100 |
| P high (in APRV) (cmH2O) | 0 to 60 | 0 to 45 |
| P high (in DuoPAP) (cmH2O) | 0 to 60 | 3 to 45 |
| P low (in APRV) (cmH2O) | 0 to 35 | 0 to 25 |
| Pat. height | | |
| (cm) | 30 to 250 | |
| (in) | 12 to 98 | |
| PEEP/CPAP (cmH2O) | 0 to 35 | 3 to 25 |
| Plimit (cmH2O) | 5 to 60 | 5 to 60 |
| P-ramp (ms) ²² | 0 to 2000 | 0 to 600 |
| | ASV, NIV, NIV-ST, SPONT, VS: max = 200 | NIV, NIV-ST, SPONT, nCPAP-PC, VS: max = 200 |
| Rate (b/min) ²³ | 1 to 80 | 1 to 80 |
| | APVcmv, PCV+: 4 to 80 | <i>PSIMV+:</i> 5 to 80 |
| | PSIMV+, NIV-ST: 5 to 80 | APVcmv, PCV+, PSIMV+PSync, nCPAP-PC, NIV-ST, APVsimv + Apnea backup: 10 to 80 |
| Set temp (°C) | INV: 35 to 41 | INV: 35 to 41 |
| | NIV: 30 to 35 | NIV: 30 to 35 |
| | HiFlowO2: 33 to 37 | HiFlowO2: 33 to 37 |
| Sex | Male, Female | |
| Sigh | On, Off | |
| SpeakValve | On, Off | |
| T gradient (°C) | -2 to 3 | -2 to 3 |
| T high ²³ (in APRV and DuoPAP) (s) | 0.1 to 40.0 | 0.1 to 40.0 |
| T low (in APRV) (s) | 0.2 to 40.0 | 0.2 to 40.0 |
| TI (s) ^{21,23} | 0.1 to 12.0 | 0.1 to 12.0 |
| TI max (s) | 0.5 to 3.0 | 0.25 to 3.0 |

¹⁷ Parameter settings and ranges can vary depending on the selected mode.
18 Only in ASV mode.
19 Only for high flow oxygen therapy.
20 In some markets, the maximum possible Flow setting may be limited.
21 In PCV+, (S)CMV, and APVcmv modes, mandatory breath timing can be controlled by using a combination of inspiratory time (TI) and Rate, or by the I:E ratio; set the method in Configuration. All other modes are controlled by using a combination of inspiratory time (TI) and Rate.
22 P-ramp is limited to one-third (1/3) of TI time. Adjustment of TI time can override the P-ramp setting.
23 Startup setting derived from IBW (adult/pediatric), body weight setting (neonatal). Does not apply in ASV mode.

| Parameter (units) | Range Adult/Ped ¹⁷ | Range Neonatal ¹⁷ |
|---|--|--|
| Trigger, flow (l/min) ²⁴ | 0.5 to 20.0 <i>APVcmv</i> , <i>PCV</i> +: 0.5 to 20.0 / Off | 0.1 to 5.0 <i>APVcmv, PCV</i> +: 0.1 to 5.0 / Off |
| Vt (ml) | 20 to 2000 | 2 to 300 |
| Vt/IBW Vt/Weight (ml/kg) ²⁵ | 5 to 12 | 5 to 12 |
| Weight (kg) | | 0.2 to 30.0 |
| ΔPcontrol (cmH2O) ²⁶ | 5 to 60 | 3 to 45 nCPAP-PC: 0 to 45 |
| ΔPinsp (cmH2O) ²⁶ | 3 to 60 | 3 to 45 |
| ΔPsupport (cmH2O) ²⁶ | 0 to 60 | 0 to 45 |

²⁴ Flow trigger is leak compensated.
²⁵ IBW is calculated using height and sex, for adult and pediatric patients. Actual body weight is used for neonates.
²⁶ Δ*Pcontrol*: Control pressure, added to PEEP/CPAP. Δ*Pinsp*: Inspiratory pressure, added to PEEP/CPAP. Δ*Psupport*: Pressure support, added to PEEP/CPAP.

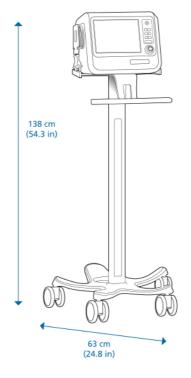
Monitoring parameters

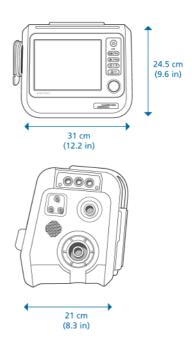
| Parameter (units | 5) | Description |
|------------------|---------------------------------|---|
| Pressure | AutoPEEP (cmH2O) | Unintended positive end-expiratory pressure |
| | PEEP/CPAP (cmH2O) | PEEP (positive end-expiratory pressure) and CPAP (continuous positive airway pressure |
| | Driving pressure, ΔP (cmH2O) | Driving pressure, calculated value reflecting the difference between Pplateau and PEE |
| | ΔPinsp (cmH2O) | Inspiratory pressure |
| | Pmean (cmH2O) | Mean airway pressure |
| | Ppeak (cmH2O) | Peak airway pressure |
| | Pplateau (cmH2O) | Plateau or end-inspiratory pressure |
| | Pprox (cmH2O) | Airway pressure at proximal patient interface |
| Flow | Flow (I/min) | HiFlowO2: The set flow of gas to the patient |
| | | nCPAP: The average flow updated every second |
| | | nCPAP-PC: The average flow during expiration, updated every breath |
| | Insp Flow (peak) (l/min) | Peak inspiratory flow, spontaneous or mandatory |
| | Exp Flow (peak) (l/min) | Peak expiratory flow |
| Volume | ExpMinVol or MinVol NIV (l/min) | Expiratory minute volume |
| | MVSpont or MVSpont NIV (I/min) | Spontaneous expiratory minute volume |
| | VTE or VTE NIV (ml) | Expiratory tidal volume |
| | VTESpont (ml) | Spontaneous expiratory tidal volume |
| | VTI (ml) | Inspiratory tidal volume |
| | VLeak (%) | Leakage percent or total minute volume leakage |
| | MVLeak (I/min) | Leakage percent or total minute volume leakage |
| | Vt/IBW or Vt/Weight (ml/kg) | Tidal volume is calculated by ideal body weight (adult/pediatric patients) or actual bod weight (neonatal patients) |
| Oxygen | Oxygen (%) | Oxygen concentration of the delivered gas |
| | O2 consumption (I/min) | The current oxygen consumption rate |
| Time | CPR timer | MMP during CPR ventilation showing duration of CPR ventilation |
| | l:E | Ratio of the patient's inspiratory time to expiratory time for every breath cycle |
| | fControl (b/min) | Mandatory breath frequency |
| | fSpont (b/min) | Spontaneous breathing frequency |
| | fTotal (b/min) | Total breathing frequency |
| | TI (s) | Inspiratory time |
| | TE (s) | Expiratory time |
| Lung mechanics | Cstat (ml/cmH2O) | Static compliance |
| | P0.1 (cmH2O) | Airway occlusion pressure |
| | PTP (cmH2O*s) | Pressure time product |
| | RCexp (s) | Expiratory time constant |
| | Rinsp (cmH2O / (l/s)) | Inspiratory flow resistance |
| | RSB (1 / (I*min)) | Rapid shallow breathing index |

| Parameter (units) | | Description |
|--------------------------|-------------------|---|
| CO2 | FetCO2 (%) | Fractional end-tidal CO2 concentration |
| | PetCO2 (mmHg) | End-tidal CO2 pressure |
| | slopeCO2 (%CO2/l) | Slope of the alveolar plateau in the PetCO2 curve, indicating the volume/flow status of the lungs |
| | V'alv (l/min) | Alveolar minute ventilation |
| | Vtalv (ml) | Alveolar tidal ventilation |
| | V'CO2 (ml/min) | CO2 elimination |
| | VDaw (ml) | Airway dead space |
| | VDaw/VTE (%) | Airway dead space fraction at the airway opening |
| | VeCO2 (ml) | Exhaled CO2 volume |
| | ViCO2 (ml) | Inspired CO2 volume |
| SpO2 | SpO2 (%) | Oxygen saturation |
| | Pulse (1/min) | Pulse |
| | SpO2/FiO2 (%) | The SpO2/FiO2 ratio (%) is an approximation of the PaO2/FiO2 ratio, which, in contrast to PaO2/FiO2, can be calculated noninvasively and continuously |
| | OSI | Oxygen saturation index |
| | PI (%) | Perfusion index |
| | PVI (%) | Pleth variability index |
| Humidifier ²⁷ | T Y-piece (°C) | Measured temperature at the Y-piece |
| | T humidifier (°C) | Measured temperature at water chamber exit |

²⁷ If HAMILTON-H900 humidifier integration is enabled, and a humidifier is connected and turned on.

Physical characteristics





Weight 4.9 kg (10.8 lb)

16.9 kg (37.3 lb) with trolley

The trolley can accommodate a maximum safe working load 28 of 44 kg (97 lb).

Dimensions See graphic above

Trolley accessories HAMILTON-H900 mounting kit, optional O2 bottle holding system, optional tubing support arm, water bottle

holder, basket

²⁸ The maximum safe working load applies to a stationary, properly load-balanced trolley.

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HAMILTON-C1