

nitrylex® classic



PRODUCT DESCRIPTION			PHYSICAL PROPERTIES							
Manufacturer	MERCATOR MEDICAL S.A. Ul. H. Modrzejewskiej 30 31-327 Kraków		Size	XS (5-6)	S (6-7)	M (7-8)	L (8-9)	XL (9-10)		
Type of the glove	Non-sterile, powder-free, examination and protective glove for single use		Length [mm]	Spec. [min]		240	240	240	240	240
Intended use	Gloves intended for use in the medical field to protect patient and user from cross-contamination, intended to be used on one individual during a single procedure.		Width [mm]	Spec.		≤80	80 ±10	95 ±10	110 ±10	≥110
Material	Nitrile		Thickness (single wall) [mm]	Middle finger	Spec. [min]	0,06				
Donning powder	None			Palm	Spec. [min]	0,05				
Colour	Blue			Cuff	Spec. [min]	0,04				
Shape	Ambidextrous, gloves fitting either hand		Elongation at break [%]	Before ageing		500				
Cuff	Beaded			After ageing		400				
External surface	Microtextured + fingertip textured		Force at break [N]	Before ageing		6,0				
Internal surface	Chlorinated			After ageing		6,0				
Packaging	10 x 100 pcs	10 x 200 pcs	Powder content [mg/glove]	Spec.		<2				
			Latex protein content [µg/g]	Spec.		Not applicable				
PRODUCT REFERENCES										
Size / reference number	XS	RD30019001	RD30096001							
	S	RD30019002	RD30096002							
	M	RD30019003	RD30096003							
	L	RD30019004	RD30096004							
	XL	RD30019005	RD30096005							
MANUFACTURING AND SAFETY STANDARDS										
AQL	Manufacturing final release: G-I inspection level AQL 1.0 in accordance with ISO 2859-1									
Latex protein content	Not applicable. Gloves do not contain natural rubber latex.									
Residual powder content	<2 mg/glove (according to EN 455-3)									
Classification	Medical Device: CE Class I (Regulation (UE) 2017/745)		Personal Protective Equipment: CE 2777 Category III (Regulation (UE) 2016/425) Type B (EN ISO 374-1)							
Product compliances	EN 455-1, EN 455-2, EN 455-3, EN 455-4, EN ISO 15223-1, EN 1041		EN ISO 374-1, EN ISO 374-2, EN ISO 374-4 / EN 374-4, EN ISO 374-5, EN 16523-1, EN ISO 21420 / EN 420							
Quality compliances	EN ISO 13485, ISO 9001									
Viral test permeation	Test in accordance with ASTM F1671 and EN ISO 374-5 (ISO 16604)									
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.									
Food contact	Gloves intended for food contact based on Regulation (EU) No 10/2011, European Regulation (EC) No 1935/2004 and with Regulation (EC) No 2023/2006 on Good Manufacturing Practice. Gloves are suitable for handling any type of food and have been tested for Overall Migration Test acc. EN 1186.									
STORAGE AND DISPOSAL										
Storage instruction	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.									
Shelf life	3 or 5 years depend on LOT number (check the packaging)									
Disposal	The product must be disposed of in accordance with local regulations.									