MERCATOR MEDICAL



nitrylex® green

The instruction below should be used in conjunction with detailed information on the packaging.

Short description of the product

Nitrile examination and protective gloves, powder-free, non-sterile for disposable use

Full description of the product

Raw material : nitrile

External surface : microtextured + fingertip textured, polymerized

Internal surface : polymerized, chlorinated

Cuff : beaded Colour : green

Shape : ambidextrous, fitting to the right and left hand Size range : XS (5-6), S (6-7), M (7-8), L (8-9), XL (9-10)

AQL : 1.0

Quantity in packaging : 100 pcs. by weight

Shelf life : 3 years (from the date of manufacturing)

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 and fluorescent light. Recommended relative humidity in the room where the gloves are stored is $60\pm20\%$.

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.

Food contact

Gloves are marked with food contact symbol XI and comply with the requirements of 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:

Extraction conditions (tested for 30 min in 40°C)	Analysis results [mg/dm²]	Test Result (limit < 10 mg/dm²)
3% acetic acid	< 3.0	Pass
10% ethanol	< 3.0	Pass
50% ethanol	3.0	Pass
95% ethanol	6.0	Pass
Isooctane (6 min in 20°C)	< 3.0	Pass

MDR classification & compliance

Gloves are classified as class I according to Annex VIII of the Regulation (EU) 2017/745 and comply to standards:

EN 455-1:2000, EN 455-2:2015, EN 455-3:2015, EN 455-4:2009, EN ISO 15223-1:2016, EN 1041:2008+A1:2013.

PPER classification & compliance

Gloves are category III Personal Protective Equipment as per Annex I of the Regulation 2016/425 and comply to standards:

EN 420:2003+A1:2009, EN ISO 374-1:2016 (Type B), EN 374-2:2014, EN 16523-1:2015, EN 374-4:2013, EN ISO 374-5:2016.

Declaration of Conformity can be found under below web address: https://mercatormedical.eu/produkty/rekawice/diagnostyczne/nitrylex-green

Notified Body 2777
responsible for EU Type
Examination (Module B)
and Module C2 On-going
Conformity:

CE2777

Satra Technology Europe Ltd Bracetown Business Park, Clonee Dublin 15, Dublin, Ireland

Intended use

These are non-sterile examination and protective gloves for single use, intended for use in medical field to: protect patient and user from cross-contamination, conducting medical examinations, diagnostic and therapeutic procedures and for handling medical contaminated material. Gloves are classified as Medical Devices Class I and as a Personal Protective Equipment Category III. Their design and labelling corresponds to the requirements of the European Regulation 2017/745 on Medical Device and the European Regulation 2016/425 on Personal Protective Equipment. Gloves should be used solely according to their intended application.

Precautions and indications for use

Dry hands before putting the gloves on. Before usage, inspect the gloves for any defect or imperfections. Use at least 1 pair of gloves for one patient and one procedure, these are disposable gloves. Do not let chemical substances get under the gloves through the cuff. If a chemical substance reaches the skin, wash it away immediately with plenty of water with soap. If the gloves get punctured, torn or broken during their use, take them off and put on the new ones. Avoid using gloves dirty in the inside as they may cause irritation leading to skin inflammation or more serious damages. The gloves should not be used in contact with open fire and to protect against any sharp tools. The gloves are not intended for welding, electric shock protection, ionizing radiation or from the effect of hot or cold objects.

This information does not reflect the actual duration of protection in the workplace and the differentiation between mixtures and pure chemicals. Degradation results indicate the change in puncture resistance of the gloves after exposure to challenge chemical. The chemical resistance has been assessed under laboratory conditions from samples taken from the palm only (except in case where glove is equal to or over 400 mm – where the cuff is tested also) and relates only to the chemical tested and to the tested specimen. It can be different if the chemical is used in a mixture. The penetration resistance has been assessed under laboratory conditions and relates only to the tested specimen.

It is recommended to check that the gloves are suitable for the intended use because the conditions at the workplace may differ from the type test depending on the temperature, abrasion and degradation.

When used, protective gloves may provide less resistance to the dangerous chemical due to changes in physical properties. Movements, snagging, rubbing, degradation caused by the chemical contact etc. may reduce the actual use time significantly. For corrosive chemicals, degradation can be the most important factor to consider in selection of chemical resistant gloves.

Gloves are suitable for special purposes as they are examination gloves where risk of injury to the wrist is considered to be minimal, gloves are shorter than EN 420 min. length requirement.

Components / hazardous components

Some gloves may contain components known to be a possible cause of allergy for person allergic to them, who may develop contact irritation and/or allergic reaction. In case of an allergic reaction, seek medical assistance immediately.

Disposal

Used gloves can be contaminated with contagious or other hazardous substances. They should be disposed of in accordance with local regulation. Gloves should be buried or burned under controlled conditions

Manufacturer

MERCATOR MEDICAL S.A. ul. H. Modrzejewskiej 30 31-327 Cracow, Poland www.mercatormedical.eu

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Permeation performance levels as per EN ISO 374-1:2016							
• Level 1 > 10 min • Level 2 > 30 min • Level 3 > 60 min • Level 4 > 120 min • Level 5 > 240 min • Level 6 > 480 min							
Test results acc. to EN 16523-1:2015		EN 374-4:2013	Test results acc. to EN 16523-1:2015		EN 374-4:2013		
Chemical	Level	Degradation [%]	Chemical	Level	Degradation [%]		
35% Ethanol	6	55.0	50% Sulphuric Acid	6	21.1		
40% Isopropanol	6	68.7	5% Ethidium Bromide	6	32.9		
10% Acetic Acid	4	53.5	3% Hydrogen Peroxide	6	44.0		
50% Benzalkonium Chloride*	6	29.5	30% Hydrogen Peroxide (P)	2	52.8		
4% Chlorhexidine Digluconate**	6	32.9	37% Formaldehyde (T)	5	20.0		
10% Phosphoric Acid	6	14.0	50% Glutaraldehyde	6	22.9		
40% Sodium Hydroxide (K)	6	2.6	0.1% Phenol	6	24.7		
12% Sodium Hypochlorite	6	22.7		•	_		

^{*}minimum detectable permeation rate: 5 µg/cm²/min

^{**}minimum detectable permeation rate: 7 μg/cm²/min

EN 3/4-4:2013 Degradation levels indicate the change in puncture resistance of the gloves after exposure to the challenge chemical.						
	Test acc. To EN 374-2:2014 – Level 2 (ISO 2859)	Test acc. To EN ISO 374-5:2016				

Test acc. To EN 374-2:2014 – Level 2 (ISO 2859)		Test acc. To EN ISO 374-5:2016			
Performance level	AQL	Protection against bacteria & fungi	Pass		
Level 3	< 0.65	Protection against viruses	Pass		
Level 2	<1.5				
Level 1	< 4.0				
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Symbols used on the packaging Personal Protective Medical device Powdered gloves Equipment Do not re-use / gloves Keep away from moisture, Powder free gloves are intended for single use store in a dry place Keep away from solar Presence of polymer coating Non-sterile gloves and fluorescent light on the inner surface of the glove Temperature limitation / Presence of cosmetic coating Lot / batch number gloves store in temperature on the inner surface of the glove 5-35°C

Catalogue number

EU Authorised Representative, symbol should be accompanied by name and address of Authorised Representative

Expiry date

Marking of gloves protecting against bacteria and fungi

Marking of gloves protecting against viruses, bacteria and fungi

Marking of type A chemical resistant gloves. Six tested chemicals shall be identified by their code letter under pictogram

Marking of type B chemical resistant gloves. Three tested chemicals shall be identified by their code letter under pictogram

Marking of type C chemical resistant gloves. One tested chemicals shall be identified by their code letter under pictogram

Indicates compliance with the requirements of Russian market

Date of manufacture Manufacturer, symbol should

Keep away from ozone

be accompanied by name and address of Manufacturer

Food contact symbol (article is suitable for food contact, for details check the instruction for use)

Package made from paper, qualify for recycling

Package is treated as municipal waste

Consult instructions for use

Additional information on inner side of package

Indicates compliance with the requirements of

Presence of external texture on the glove

Gloves made from nitrile

Gloves made from vinyl

Raw material natural rubber latex

50 gloves by weight

100 100 gloves by weight

200 200 gloves by weight by weight

Do not use, if package is damaged









Ukrainian market

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■ HOW TO PUT THE GLOVES ON? ■













■ HOW TO TAKE THE GLOVES OFF? ■











