











Blender Transition Kits SPECIFICATIONS

PRODUCT	 OJR410B	 OJR412B	 OJR414B	 OJR416B	 OJR418B
NASAL CANNULA SIZE	XS	S	M	L	XL
Description	F&P Optiflow™ Junior 2 Blender Transition Kit – XS	F&P Optiflow™ Junior 2 Blender Transition Kit – S	F&P Optiflow™ Junior 2 Blender Transition Kit – M	F&P Optiflow™ Junior 2 Blender Transition Kit – L	F&P Optiflow™ Junior 2 Blender Transition Kit – XL
Weight (per kit)	31.2 g	31.6 g	33.8 g	38 g	38.3 g
FLOW RATES* (L/min) - MR850					
 BC151 BC161 BC153 BC163	4-8	4-9	4-10	4-15	4-15
COMPATIBLE SPARES					
F&P Wigglepad™ 2	 WJR110		 WJR112		
PRODUCT SPECIFICATIONS					
Compatible circuits	BC151, BC161, BC153, BC163				
Connection	Easy-click connector, adaptor				
Compatible humidifier	MR850 (in invasive mode)				
Quantity	Box of 5				
Box components	F&P Optiflow Junior nasal cannula, proprietary adaptor, OJR215 pressure relief manifold, user instructions				

* Flow rates above describe technical capability of the product when used at sea level.
Ensure clinical judgement is used when prescribing flow rates.

Blender Transition Kits | SPECIFICATIONS

PERFORMANCE SPECIFICATIONS	
Ambient range	18-26 °C / 64-79 °F
Usage	Single patient use
Duration of use	7 days
Recommended gas source	Medical gas
Shelf life	3 years
Nominal relief pressure	40 cmH ₂ O
COMPONENTS AND COMPOSITION	
Cannula predominant materials	Thermoplastic Elastomer; Hydrocolloid; ABS; Stainless steel
Adaptor predominant materials	Polypropylene
Manifold predominant materials	Thermoplastic Elastomer White; ABS; Stainless steel; Silicon rubber
Materials not present	Not manufactured with natural rubber latex, PVC or Phthalates (DEHP, DBP, BBP)
Manufacturing mode	Produced in a controlled working environment
Disposal	Incineration or according to hospital protocol for cannula, manifold and adapter; Clamshell packaging and label recyclable PET
REGULATORY	
Classification	AU-IIa, EU-IIa, Canada-II, US-II
Country of origin	New Zealand
Notified body	TÜV SÜD Product Services GmbH CE0123
Standard	Medical tapers used comply with ISO 5356-1

Please note that the information in this specifications sheet (including product information and images) is summarized and provided for illustrative purposes only. Please refer to the relevant user instructions for more information and confirm details with your local Fisher & Paykel Healthcare representative prior to placing an order. Information subject to change without notice.