

Lumis[™] 100

VPAP ST



iBR: Intelligent Backup Rate

Provides back up breaths intelligently



















Restried Lunis 170

Lumis™ 100 VPAP ST

The Lumis[™] 100 VPAP ST device is indicated to provide non-invasive ventilation for patients weighing more than 13 kg with respiratory insufficiency or obstructive sleep apnoea (OSA). It is intended for home and hospital use.

The humidifier is intended for single patient use in the home environment and re-use in a hospital/institutional environment.



iBR: Intelligent Backup Rate

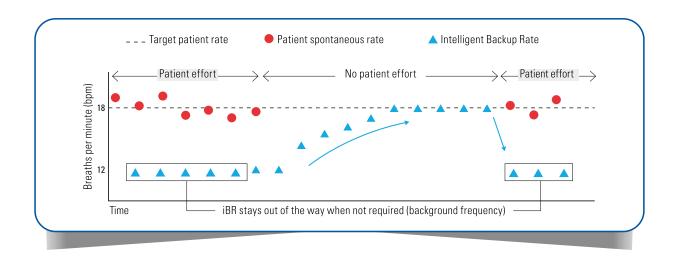
iBR is designed to enhance the conventional approach to backup breaths. It maximises your patient's opportunities for spontaneous breathing by delivering support only when it's needed and tailoring that support to meet their real needs.

What does iBR do?

Provides back up breaths intelligently: provides back up breaths within 2 boundaries: target patient rate and background frequency which is two-thirds of the target rate. iBR will provide backup breaths only if there's an apnoes or lack of effort, but won't provide unnecessary support for pauses caused by cough or sigh.

Reacts swiftly and enhances comfort: designed to safely and quickly return your patient to target rate as iBR provides minimal pressure support and best synchronization at the time of backup breaths. This improves patient's comfort and compliance.

Time saving personalisation: automatically determines the most suitable iBR for your patient based on their spontaneous stable awake rate, giving you peace of mind and ensuring your patient receives personalised ventilation.



ResMed.com/Lumis Awaken your best journey.

ResMed NIV technologies

A key requirement for sucess in NIV is to improve patient comfort, compliance and treatment efficacy though excellent patient ventilator synchrony. Lumis™ 100 VPAP ST features 3 unique ResMed technologies that work together to achieve this synchrony.



Provide patient— ventilator synchrony, even in the presence of significant leak.



Set min and max limits on either side of the patient's ideal inspiratory time.



Optimise settings according to the patient's condition, using five trigger and cycle sensitivity levels.

5 Trigge

Easy-to-read, front-facing colour screen take you back to the top of the menu with a single scroll menu.

EPAP Start EPAP Ramp down

Lumis ramps down the pressure delivered over a 15-minute period

Intuitive and simple to navigate

The user interface on Lumis ventilators has been designed with you and your patients in mind: it's intuitive and simple to navigate. It's easy to view and personalise patient control settlings, as well as gain valuable insight into their progress with a sleep report at the end of every session.

Automatic humidification

When used with a HumidAir heated humidifiers and ClimateLineAir heated tube, Lumis delivers automatic humidification Climate Control Auto. This mode comes pre-set with the temperature and humidity levels designed for optimal comfort, so you can set your patients up to receive the benefits of humidification instantly- no settings to change and no complicated menus to navigate.

Customisation for comfort

For patients who need greater levels of pressure support (e.g. COPD patients), turning the ventilator off at the end of therapy can be quite abrupt. Lumis's optional Ramp Down feature gradually reduces the pressure support delivered to help ease patients into spontaneous breathing.

Reference: Lumis series clinical guide

Product code

Lumis™100 VPAP ST 28126

Technical specificationsUnits are expressed in cm H20 and hPa.1cm H20 is equal to 0.98 hPa.

90W power supply unit AC input range: DC output: Typical power consumption: Peak power consumption:	100-240V, 50-60Hz 1.0-1.5A, Class II 115V, 400Hz 1.5A, Class II (nominal for aircraft use) 24V3.75A 53W (57VA) 104W (108VA)
Environmental conditions Operating temperature:	+5°C to +35°C Note: The airflow for breathing produced by this therapy device can be higher than the temperature of the room. Under extreme ambient temperature conditions (40°C) the device remains safe.
Operating humidity: Operating altitude: Storage and transport temperature: Storage and transport humidity:	10 to 95% relative humidity, non-condensing Sea level to 8,500'(2,591 m); air pressure range 1013 hPa to 738 hPa -20°C to +60°C 5 to 95% relative humidity, non-condensing

Classification: EN60601-1:2006/A1:2013

Class II (double insulation), Type BF, Ingress protection IP22.

Physical - device and humidifier	
Dimensions (H x W x D):	116 mm x 255 mm x 150 mm
Air outlet (complies with ISO 5356-1:2015):	22 mm
Weight (device and cleanable humidifier):	1268 g
Housing construction:	Flame retardant engineering thermoplastic
Water capacity:	To maximum fill line 380 mL
Cleanable humidifier - material:	Injection moulded plastic, stainless steel and silicone seal
Air filter	
Standard:	Material: Polyester non woven fibre
	Average arrestance: >75% for ~7micron dust
Hypoallergenic:	Material: Acrylic and polypropylene fibres in a polypropylene
	carrier Efficiency: >98% for ~7-8 micron dust; >80% for ~0.5 micro
	dust
Aircraft use	
ResMed confirms that device meets the Federal Aviation Adminis	ration (FAA) requirements (RTCA/DO-160, section 21, category M)
for all phases of air travel.	
Operating pressure range	
S, ST, T, PAC	2 to 25 cm H20 (2 to 25 hPa)
CPAP	4 to 20 cm H20 (4 to 20 hPa)
Supplemental oxygen	
Maximum flow:	15 L/min (S, ST, T, PAC, CPAP)

Disclaimer: This device should only be used under the supervision of a registered medical practitioner. The information herein is intended solely for healthcare professionals and does not constitute medical advice. A healthcare professional must always rely on his/her own professional clinical judgment when deciding whether to use a particular product to treat a particular patient. ResMed does not provide medical advice and recommends that healthcare professionals be trained in the use of any particular product before use on a patient.





