

Powder Free Synthetic Examination Gloves

DERMAGRIP Powder Free Nitrile, Ultra

DERMAGRIP Powder Free Nitrile, Ultra LT

DERMAGRIP SAFEDON Powder Free Nitrile, Ultra

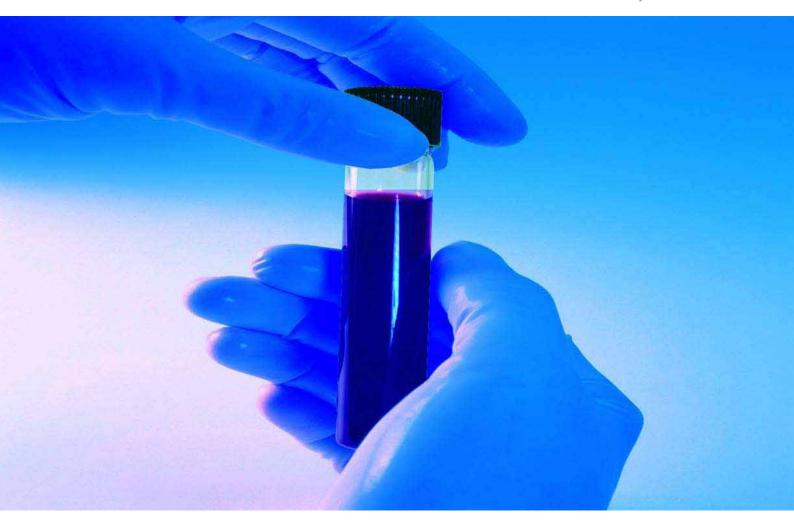
DERMAGRIP SAFEDON Powder Free Nitrile, Ultra LT

DERMAGRIP Powder Free Nitrile

DERMAGRIP Powder Free Nitrile, Sterile

DERMAGRIP Powder Free Nitrile Extended Cuff

DERMAGRIP Powder Free Vinyl



Latex Free

Powder Free

Enhanced Comfort





No Need to Compromise Comfort for Protection

Medical gloves provide protection from unwanted and dangerous substances for both healthcare professionals and patients. **DERMAGRIP®** Ultra Nitrile Examination Gloves meet and exceed international product standards to provide uncompromising protection to its

Dermagrip® Ultra Nitrile Examination Gloves are formulated with 100% nitrile (Acrylonitrile-butadiene) providing an alternative solution for individuals who are allergic to natural rubber latex.



PRODUCT INFORMATION

Type: Powder free and non-sterile

Primary Material: Acrylonitrile-butadiene

Dimensions:

Glove Size	Reorder#	Palm Width (mm)	Length (mm)
Extra-small	D1500-18/19	≤ 80	240 min.
Small	D1501-18/19	80 ± 10	240 min.
Medium	D1502-18/19	95 ± 10	240 min.
Large	D1503-18/19	110 ± 10	240 min.
Extra-large	D1504-18/19	≥ 80	240 min.

Thickness:

Location of Thickness Measurements		Single Wall (mm)
Finge	r (at 15 mm from the extreme tip)	0.09 min.
Palm	(at center of palm)	0.06 min.
Cuff	(at 25 mm from cuff end)	0.05 min.

Physical Properties:

Parameters	Before Aging	After Aging
Tensile Strength (MPa)	14 min.	14 min.
EN 455-2 Force At Break (N)	6 min.	6 min.

Powder: No powder lubricant added

Color: Blue (PMS 285 U)

Design and Feature: Ambidextrous, textured surface at fingers and beaded

Packing: 100 gloves per dispenser, 10 dispensers per carton OR 200 gloves per dispenser, 10 dispensers per carton

Pre-shipment Quality Inspection:

Criteria	Insp. Level	AQL
Dimension	N=13	Median
Physical Properties	N=13	Median
1000ml Water Leak Test	G-I	1.0
Visual Inspection for Major Defects	G-I	2.5
Visual Inspection for Minor Defects	G-I	4.0
Powder Free Residue	N=5	N/A

Product Conformance:

- Medical Device: in compliance with European Medical Device Directive 93/42/EEC (CE Class I).
- EN 455 Parts 1, 2, 3 and 4.
- Personal Protective Equipment of Complex Design Category III, in compliance with 89/686/EEC, type tested to EN 420:2003+A1:2009, EN 374-2:2003, EN 374-3:2003 & EN 388:2003, CE 0086.
- ASTM D6319.

Quality Assurance: Manufacturing process is in compliance with US FDA Quality System Regulation (QSR) and ISO9001 and ISO13485 Quality Systems



WRP Asia Pacific Sdn Bhd 147817V

Lot 1, Jalan 3, Kawasan Perusahaan Bandar Baru Salak Tinggi 43900 Sepang, Selangor Darul Ehsan **MALAYSIA**

Office +60-3-8706 1486 Facsimile +60-3-8706 1557

MK5 8PP, ÚK

COUNCIL DIRECTIVE 93/42/EEC
COUNCIL DIRECTIVE 89/686/EEC

10000

customer_wrp@wrpworld.com

EC REP REMESCO Handelges.m.b.H.

Grinzinger Allee 5/25, 1190 Vienna **AUSTRIA**

Office +43-1-328-50-88 Facsimile +43-1-328 29-49 office@remesco.com Distributed by: