PROFEEL® **NON-LATEX SURGICAL GLOVES**

DHD™ Polyisoprene Powder Free



Damp Hand Donning - DHD [™] • Polyisoprene: Non- Latex • Powder Free





A new era of synthetic gloves technology

- This latex-free and powder-free glove represents the latest innovation in synthetic glove technology.
- It is manufactured using a proprietary synthetic polymer that is chemically identical to natural rubber and incorporates our state of the art damp hand donning technology.
- With WRP's new PROFEEL® DHD™ Polyisoprene Surgical Gloves, you can enjoy the fit, feel and comfort of natural rubber latex in a 100% latex free product.
- Powder free feature eliminates powder-induced irritation and powder stains
- Does not contain any Natural Rubber Latex benefiting individuals with Type I Natural Rubber Protein Allergies
- Free from any unpleasant odours or smells
- Environmentally friendly and biodegradable
- A proprietary manufacturing process that ensures the exterior of our gloves consistently match the smooth grip of the human hand
- Damp Hand Donning (DHD)- The inner surface of the glove is coated with a proprietary polymer to enhance damp hand donning
- Excellent levels of tear resistance and barrier integrity not found in other synthetic material
- Beaded cuff prevents tearing while donning and helps prevent roll back

PRODUCT INFORMATION

Article No. : WRP SG PI PF 135

Style No. : WRP SG PI PF 135 - PI DHD

Material : Polyisoprene Rubber

Coating : Damp Hand Donning (DHD)

 Weight per pair, size 7.5, g
 : 24.3 ± 0.6

 Sizes
 : 5.5 - 9.0

 Finger Thickness
 : 0.20 ± 0.02

 Length (mm)
 : 290 ± 10

 Colours
 : White

Glove Size	Palm Width (mm)
5 ½	72 ± 4
6	77 ± 5
6½	83 ± 5
7	89 ± 5
7 ½	95 ± 5
8	102 ± 6
8 ½	108 ± 6
9	114 ± 6

PRODUCT CONFORMANCE:

Medical Device: In compliance with European Medical Device Directive 93/42/EEC (Class IIa), EN 455: Part 1/2/3/4, EU Regulation 2016/425, EN 420:2003 +A1 2009, EN ISO 374-1:2016, EN ISO 374-2:2014, EN ISO 374-4:2013, EN 374-5:2016, EN 16523-1:2015, ASTM D3577, FDA 510 K

QUALITY ASSURANCE:

Manufacturing process is in compliance with US FDA Quality System Regulation (QSR), ISO 9001 Quality Mangement System and BS EN ISO 13485 Quality Systems



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