



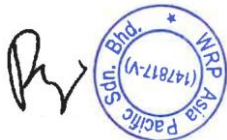
SPECIFICATION SHEET

DERMAGRIP[®] Chemo Protection Gloves

NR LATEX HIGH PROTECTION EXAMINATION GLOVES, POWDER FREE, STERILE

Size	Reorder #	Dimension		
		Target weight (g/pc)	Palm width (mm)	Length (mm)
Small (6.5)	D3165-19	23.0 ± 3	83 ± 5	Min.290
Medium (7.5)	D3175-19	25.0 ± 3	95 ± 5	Min.290
Large (8.5)	D3185-19	27.0 ± 3	108 ± 6	Min.290
Extra Large (9.0)	D3190-19	29.0 ± 3	114 ± 6	Min.290
		<i>*non-release criteria</i>		
Product Part No.		312xx.109422020		
Design and Feature		Hand specific, straight fingers and beaded cuff		
Type		Powder free, extra-thick, hand specific and sterile exam gloves		
Material		Natural rubber latex		
Colour		Blue (PMS 307U)		
Surface Treatment		Chlorination (i.e., SBS-HP)		
Protein Content		This latex glove contains 50 microgram or less of total extractable protein per gram		
Powder Residue (mg/glove)		≤ 2		
Single-wall Thickness (mm) <i>*All sizes</i>	Finger	<u>Specification</u>		
	Palm	0.48 ± 0.03		
	Cuff	0.41 ± 0.03		
		0.28 ± 0.03		
Physical Properties	Tensile Strength (MPa) Ultimate Elongation (%) Stress at 500% (MPa) Force at Break (N)	Unaged	Aged	
		Min. 18	Min. 14	
		Min. 650	Min. 500	
		Max. 5.5 ≥ 6	N/A ≥ 6	
Packing Mode		1 pair of gloves per inner wallet 1 inner wallet per inner polybag 10 inner polybag per outer polybag 10 outer polybag per carton		
Glove Marking		The cuff of each glove is marked with 'Hand & Size'		
Lot # Identification on Finished Goods		Lot Number Structure : YMMPPPPSS (9 Digits) Y = Year of packing (e.g. 3 for year 2023, 4 for year 2024, etc.) MM = Month of packing (e.g. 09 for Sept., 10 for Oct., etc.) PPPP = WRP's Packing Work Order# (PWO) SS = Size (65=6.5, 75=7.5, 85=8.5 and 90=9.0)		
Shelf Life Claim		3 years upon manufacturing date		
Pre-shipment Inspection <i>*Single-Normal Sampling Plan</i>	Dimension	N=13, Median; S-2, AQL 4.0		
	Physical Properties	N=13, Median; S-2, AQL 4.0		
	1000ml Water Leak	G-I, AQL 0.65		
	Protein Content	N=8; N=3		
Powder Residue	N=5			
Major Visual Inspection	G-I, AQL 1.5			
Minor Visual Inspection	G-I, AQL 4.0			
Product Conformance		<ul style="list-style-type: none"> Medical Device: in compliance with MDR (EU)2017/745 (CE Class I) Personal Protective Equipment of Complex Design Category III, in compliance with European Regulation (EU) 2016/425, Regulation 2016/425 on personal protective equipment, as amended to apply in GB, tested to EN ISO 21420:2020, EN ISO 374-1:2016+A1:2018 Type B, EN ISO 374-2:2019, EN ISO 374-4:2019, EN ISO 374-5:2016, EN 16523-1:2015+A1:2018, CE2797 EN 455 Part 1/2/3/4 ASTM D3578, ASTM D6978 (Chemotherapy testing) Health Canada Compliance 		
Quality Assurance		<ul style="list-style-type: none"> MDSAP ISO 13485:2016 Quality Management System ISO 9001:2015 Quality Management System EN ISO 13485:2016 ISO 14001:2015, Environmental Management System 		

WRP ASIA PACIFIC SDN. BHD.



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