

Technical Sheet

Sensicare® PI Evolution

Sterile Synthetic Polyisoprene formulation Powder-Free Surgical Gloves with Synthetic Polymer Coating

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Reference & Size	MSG9555 5,5	MSG9560 6,0	MSG9565 6,5	MSG9570 7,0	MSG9575 7,5	MSG9580 8,0	MSG9585 8,5	MSG9590 9,0	
Primary Material	Powder-Free Synthetic Polyisoprene formulation with Synthetic Polymer Coating 0.26 milligrams/glove of powder in accordance with ASTM D6124 and ISO 21171								
Donning Agent	Synthetic Polymer Coating (Inner surface coated for dry and damp hand donning)								
Color	White								
Grip	Smooth								
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 (ZDEC) and Zinc Dibutyldithiocarbamate (ZDBC) Residual chemical levels below detectable level according to UPB/P/003a test method								
Leachable Protein (per EN455-3 using ASTM D5712 (Modified Lowry Protein Method)	No natural rubber latex proteins and allergens.								
Thickness (per ASTM D3577 ≥ 0,10 mm)	Finger Tip 0,23 mm Palm 0,22 mm Cuff 0,17 mm								
Cuff Length (per EN455-2 ≥ 270 mm size 7.5)	291 mm								
Force @ Break Before Accelerated Aging (per EN455-2 ≥ 9 N)	18.3 N								
Force @ Break After Accelerated Aging (per EN455-2 ≥ 9 N, 7 days 70°C in an oven)	14.8 N								
Freedom from Holes (per EN 455 AQL 1.5)	0.65 AQL Final Inspection								
Viral Penetration	Tested and passed, in accordance with ASTM F 1671								
Chemical Resistance	The resistance to some chemicals has been assessed in accordance with EN 374-3 Results and recommendations for use with chemicals can be obtained on request								
Sterilization	Gamma Irradiation, Sterility Assurance Level 10-6								
Expiration Date	35 Months from Date of Manufacture								

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Manufacture and Expiration Dates are printed on packaging (YYYY-MM format)

Revision Date: 09/11/15



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Polyethylene peel pouch material protects product during transport and storage from moisture and ozone and prevents tearing when opening to maintain a sterile Packaging environment

Long pack configuration for easier aseptic donning.

50 pairs per dispenser box / 4 boxes per carton / 200 pairs per carton

Medline manufacturing locations are certified to EN ISO 13485 Regulations and Quality Standards

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

Protect from freezing. Avoid excessive heat. Keep dry. Product should be shielded from Storage Recommendations direct sunlight, fluorescent lighting, X-rays, moisture and ozone. Do not store in temperatures above 40°C.

Country of Origin Malaysia

Legal Manufacturer Medline Industries, Inc.

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