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Technical Data Sheet

Signature Latex OR

Sterile Low-Protein Latex Powder-Free Surgical Gloves with Synthetic Polymer Coating

Reference & Size

MSG7255	MSG7260	MSG7265	MSG7270	MSG7275	MSG7280	MSG7285	MSG7290
5,5	6,0	6,5	7,0	7,5	8,0	8,5	9,0

Primary Material

Natural Rubber Latex with Synthetic Polymer Coating

Powder free in accordance with EN455-3

Caution: This product contains natural rubber latex, which may cause allergic reactions, including anaphylactic responses.

Donning Agent

Synthetic Polymer Coating (Inner surface coated for dry and damp hand donning)

Color

Cream (Provides contrast when using a dark colored underglove)

Grip

Micro textured

Former (Mold) Design

Anatomical to replicate curved hand shape and minimize hand fatigue

Cuff Design

Tapered, beaded cuff design to prevent rolldown

Chemical Additives (Accelerators)

Zinc Diethyldithiocarbamate (ZDC)

Leachable Protein

(per EN455-3 using ASTM D5712 (Modified Lowry Protein Method)

Below 50 micrograms/gram of total extractable protein
Caution: Safe use of these gloves by or on Latex-sensitized individuals has not been established

Thickness

(per ASTM D3577 ≥ 0,10 mm)

Fingertip	0.22 mm
Palm	0.19 mm
Cuff	0.17 mm

Document Number: EU1-00004-F-00025 Document Title: FINT.441 - Technical sheet Revision: 3

CE
0344 Medical Device
2777 Personal Protective Equipment

Retention period: End of life and support p

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Length (mm) & Width (mm) Per EN455-2	<table><tr><td>MSG7255</td><td>MSG7260</td><td>MSG7265</td><td>MSG7270</td><td>MSG7275</td><td>MSG7280</td><td>MSG7285</td><td>MSG7290</td></tr><tr><td>289</td><td>290</td><td>290</td><td>290</td><td>289</td><td>290</td><td>289</td><td>290</td></tr><tr><td>72</td><td>78</td><td>85</td><td>92</td><td>97</td><td>104</td><td>110</td><td>114</td></tr></table>
Force @ Break Before Challenge Testing (per EN455-2 ≥ 9 N)	17.9 N
Force @ Break After Challenge Testing (per EN455-2 ≥ 9 N, 7 days 70°C in an oven)	16.5 N
Freedom from Holes (per EN 455 AQL 1.5)	0,65 AQL Before Packaging 0,65 AQL Final Inspection
Viral Penetration	Tested and passed, in accordance with ISO16604 (EN ISO 374-5:2016)
Chemical Resistance	The resistance to chemicals has been assessed in accordance with EN 16523-1 Results and recommendations for use with chemicals can be obtained on request
Sterilization	E-Beam Radiation, Sterility Assurance Level 10^{-6}
Expiration Date	24 Months from Date of Manufacture Manufacture and Expiration Dates are printed on packaging (YYYY-MM format)
Packaging	Polyethylene peel pouch Full length pack 50 pairs per dispenser box / 4 boxes per carton / 200 pairs per carton
Regulations and Quality Standards	Guilin HBM Health Protections Inc. manufacturing locations are certified to EN ISO 13485 by DEKRA Product meets requirements of the EU Medical Device Directive (93/42/EEC) Product meets requirements of European harmonized standards EN 455-1, -2, -3 and -4
PPE Certification	Under the requirements of Personal Protective Equipment Regulation (EU)2016/425 Category III. Complies with standards EN ISO 21420, EN ISO 374-1, EN ISO 374-2, EN 16523-1, EN ISO 374-4, EN ISO 374-5 and ISO 16604.
Storage Recommendations	Protect from freezing, avoid excessive heat. Shield from direct sunlight, fluorescent lighting, x-rays, moisture and ozone.
Country of Origin	China
Manufacturer's Address	Guillin HBM Health Protections, Inc. Add. No. 1-2 Shuijing East Road, Economic and Technological Development Area Guillin, China

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