

Instruction for Use KL03/IFU-D16



01 Product Description

Product Name: Powder Free Sterile Latex Surgical Gloves

as an Under Glove

Material : Natural Rubber Latex

Colour : Dark Green
Shape : Anatomic
Cuff : Beaded

External Surface: Chlorinated and Siliconized

Internal Surface: Polymer Coated

Size : 5.5, 6.0, 6.5, 7.0, 7.5, 8.0, 8.5 and 9.0 Sterilization : Ethylene Oxide (EO) / GAMMA (R)

Shelf Life : 3 years

Basic UDI-DI : 806363LSGFM4

02 Intended Use

This disposal medical device is made up of natural rubber latex which is anatomically shaped with thumb position towards the Palm side of the index finger which reduces the fatigue on the hands, intended to be worn on the hands usually in surgical settings to provide barrier against potentially infectious fluids and other contaminants.

03 Product Classification & Standard compliance

Medical Device Classification: Class IIa, Rule#06

Conformity assessment route: Annex II section 4 of council

directive 93/42/EEC

Regulatory Authority : DNV Product Assurance As

Notified Body Number : 2460

Standard Compliance :

EN 455-1:2020+A2:2024, EN 455-2:2024, EN 455-3:2023, EN 455-4:2009, EN ISO 15223-1:2021, EN 1041:2013, EN 566-1:2024, EN ISO 11135: 2014/A1:2019, EN ISO 11137-1:2015/A2:2019, EN ISO 11137-2:2015/A1:2023, EN ISO 11737-1:2018/A1:2021, EN ISO 11737-2:2020, ISO 11607-1:2020/A1:2023, ISO 11607-2:2020/A1:2023, EN ISO 10993-1:2020, EN ISO 10993-5:2009, EN ISO 10993-7:2008, EN ISO 10993-10:2021, EN ISO 10993-11:2017, EN ISO 10993-23:2021, ISO 10282:2023, ASTM D 3577:2019, ISO 13485:2016, ISO 14001:2015, ISO 9001:2015, IS 13422:2024.

04 Storage Instruction

Gloves must be stored in cool dry environment which is dust free. Cartons and Boxes must be stored unopened until required. Recommended storage temperature is 5°C-35°C. Avoid exposure to direct light, heat and excessive humidity.

As ozone is deleterious, storage area should not contain any equipment capable of generating ozone such as ultraviolet light, fluorescent lights, mercury vapour lamps, photocopier, high voltage equipment, x-ray units, electric motors and electro surgical equipments.

05 Indication For Use

- Dry hands thoroughly before donning.
- Protective gloves should only be used for the intended application and in the correct size.
- These are sterile gloves for single use only.
- Users should take care when using the gloves. Using them solely according to their intended application.
- Before usage, inspect the gloves for any defect or imperfection.

06 Contraindication

- Latex gloves are made of Natural rubber latex, which may cause allergic reactions including anaphylaxis response if the user is allergic to latex.
- Gloves contain Natural Latex; persons who are sensitive to Latex should consult a physician before using.

07 Precautions

- Do not use if package is damaged or wet.
- Risk of reuse: May cause infection, allergic reaction and poor barrier protection.
- Gloves shall not be worn where there is a risk of entanglement by moving parts of machines is needed.
- The results do not reflect the actual duration of protection in the workplace due to other factors influencing the performance, such as temperature, abrasion, degradation etc.

08 Warnings

- Device disposal should be done as per local regulatory norms.
- Do not re-sterilize.
- The product contains Natural Rubber Latex which may cause allergic reactions including anaphylactic responses to some individuals.
- The gloves not intend to prevent Electrical shock care should be taken to have proper earthing in Medical Device Electrical appliances user.
- Necessary caution to be practiced against probable Electrical Hazards.

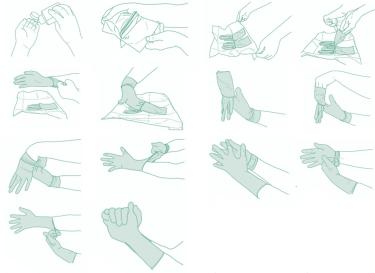


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Surgicare

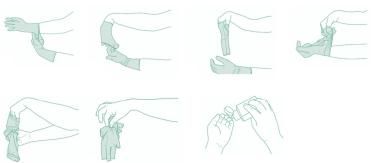
09 Directions for use

Glove Opening and Donning Procedure:



- Remove the walleted gloves (inner wrapper) from the Pouch (outer wrapper) by Peel open from the corner for Paper Pouch (Peel down to open pouch).
- Open the walleted glove to see "Left" and "Right" compartment.
- Pinch back upper and lower flaps of the inner wrapper.
- Using the middle flaps, open the wrapper touching only the 1 inch margin for safety.
- Be sure wrapper does not close over gloves after opening to avoid contamination.
- Using the thumb and the first two fingers of the nondominant hand, pinch the cuff of the folded edge of the glove cuff for the dominant hand, touching only the inside surface of the glove.
- Slide dominant hand in to the gloves keeping hand point downwards and pull up to wrist.
- Using the glove hand insert the 4 fingers under the cuff of the other glove and pull the glove up to the arm.
- Adjust the gloves as necessary.

Glove Removal Procedure:



- Take hold of the first glove at the wrist.
- Fold it over and peel it back, turning it inside out as it goes. Once the glove is off, hold it with your gloved hand.
- To remove the other glove, place your bare fingers inside the cuff without touching the glove exterior. Peel the glove off from the inside, turning it inside out as it goes. Use it to envelope the other glove.

10 Explanation of Symbols

444

Manufacturer

EC REP

Authorized representative in the European Community

AMSTERMED B.V., Saturnusstraat 46-62,

Unit 032, 2132 HB Hoofddorp,

The Netherlands

Date of Manufacture / Country of Manufacture

2

Use by date

LOT

Lot No

REF

Reference Number / Catalogue Number

SN

Serial Number

C€ 2460

CE Logo

STERILE EO

Sterilization using Ethylene oxide

STERILE R

Sterilization using Irradiation



Single Sterile Barrier System



Do not re-sterilize



Do not use package is damaged or wet



Keep away from sunlight



Keep dry



Temperature limit



This way up or end up



Keep away from Ozone



Single Use



Instruction for Use



Caution



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Latex Caution



Medical Device



Unique Device Identifier



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EN ISO 374-1/Type A EN ISO 374-1/Type B EN ISO 374-1/Type C







Performance Level

Protection index

1

2

3

4

5

6



TYPE A Permeation min. 6 chemicals, level 2 (table 1)

TYPE B Permeation min. 3 chemicals, level 2 (table 1)

TYPE C Permeation min. 1 chemical, level 1 (table 1)



Protective gloves protecting from viruses, bacteria and fungus



Protective gloves protecting from bacteria and fungus

Time

>10 min

>30 min

>60 min

>120 min

>240 min

>480 min

Table 1

Chemicals

- Methanol
- В Acetone
- \mathbf{C} Acetonitrile
- D Dichloromethane
- Е Carbon disulphide
- F Toluene
- G Dimethylamine
- Н Tetrahydrofuran
- Ethyl acetate
- n-Heptane
- Sodium hydroxide 40% K
- Sulphuric acid 96%
- Nitric acid 65% Μ
- Acetic acid 99%
- O Ammonium hydroxide 25%
- Hydrogen peroxide 30%
- Hydrofluoric acid 40%
- Formaldehyde 37%

EN ISO 374-1/Type B











In Compliance with the Harmonized Standards Personal Protective Equipment, risk category III under

Regulation (EU)2016/425 EN ISO 21420: 2020 EN ISO 374-1: 2016 EN ISO 374-2:2019 EN 16523-1:2015+A1:2018

EN ISO 374-4: 2019 EN ISO 374-5: 2016

Regulatory Authority : SGS Fimko Oy, Takomotie 8, FI-00380, Helsinki, Finland.

Notified Body Number : 0598

Warnings as per

- This information does not reflect the actual duration of protection in the work place and the differentiation between mixtures and pure chemicals.
- The Chemical resistance has been assessed under laboratory conditions from samples taken from the palm only (except in cases where the glove is equal to or over 400mm - where the cuff is tested also) and relates only to the chemical tested. It can be different if the chemical is used in a mixture.
- Tt is recommended to check that the gloves are suitable for the intended use because the conditions at the work place may differ from the type test depending on temperatures, 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.
- Before usage, inspect the gloves for any defect or imperfections.
- For Single use only.
- The Penetration resistance has been assessed under laboratory conditions and relates only to the tested specimen.

Cest Results as per EN ISO 374-1 & 5

1. EN ISO 21420 : 2020

Protective Gloves - General Requirement and test methods

	Туре	Declared Size	5.5	6.0	6.5	7.0	7.5	8.0	8.5	9.0
		Length (mm)	268.7	285.3	282.3	285.3	300.7	299.0	291.7	292.0
		Circumference (mm)	145.3	160.7	168.3	181.7	196.3	203.3	216.3	229.3
Glove Dexterity Performance Level 5					•	•				
pH value 7.0										

Polyaromatic Hydrocarbons content

	Results
Benz (a) anthracene	Not Detected
Chrysene	Not Detected
Benzo (b) Fluoranthene	Not Detected
Benzo (k) Fluoranthene	Not Detected
Benzo (a) pyrene	Not Detected
Dibenz (a,h) anthracene	Not Detected
Benzo (e) pyrene	Not Detected
Benzo (j) Fluoranthene	Not Detected



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2. EN ISO 374-2: 2019

Protective gloves against dangerous chemicals and micro-organisms - Part 2 : Determination of resistance to penetration

Resistance to Penetration	Status
Air Leakage	Pass
Water Leakage	Pass

Clause	Test Name	Test	Performance level	
7.2	Air leak test (Air Pressure Used: 0.5 kPa)	Sample size 6.0 6.5 7.0	Leakage No Leakage No Leakage No Leakage	Pass
		7.5 8.0 8.5	No Leakage No Leakage No Leakage	
7.3	Water leak test	6.0 6.5 7.0 7.5 8.0	Leakage No Leakage No Leakage No Leakage No Leakage No Leakage	Pass

3. EN ISO 374-4: 2019

Protective gloves against dangerous chemicals and micro-organisms - Part 4: Determination of resistance to degradation by chemicals

	Chemical / CAS No	Exposure Duration	Test Results (Per change in puncture	Observation	
	Sodium hydroxide 40% 1310-73-2	60±5 minutes	Glove sample 1 2 3 Mean Standard Deviation	Result (%) 1.6 1.6 4.2 2.4 1.507	No change
	Sulphuric acid 96% 7664-93-9	60±5 minutes	Glove sample 1 2 3 Mean Standard Deviation	Result (%) 97.3 96.2 96.6 96.7 0.558	Severe Swelling & Colour change
	Nitric acid 65% 7697-37-2	60±5 minutes	Glove sample 1 2 3 Mean Standard Deviation	Result (%) 51.1 51.1 50.8 51.0 0.159	Severe Swelling & Colour change
	Hydrogen 60±5 peroxide 30% minutes 7722-84-1		Glove sample 1 2 3 Mean Standard Deviation	Result (%) 41.4 43.3 42.6 42.4 0.971	Moderate swelling
			Glove sample 1 2 3 Mean Standard Deviation	Result (%) 11.2 9.0 10.6 10.3 1.131	Slight swelling
	Formaldehyde 37% 50-00-0	60±5 minutes	Glove sample 1 2 3 Mean Standard Deviation	Result (%) 12.5 11.9 11.5 12.0 0.515	Slight swelling

4. EN 16523-1: 2015+A1:2018

Determination of material resistance to permeation by chemicals - Part 1 : Permeation by Liquid chemical under conditions of continuous contact.

Chemical CAS No	Loop System /collection medium	Analytical Technique used	Mean thickness (mm)	NBT at NPR 1.0 µg cm ⁻² min ⁻¹ (minutes)	Performance level accordance to BS EN ISO 374-1:2016+ A1.2018 Table 1	Observation
Sodium hydroxide 40% 1310-73-2	Closed loop /Grade 3 water/350 rpm	Continuous measurement with conductivity electrode	0.17 0.16 0.18	>480 >480 >480 >480	6	No change
Nitric acid 65% 7697-37-2	Closed loop /Grade 3 water/350 rpm	Continuous measurement with conductivity electrode	0.18 0.16 0.17	54 56 59	2	Severe Swelling & Colour change
Hydrogen peroxide 30% 7722-84-1	Closed loop /Grade 3 water	Periodic measurement with HPLC	0.18 0.19 0.16	>480 >480 >480 >480	6	Slight Swelling
Formaldehyde 37% 50-00-0	Closed loop /Grade 3 water	Periodic measurement with HPLC	0.17 0.17 0.16	31 33 35	2	Slight Swelling
Sulphuric acid 96% 7664-93-9	Closed loop /Grade 3 water/350 rpm	Continuous measurement with conductivity electrode	0.16 0.17 0.18	15 14 17	1	Severe Swelling & Colour change
Acetic acid 99% 64-19-7	Closed loop /Grade 3 water/350 rpm	Continuous measurement with conductivity electrode	0.16 0.17 0.17	5 4 6	0	Slight Swelling

5. EN ISO 374-5:2016

Protective gloves against dangerous chemicals and microorganisms - Part 5: Terminology and Performance Requirements for microorganisms risks

Protected against viruses according to ISO 16604 procedure B

Specimen	Observe the Presence of Plaques on Agar Plate (PFU)	Results at 20 Kpa	Assay Titer Ratio
1	<1	Pass	
2	<1	Pass	
3	<1	Pass	1.3
Negative control	No	NA	
Positive control	Yes	NA	

Plate exposes during testing (Settle Plate): No plaque formation observed.

➤ Declaration of conformity is supplied with the product.