

Nasal Cannula | MR850 | SPECIFICATIONS



| PRODUCT | OJR410  | OJR412  | OJR414  | OJR416  | OJR418  |
|-----------------------------------------------------------------------------------------|----------------------------------------------------------------------------------------------------------|------------------------------------------------------------------------------------------|--------------------------------------------------------------------------------------------|--------------------------------------------------------------------------------------------|--------------------------------------------------------------------------------------------|
| Nasal cannula size | XS | S | M | L | XL |
| Cannula weight | 6.7 g | 7.1 g | 9.3 g | 13.5 g | 13.8 g |
| MAXIMUM FLOW RATES* (L/min) | | | | | |
| MR850  | 0.5 - 8 | 0.5 - 9 | 0.5 - 10 | 0.5 - 23 | 0.5 - 25 |
| COMPATIBLE SPARES | | | | | |
| F&P Wigglepad™ 2 |  WJR110 | |  WJR112 | | |
| PRODUCT SPECIFICATIONS | | | | | |
| Compatible with | RT330 Circuit, BC115-05 Pressure Relief Manifold | | | | |
| Connection | Easy-click connector | | | | |
| Compatible humidifier | MR850 (in invasive mode) | | | | |
| Quantity | Box of 20 | | | | |
| Box components | F&P Optiflow Junior 2 nasal cannula, user instructions | | | | |
| PERFORMANCE SPECIFICATIONS | | | | | |
| Ambient range °C | 18-26 °C / 64-79 °F | | | | |
| Humidifier mode | Invasive | | | | |
| Usage | Single patient use; Maximum 7 days | | | | |
| Duration of use | 7 days | | | | |
| Recommended gas source | Medical gas | | | | |
| COMPONENTS AND COMPOSITION | | | | | |
| Predominant materials | Thermoplastic Elastomer; Hydrocolloid; ABS; Stainless steel | | | | |
| Materials not present | Not manufactured with natural rubber latex, PVC or Phthalates (DEHP, DBP, BBP) | | | | |
| Manufacturing mode | Non-invasive device; Produced in a controlled working environment | | | | |
| Disposal | Incineration or according to hospital protocol for cannula; Clamshell packaging and label recyclable PET | | | | |
| REGULATORY | | | | | |
| Classification | AU-IIa, EU-IIa, Canada-II | | | | |
| Country of origin | New Zealand | | | | |
| Notified body | TÜV SÜD Product Services GmbH CE0123 | | | | |

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

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.