

RT266 | SPECIFICATIONS

Infant Ventilator Circuit (0.3 to 4 L/min) Dual Heated with MR290 Autofeed Chamber



CIRCUIT COMPONENTS AND COMPOSITION

Pack Components	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 nitric oxide port, swivel wye-piece, MR290V autofeed humidification chamber, pressure line, Dräger adaptors, label kit change out

Materials	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.
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Quantity	10 circuits per carton
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Carton Dimensions and Weight	Length: 531 mm; Width: 204 mm; Height: 392 mm; Weight: 3.97 kg
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Carton Material	Cardboard box
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Manufacturing Mode	Produced in a Controlled Working Environment
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Disposal	Incineration, or According to Hospital Protocol
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PERFORMANCE SPECS

Resistance to Flow (with and without accessories)	Inspiratory limb: 0.10 ± 0.05 cmH ₂ O Expiratory limb: 0.07 ± 0.05 cmH ₂ O @ 2.5 L/min
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Flow Rate	0.8 - 4 L/min at Ambient Temperature 20 - 26 °C 0.3 - 4 L/min at Ambient Temperature 23 °C
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Circuit Length	Inspiratory: 1.6m Expiratory: 1.6m
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Minimum Tube Internal Diameter	10.2 mm
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Ambient Range	20-26 °C / 68-79 °F
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Compliance (with and without accessories)	1.12 ± 0.11 mL/cmH ₂ O @ 60 cmH ₂ O
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Humidifier Compatibility	MR850 humidifier
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Humidifier Mode	Invasive
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Compressible Volume	758 mL
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Duration of Use	7 days
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Use	Single patient
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Recommended Gas Source	Air/Oxygen
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Interface Connections	ISO 5356-1 Conical Connectors
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Shelf Life	5 years
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Gas Leakage	<30 mL/min @ 60 cmH ₂ O
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Storage Temperature	Lower temp limit: -10 °C Upper temp limit: 50 °C
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Tidal Volume	<185 mL
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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 ₂ O
Compressible Volume	280 mL
Resistance to Flow	At 60 L/min: 0.52 cmH ₂ O
REGULATORY	
Classification	AU IIa; EU IIa; CA II; USA II. For more Regulatory information visit: www.fphcare.com/regulatory
Country of Origin	New Zealand or Mexico
Intended Use	This device is intended to deliver respiratory gases to infant patients who are dependent on mechanical respiratory support.
Notified Body Identification Number	TÜV SÜD Product Services GmbH CE0123
GTIN Number	09420012431103 (EA) 09420012424372 (PAC)
UNSPSC Number	42272224
GDMN Code	37706
Biocompatibility	Meets standards: ISO 10993-3, ISO 10993-5, ISO 10993-6, ISO 10993-10