

## RT269 | SPECIFICATIONS

Infant Ventilator Circuit Dual Heated with MR290 Autofeed Chamber for SLE6000



### CIRCUIT COMPONENTS AND COMPOSITION

<b>Pack Components</b>	1.1 m heated inspiratory limb, 0.3 m incubator extension, 1.6 m heated expiratory tube with Evaqua™ 2 technology, 0.6 m humidifier connection tube with SLE6000 connector and nitric oxide port, swivel wye-piece, MR290V autofeed humidification chamber, pressure line, SLE restrictor, adaptor kit, label kit change out

<b>Materials</b>	Polypropylene, polycarbonate, thermoplastic elastomer, linear low-density polyethylene, low-density polyethylene, high-density polyethylene, polyether ester elastomer, silicone, nylon, brass, styrene resin, cyanoacrylate, DEHP-free PVC. Not made with natural rubber latex.
<b>Quantity</b>	10 circuits per carton
<b>Carton Dimensions and Weight</b>	Length: 530 mm; Width: 210 mm; Height: 400 mm; Weight: 3.98 kg
<b>Carton Material</b>	Cardboard box
<b>Manufacturing Mode</b>	Produced in a Controlled Working Environment
<b>Disposal</b>	Dispose of product according to Hospital protocol. User may be exposed to breathing tract fluids during disposal.

### PERFORMANCE SPECS

<b>Resistance to Flow</b>	Inspiratory limb: 0.09 ± 0.04 cmH <sub>2</sub> O Expiratory limb: 0.07 ± 0.04 cmH <sub>2</sub> O @ 2.5 L/min
<b>Flow Rate</b>	4 - 15 L/min
<b>Circuit Length</b>	Inspiratory: 1.6 m Expiratory: 1.6 m
<b>Minimum Tube Internal Diameter</b>	10.2 mm
<b>Ambient Range</b>	20 - 26 °C / 68-79 °F
<b>Compliance</b>	1.30 ± 0.15 mL/cmH <sub>2</sub> O @ 60 cmH <sub>2</sub> O
<b>Humidifier Compatibility</b>	MR850 humidifier
<b>Humidifier Mode</b>	Invasive
<b>Compressible Volume</b>	762 mL
<b>Duration of Use</b>	7 days
<b>Use</b>	Single patient
<b>Interface Connections</b>	ISO 5356-1 Conical Connectors
<b>Shelf Life</b>	5 years
<b>Gas Leakage</b>	< 30 mL/min @ 60 cmH <sub>2</sub> O
<b>Storage Temperature</b>	Lower temp limit: -10 °C Upper temp limit: 50 °C

CHAMBER	MR290
Interface Connections	ISO 5356-1 Conical connector (22 mm Male)
Maximum Chamber Operating Pressure	8 kPa
Maximum Peak Flow	180 L/min for 30 secs
Gas Leakage	< 10 mL/min @ 8 kPa
Materials	ABS, Polystyrene, Polyethylene, Thermoplastic Elastomer, Aluminium, Ink, Polypropylene, Adhesive, Polycarbonate, Silicone, DEHP-free PVC
Compliance	0.4 mL/cmH <sub>2</sub> O
Compressible Volume	280 mL
Resistance to Flow	At 60 L/min: 0.52 cmH <sub>2</sub> O
REGULATORY	
Classification	AU IIa; EU IIa; CA 2; USA II. For more Regulatory information visit: <a href="http://www.fphcare.com/regulatory">www.fphcare.com/regulatory</a>
Country of Origin	New Zealand
Intended Use	This device is intended for the delivery of breathing gases to infant patients requiring respiratory support.
Notified Body Identification Number	TÜV SÜD Product Services GmbH CE0123
GTIN Number	09420012458223 (EA) 09420012458230 (PAC)
UNSPSC Number	42272224
GDMN Code	37706
Biocompatibility	Meets standards: ISO 10993-3, ISO 10993-5, ISO 10993-6, ISO 10993-10,