

A member of Top Glove Group: The World's Largest Manufacturer of Gloves

FACTORY 36 : 2, Jalan 8, Pengkalan Chepa 2 Industrial Zone, 16100 Kota Bharu, Kelantan D.N., Malaysia.

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BRAND NAME:

dermagel micropren

PRODUCT DESCRIPTION		PHYSICAL PROPERTIES								
Type of the glove	Sterile powder-free surgical and protective gloves for single use	Size	6.0	6.5	7.0	7.5	8.0	8.5	9.0	
Intended use	Sterile, powder-free, polychloroprene surgical gloves intended to be worn on hand of healthcare personnel, operating room personnel and similar personnel to prevent contamination between the healthcare or similar personnel and the patient's body, fluids, waste, or environment during surgical procedure. Chemo-risk gloves designed for protection while working with the administration of chemotherapy drugs. Single use.	Length [mm]	EN 455-2 normative value	260	260	270	270	270	280	280
			Spec. [min]	290	290	290	290	290	290	290
		Width [mm]	EN 455-2 normative value	77	83	89	95	102	108	114
				±5	±5	±5	±5	±6	±6	±6
Material	Synthetic polychloroprene	Thickness single wall [mm]	Middle finger [min]	0,18						
			Palm [min]	0,16						
			Cuff [min]	0,15						
Donning powder	None	Force at break [N]	Before aging	9,0						
			EN 455-2 normative value	9,0						
Colour	Tan	Powder content [mg/glove]	After aging	9,0						
			EN 455-2 normative value	9,0						
Shape	Anatomic, curved fingers, hand specific									
Cuff	Beaded									
External surface	Textured, polymerized									
Internal surface	Layered polymer									
Packaging	1 pair per pouch, 50 pairs per dispenser, 200 pairs per carton									
MANUFACTURING AND SAFETY STANDARDS										
Manufacturer	Terang Nusa (Malaysia) Sdn. Bhd. 2, Jalan 8, Pengkalan Chepa 2, Industrial Zone 16100 Kota Bharu, Kelantan, Malezja									
EC Representative	Ulma International GmbH Pfaffenweg 35, 89231 Neu-Ulm, Niemcy									
Importer	Mercator Medical S.A. H. Modrzejewskiej 30 street 31-327 Cracow, Poland									
AQL	Manufacturing final release: G-I inspection level AQL 0.65 in accordance with ISO 2859-1.									
Latex protein content	N/A. Product does not contain natural rubber latex.									
Sterilization	Gamma (R)									
Classification	Medical Device: class IIa Rule classification acc. to declaration of conformity					Personal Protective Equipment: Category III (Regulation (EU) 2016/425) Type B (EN ISO 374-1)				
Conformity assessment body	TÜV SÜD Product Service GmbH, No 0123 Ridlerstraße 65, 80339, Munich, Germany					SATRA Technology Europe Limited, No 2777 Bracetown Business Park, Clonee, D15YN2P, Ireland				
Product compliances	EN 455-1, EN 455-2, EN 455-3, EN 455-4, EN ISO 15223-1, EN ISO 20417, EN 556-1, EN ISO 11737-1, EN ISO 11737-2, EN ISO 11137-1, EN ISO					EN ISO 374-1, EN ISO 374-2, EN ISO 374-4, EN ISO 374-5, EN 16523-1, EN ISO 21420				

**“TO PREVENT CORRUPTION & BRIBERY. CORRUPTION & BRIBERY IS A CRIME.
BE HONEST AND NO CHEATING”**

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	11137-2, EN ISO 14971, EN ISO 10993-1, EN ISO 10993-5, EN ISO 10993-10, EN ISO 10993-11, EN ISO 10993-23, EN ISO 11607-1, EN ISO 11607-2					
Quality compliances	EN ISO 13485, ISO 9001					
Viral test permeation	Test in accordance with EN ISO 374-5 (ISO 16604).					
Bacteria and fungi permeation	Test in accordance with EN ISO 374-5 (EN ISO 374-2).					
Chemotherapy drugs permeation test	Test in accordance with ASTM D6978.					
Chemical substances permeation test	Test in accordance with EN 16523-1.					
Biocompatibility/ biological evaluation	Test in accordance with EN ISO 10993-5. No cytotoxic evidence observed. Test in accordance with EN ISO 10993-10. No skin irritation and sensitization evidence observed. Test in accordance with EN ISO 10993-11.					
REACH	The product does not contain substances listed in candidate list according to Regulation (EC) 1907/2006.					
STORAGE AND DISPOSAL						
Long-term storage instructions	It is recommended to store the gloves in dry place, in the temperature of 5-35°C and to protect them against direct sunlight. Keep the gloves in a distance of not less than 1m from heating devices, sources of fire and ozone. Do not keep in direct vicinity of solvents, oils, fuels and lubricants.					
Transport instructions	Transport in conditions ensuring an appropriate hygienic standard, protecting the product against dirt. The product is not thermolabile - changing conditions regarding temperature or humidity in the short-term transport period do not affect the usability of the product or its properties or safety of use in any way. The product does not require transport in controlled conditions in terms of temperature and humidity (confirmed on the basis of accelerated aging tests and risk analysis).					
Shelf life	5 years from manufacturing date					
Product disposal	Used product should be treated as contaminated material, therefore local regulations regarding the disposal of such materials should be applied.					
Packaging disposal	Master packaging (carton, dispenser) is made of homogeneous material, does not contain any foil elements, does not contain any different type of materials, does not need to be separated into fractions. Packaging is 100% recyclable, in accordance with local regulations. Unit packaging (outer envelope and inner envelope) is to be regarded as contaminated medical material and local regulations for the handling of such materials must be followed.					
PRODUCT REFERENCES						
Size / REF number						
6.0	6.5	7.0	7.5	8.0	8.5	9.0
RC40001060_0687	RC40001065_0687	RC40001070_0687	RC40001075_0687	RC40001080_0687	RC40001085_0687	RC40001090_0687