

RESmart[®] BPAP 25A Auto Bi-Level

- Full Efficacy Data Reporting
- Innovative tracking technology ensures accurate and comfortable therapy
- 4-25 cmH₂O therapy pressure
- Real time audible alert for tubing/mask leak
- REslex[®] exhalation relief with four comfort settings
- Remote compliance data access with iCode[®]
- SD Card standard
- Integration with cloud based iCode Connect[™]
- · Embedded memory stores last night's full raw data, 365 nights of user data and more than 30 years of therapy records
- Last night's compliance-therapy snapshot to assist patient with compliance
- · Automatic leak and altitude compensation
- Inspiration trigger for automatic start-up
- Quiet operation

Ergonomic Design

- Integrated heated humidifier
- DC 24V powered and infrared controlled humidifier for patient safety and comfort
- · Delay-off feature protects the device from humidity hazard
- Backlit LCD display for operation in dark
- Light user buttons with lockup capability

3B Medical, Inc. 799 Overlook Drive • Winter Haven, FL 33884 (863) 226-6285 · info@3bproducts.com www.3Bproducts.com



B7000 Bundled RESmart[®] BPAP + Humidifier **B5500 RESmart® BPAP Integrated Humidifier** M1020 RESmart[®] BPAP Air Filter (2 Pack) M1019B RESmart[®] Air Filter Cover – BPAP M4510 Humidifier Water Chamber

Dimensions: 8.66 × 7.6 × 4.4" (220 x 194 x 112mm), 12.3" (313 x 194 x 112mm) (with Integrated Heated Humidifier)

Weight: ___ lbs. (2.2 kg), <4.8 lbs. (3 kg) (with Integrated Heated Humidifier)

Environmental

	Operating	Transport and Storage
Temperature	5 to 35°C	-20 to 55°C
Relative Humidity	≤ 80% Non-condensing	≤ 93% Non-condensing
Atmospheric		

Pressure 860 to 1060 hPa 500 to 1060 hPa

Standards Compliance:

IEC 60601-1 General Requirements for Safety of Medical **Electrical Equipment**

IEC 60601-1-2 Electromagnetic Compatibility

ISO 17510 Sleep Apnea Breathing Therapy

ISO 8185 General Requirements for Humidification Systems

Electrical:

AC Power Consumption: 100 - 240V AC, 50/60Hz, Max 95VA Type of Protection Against Electric Shock: Class II Equipment Degree of Protection Against Electric Shock: Type BF Applied Part

Degree of Protection Against Ingress of Water: Drip Proof, IPX1

Mode of Operation: Continuous

BPAP 25 4-2	ssure 5cmH ₂ O 5cmH ₂ O	Functi S, CPA S, CPAP, A	Р	ito Pressure No available
Pressure IPAP EPAP CPAP Ramp Duration Rise Time	4 to 25 4 to 20	minutes	±1 cm ±1 cm ±1 cm ±10% ±25% ³	H ₂ O** H ₂ O** of the setting
* Limited to 25 cmH ₂ O when using the Auto feature (only available under AutoS mode).				

** Dynamic pressure accuracy is ±1 cmH2O measured at the patient end of

the circuit. Static pressure accuracy is ±0.5 cmH2O measured at the patient end of the

**** The range of values corresponds to tenths of seconds (e.g., a setting of 4 indicates a Rise Time of 0.4 seconds). **** Measured at the patient end of circuit.

Disposal: Dispose of the device in accordance with local regulations

Manufacturer: BMC Medical Co., Ltd.

NOTE: specifications subject to change without prior notice.