

DATA SHEET

DERMAGRIP®

Nitrile Examination Gloves **Ultra LS**

Powder Free & Latex Free Examination Glove, non-sterile

Reorder#/Size		XS : D1100-24 S : D1101-24 M : D1102-24 L : D1103-24 XL : D1104-24	
Design and Features		Ambidextrous, textured finger-tips and beaded cuff	
Type		Powder free and non-sterile examination glove	
Material		100% Nitrile (Acrylonitrile butadiene)	
Colour		Blue	
Surface Treatment		Chlorination	
Powder Residues (mg/glove)		≤ 2 (powder free)	
Dimensions – Palm width (mm)		<u>Palm width (mm)</u> XS : <80 S : 80 ± 10 M : 95 ± 10 L : 110 ± 10 XL : ≥110	<u>Length (mm)</u> XS : Min. 240 S : Min. 240 M : Min. 240 L : Min. 240 XL : Min. 240
Single wall thickness (mm)	Finger Palm Cuff	<u>Specification</u> min. 0.05 min. 0.05 min. 0.05	<u>Typical Value</u> 0.10 0.07 0.06
Physical Properties	Tensile Strength (MPa) Elongation (%) Force at break (N) – specification Typical Force at break (N)	Before aging : min. 14 : min. 500 : min. 6 : 7.0	After aging : min. 14 : min. 400 : min. 6 : 7.0
Packing mode		200 gloves (by weight) per dispenser (XL: 180 gloves by weight) 10 dispensers per carton	
Glove marking		No marking	
Lot number		Lot number structure: YMMPPPPSS Y = year of packing P = internal reference no. MM = month of packing SS = size	
Shelf life		5 years from date of manufacture	
Pre-shipment Quality Inspection	Dimensions Physical Properties 1000ml Water Leak Major Visual Inspection Minor Visual Inspection Powder Residues	N=13 (EN 455-2) N=13 (EN 455-2) G-I, AQL 1.5 (EN 455-1) G-I, AQL 2.5 G-I, AQL 4.0 N=5 (EN 455-3)	
Product Conformance		- Medical Device in compliance with European Medical Device Regulation (EU) 2017/745 (CE Class I): EN 455 Parts 1, 2, 3 & 4 - Personal Protection Equipment in compliance with European Regulation (EU) 2016/425 (category III, type tested to EN ISO 21420:2020, EN ISO 374-1:2016 Type B, EN ISO 374-5:2016), CE 2777 - Regulation 2016/425 on personal protective equipment, as amended to apply in GB (in UK), UKCA 0086 - ASTM D6319	
Quality Assurance		- ISO 9001 - ISO 13485 Quality Management System - ISO 14001 Environmental Management System - US FDA Quality System Regulation (QSR)	

Note: The above information is a guideline of typical performance values and characteristics of the product and not to be used as actual product specifications.