KIMTECH™

DATA PACK

Kimtech™ G3 Sterile Latex Gloves

56843 / 56844 / 56845 / 56846 / 56847
56848 / 56849 / 56842

Former Product Codes:
HC1360S/ HC1365S / HC1370S / HC1375S / HC1380S
HC1385S / HC1390S / HC1310S

Publication code:
4560.01 EN 04.3

Table of Content

[1]For other languages please visit the product page on www.kimtech.eu
[2]Certificate of Analysis / Irradiation are available on a lot by lot basis, please visit: www.kimtech.eu/ressources/certifcates

KIMTECH

EU Declaration of Conformity

Version	Revision Date:	DoC #:	Date of last issue: -
1.0	21.11.2022	100000019626	Date of first issue: 21.11.2022

The manufacturer, and his authorised representative established in the Community, Kimberly-Clark Europe Ltd., confirms that the PPE models, as described, are in conformity with the provisions of Regulation (EU) 2016/425 for category

Style	Product Code(s)	Product Description
Gloves	56843, 56844, 56845, 56846, 56847, 56848, 56849, 56842	KIMTECH* G3 Sterile Latex Gloves

Personal Protective Equipment, the European harmonised standard:
Category III PPE
Subject to the procedures set out in Module D of the The Regulation (EU) 2016/425 EC under the supervision of Notified Body.

Harmonized Standards
: EN ISO 21420:2020 (Protective gloves – General requirements and test methods)
: EN ISO 374-1:2016+A1:2018 (Protective gloves against chemicals and micro-organisms) as a Type C glove against reagent K
EN ISO 374-5:2016: (Protective gloves against chemicals and micro-organisms) with EN 374-2:2019 performance level 2 and including Viral Penetration

Is identical to the tested samples which are the subject of:
EU type-examination certificate: 0598/PPE/22/3707
Granted to Kimberly - Clark Europe Ltd, based on Technical File by the Notified Body:PPE.TG.EU.330.v04

Signed on behalf of the manufacturer in the European Community.

Christelle Bouvier		Revision Date: 21.11.2022
Senior Regulatory Affairs Manager		
Kimberly-Clark Europe Ltd.		

As requested by the (EU) 2016/425, addresses of the parts involved as follows:

Kimberly-Clark Europe Limited	
Walton Oaks, Dorking Road, Tadworth, Surrey, KT20 7NS, United Kingdom	
Telephone: +44 1737 736000	Fax: +44 1737 736670
SGS FIMKO OY (0598)	
Takomotie 8,HELSINKI, 00380, Finland	
Telephone: +358 9 696 361	Fax:
TÜV SÜD Product Service GmbH Zertifizierstellen (0123)	

EU Declaration of Conformity

Version	Revision Date:	DoC #:	Date of last issue: -
1.0	21.11.2022	100000019626	Date of first issue: 21.11.2022

Ridlerstraße 65, MÜNCHEN, 80339, Germany,	
Telephone: +49 (89) 50084261	Fax:

UK Declaration of Conformity

Version	Revision Date:	DoC #:	Date of last issue: -
1.0	21.11.2022	100000050945	Date of first issue: 21.11.2022

The manufacturer, and his authorised representative established in the United Kingdom, Kimberly-Clark Europe Ltd., confirms that the PPE models, as described, are in conformity with the provisions of Regulation (EU) 2016/425 as brought into UK law and amended.

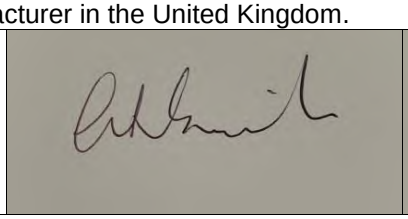
Style	Product Code(s)	Product Description
Gloves	56843, 56844, 56845, 56846, 56847, 56848, 56849, 56842	KIMTECH* G3 Sterile Latex Gloves

Personal Protective Equipment:
Category III PPE
Subject to the procedures set out in Module D of the Regulation (EU) 2016/425 as brought into UK law and amended under the supervision of Approved Body

UK Designated Standards:
: EN ISO 21420:2020 (Protective gloves – General requirements and test methods)
: EN ISO 374-1:2016+A1:2018 (Protective gloves against chemicals and micro-organisms) as a Type C glove against reagent K
EN ISO 374-5:2016: (Protective gloves against chemicals and micro-organisms) with EN 374-2:2019 performance level 2 and including Viral Penetration

Is identical to the tested samples which are the subject of:
UK type-examination certificate:0120/PPE/221008
Granted to Kimberly - Clark Europe Ltd, based on Technical File examination by the Approved Body:PPE.TG.EU.330.v04

Signed on behalf of the manufacturer in the United Kingdom.

Liz Brigden		Revision Date: 21.11.2022
KCP EMEA Regulatory Affairs Associate Director		
Kimberly-Clark Europe Ltd.		

As requested by the (EU) 2016/425 as brought into UK law and amended, addresses of the parts involved as follows:

Kimberly-Clark Europe Limited	
Walton Oaks, Dorking Road, Tadworth, Surrey, KT20 7NS, United Kingdom	
Telephone: +44 1737 736000	Fax: +44 1737 736670
SGS United Kingdom Limited (0120)	
Rossmore Business Park,Ellesmere Port,South Wirral, CH65 3EN, Cheshire, United Kingdom	
Telephone: +44 (0) 1934 522917	Fax:

UK Declaration of Conformity

Version	Revision Date:	DoC #:	Date of last issue: -
1.0	21.11.2022	100000050945	Date of first issue: 21.11.2022

TUV SUD BABT UNLIMITED (0168)	
Octagon House, Concorde Way, Segensworth North,Fareham, PO15 5RL, Hampshire, United Kingdom	
Telephone: +44 1489 558100	Fax:

Kimberly-Clark Professional*1400 Holcomb Bridge Rd.Roswell, GA 30076 USA

CERTIFICATE OF ANALYSIS

Product Description : **Kimtech* G3 Sterile Latex Gloves, Hand Specific, 12" Pair packed**

K-C Code : 56843-40, 56844-40, 56845-40, 56846-40, 56847-40, 56848-40, 56849-40, 56842-40

Lot # : 420123
Batches : SM30012XX to SM30312XX
SM30012VX to SM30312VX

Total Cases per Lot : 5,026
Date of Manufacture : Jan-23
Expiration Date : 2027-12

Physical Test Data**

		Visual Defects				Elongation (%)	Tensile (MPa)
	Watertight	Critical Visual	Major	Minor	Dimensions	Pre Aging	Pre Aging
Sample Size :	2675	2675	2675	2675	1020	380	380
AQL Level :	1.5	1.5	2.5	4.0	2.5	2.5	2.5
Failures Allowed per AQL :	71	65	103	149	51	19	19
Failures :	33	0	0	0	0	0	0
Inspection Results :	Accept	Accept	Accept	Accept	Accept	Accept	Accept
					Averages:	853	29.04

Test Methods : Water tight test ASTM D 5151, EN 455-1, Elongation and Tensile ASTM D 412, ASTM D 3577, EN 455-2, Dimension ASTM D 3577, EN 455-2

Particle Test Data**

Particle Size (µm)	Min	Max	Standard Deviation	Average Particles/cm²
0.5 - 1.0	253	1139	278	680
1.0 - 2.0	32	275	52	91
2.0 - 5.0	10	85	17	28
5.0 - 10.0	1	4	1	2
10.0 - 20.0	0	2	0	1
>20	0	0	0	0
Total per Sample	296	1261	310	802

Test Method : IEST-RP-CC005.4

Extractable Ion Test Data**

	Anions Results						
	Fluoride F^-	Chloride Cl^-	Nitrite NO_2^-	Bromide Br^-	Nitrate NO_3^-	Phosphate PO_4^{-3}	Sulfate SO_4^{-2}
µg/g glove	0.463	45.856	1.388	1.388	3.795	2.312	5.480
µg/cm²	0.004	0.376	0.012	0.012	0.031	0.019	0.045

	Cations Results				Trace Element Results	
	Sodium Na^+	Ammonium NH_4^+	Potassium K^+	Magnesium Mg^{+2}	Calcium Ca^{+2}	Zinc Zn
µg/g glove	0.974	1.516	1.202	0.925	2.359	42.04
µg/cm²	0.008	0.012	0.010	0.008	0.019	0.34

Test Method : IEST-RP-CC005.4

Endotoxin Data**

Test Result:	BD	**Endotoxin Units/ device**
Specification:	≤ 20	**Endotoxin Units/ device**

Detection Limit is ≤ 0.4 EU/device : BD = Below Detection Limit
Test Method : Limulus Amebocyte Lysate (LAL) Test: Kinetic Turbidimetric Technique

Sterility Data

This product conforms to standards for sterility ISO11137 with sterility assurance level (SAL) 10^{-6}. A Certificate of Irradiation is also available to assure that the product has received the specified radiation dosage.

**Testing performed at final quality inspection gate prior to sterilization.

Review By : 14 Feb 2023

(QA Sr. Manager)

FORM-21963/5

STERIS

http://www.steris-ast.com

Certificate of Irradiation

Date Issued: 25-Jan-2023 MY03S12721817-1-1

This is to certify that Synergy Sterilisation (M) Sdn. Bhd., a STERIS Company has where appropriate delivered an irradiation process in accordance with the current certified standards:

EN ISO 11137-1 Sterilisation of Health Care Products
EN ISO 9001 Quality Management System
EN ISO 13485 Quality System - Medical Devices

Safeskin Medical & Scientific (Thailand) Ltd
200 MOO 8 KANCHANAVANICH ROAD
PRIK SADAO
Amphur Sadad
SONGKHLA 90120
THAILAND

Order Information

Account Number:	101195
Synergy Health Sales Part Reference:	1126467
Customer Reference Number:	4027021145
Product Description:	KIMTECH*G3 STERILE LATEX GLOVES,HAND SPECIFIC,12" PAIR PACKED
Validation Reference:	0.0764 Rev.02
Quantity Received:	1062
Customer Minimum Specification kGy:	25.0
Customer Maximum Specification kGy:	50.0
Customer Unit Lot/Batch Number:	SEE BELOW

26/01/2023
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QA Executive
Synergy Sterilisation (M) Sdn. Bhd
+60(0)44152111

STERIS

http://www.steris-ast.com

Certificate of Irradiation

Date Issued: 25-Jan-2023 MY03S12721817-1-1

This is to certify that Synergy Sterilisation (M) Sdn. Bhd., a STERIS Company has where appropriate delivered an irradiation process in accordance with the current certified standards:

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Safeskin Medical & Scientific (Thailand) Ltd
200 MOO 8 KANCHANAVANICH ROAD
PRIK SADAO
Amphur Sadad
SONGKHLA 90120
THAILAND

Other Process Details:

Kimtech* G3 Sterile Latex Gloves, Hand Specific,12" Pair Packed
KC Code : 56843-40, 56844-40, 56845-40, 56846-40, 56847-40, 56848-40, 56849-40, 56842-40

Catalog Number(s)	Lot No./Batch No.	Quantity
56843-40	420123/SM30122XX	8
56843-40	420123/SM30112XX	36
56845-40	420123/SM30132XX	141
56845-40	420123/SM30122XX	124
56845-40	420123/SM30112XX	184
56845-40	420123/SM30102XX	82
56846-40	420123/SM30112XX	105
56846-40	420123/SM30102XX	32
56847-40	420123/SM30142XX	36
56847-40	420123/SM30122XX	104
56847-40	420123/SM30112XX	38
56848-40	420123/SM30132XX	124
56848-40	420123/SM30112XX	36
56849-40	420123/SM30112XX	12

Irradiation Data

Date and Time of Irradiation:	20-Jan-2023 08:08
Reference Dose Range kGy:	29.8 - 31.6

26/01/2023
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STERIS

http://www.steris-ast.com

Certificate of Irradiation

Date Issued: 25-Jan-2023 MY03S12721817-1-1

This is to certify that Synergy Sterilisation (M) Sdn. Bhd., a STERIS Company has where appropriate delivered an irradiation process in accordance with the current certified standards:

EN ISO 11137-1 Sterilisation of Health Care Products
EN ISO 9001 Quality Management System
EN ISO 13485 Quality System - Medical Devices

Safeskin Medical & Scientific (Thailand) Ltd
200 MOO 8 KANCHANAVANICH ROAD
PRIK SADAO
Amphur Sadad
SONGKHLA 90120
THAILAND

Calculated Minimum Dose kGy:	25.7
Calculated Maximum Dose kGy:	33.8

Irradiation Release Authorised By Synergy Health Sterilisation UK, a STERIS Company

26/01/2023
NOR AZWIN BT. YUSUF
QA Executive
Synergy Sterilisation (M) Sdn. Bhd
+60(0)44152111

Kimberly-Clark
PROFESSIONAL®

Summary of Validation of Kimtech Pure G3 (formerly Safeskin Critical) Latex gloves performed in 2008.

The validation study was performed according to the guidelines established by ANSI/AAMI/ISO 11137:2006 "American National Standard, Sterilization of Healthcare Products—Requirements for Validation and Routine Control—Radiation Sterilization.". The need to re-establish the dose arose from the fact that the packaging was significantly changed. The maximum dose study performed previously established the maximum dose at 50 kGY. The product was tested to be non-bacteriostatic/fungistatic.

Three batches of above-referenced gloves were assayed for bioburden levels. Method 1 was used. The overall average was **24.67 CFU/device**. No single batch was shown to be more than twice this overall average and therefore this overall average was used to calculate the sub process verification dose.

With reference to table 5 of the ISO11137:2006 (this was the current version at that time), the nearest value listed equal to or greater than the bioburden level is 24.67 CFU. Therefore the sub-process dose required for the sterility assurance level of 10^{-2} is 6.3 kGy.

One hundred units were irradiated at this dose and subsequently individually tested for sterility at **6.3+/-10% kGy**.

After full incubation period, there were no growths found. Therefore statistical verification is accepted. Referring to Table 5, to achieve the desired Sterility Assurance Level of 10-6, the minimum dose required is 19.1 kGy.

Submitted by:

Ruthlyn M. Reyes

Ruthlyn M. Reyes
KCP Operations

Date: January 19, 2010

Test Method for Analyzing Liquid Particle Counts

This test method is used to analyze the mobile particle contaminants from cleanroom gloves.

1. Scope
   - 1.1. The test method covers the average particulate contamination found on gloves designated for cleanroom applicability.
   - 1.2. The average contaminant concentration will be reported in particles per cm^2 in two ways:
     - 1.2.1. By size grouping, 0.5 to 1.0 microns, 1.0 to 2.0 microns, 2.0 to 5.0 microns, 5.0 to 10.0 microns, 10.0 to 20.0 microns, greater than 20.0 microns, and a total particle count greater than 0.5 microns.
     - 1.2.2. Statistical analysis of each grouping consisting of Minimum Value, Maximum Value, Standard Deviation, and Average Value, for each group of individual gloves.
   - 1.3. The safe and proper use of gloves is beyond the scope of this test method.
   - 1.4. This test method does not purport to address all of the safety concerns, if any, associated with its use. It is the responsibility of the user of this Test Method to establish appropriate safety and health practices and determine the applicability of regulatory limitations prior to use.

2. Referenced Documents
   - 2.1. IEST-RP-CC005.3 Recommended Practice for Gloves and Finger Cots Used in Cleanrooms and Other Controlled Environments
   - 2.2. Work Instruction
3. Apparatus
   - 3.1. Analytical Balance, capable of readability and repeatability to 0.1 mg
   - 3.2. Particle Measuring Systems CLS-900 Liquid Particle Counting System
   - 3.3. 2000 mL glass beaker or 1000mL glass conical flask
   - 3.4. Stainless Steel Forceps, 10" length
   - 3.5. 250 ml Volumetric Flask
   - 3.6. 500 ml Volumetric Flask
   - 3.7. High Purity Deionized Water System, capable of producing 18.2 MOhm quality water
   - 3.8. Point of Use Filter, 0.2 micron size
   - 3.9. Orbital Shaker, ¾" orbit, capable of 200 rpm
   - 3.10. Circular Die, 1.5 inch diameter, calibrated

4. Procedure
   - 4.1. Test Preparation
     - 4.1.1. Prior to extraction, all Erlenmeyer flasks will be cleaned no less than five times with high purity deionized water filtered to 0.2 microns at point of use.
     - 4.1.2. All related equipment (forceps, volumetric flasks, etc.) must be rinsed with high purity deionized water prior to use.
   - 4.2. Extraction
     - 4.2.1. Randomly pull a glove from the package.
     - 4.2.2. Place glove finger-first into the one liter Erlenmeyer flask and hold open by cuff using the rinsed forceps.
     - 4.2.3. Empty into the inside of the glove 500 ml high purity filtered deionized water.
     - 4.2.4. Allow the glove to settle into the Erlenmeyer flask.
     - 4.2.5. Place an additional 250 ml high purity filtered deionized water over the glove within the Erlenmeyer flask.
     - 4.2.6. Allow the Erlenmeyer flask with glove to agitate on the shaker for 10 minutes ± 10 seconds at a rate of 150 rpm ± 10 rpm.
     - 4.2.7. Using clean tongs, immediately remove the glove from the container. Drain any trapped liquid into the beaker by manipulating the fingers on the glove, with the tongs
     - 4.2.8. Dispose of the glove.
     - 4.2.9. Repeat the extraction two additional times to complete the set.
     - 4.2.10. Prepare a process blank, using all the steps in section 4.2, without placing the glove in the Erlenmeyer flask.

4.3. Measurement

4.3.1. Follow the Work Instruction for the Liquid Particle Counter for analyzing the solutions.

4.4. Glove Surface Area

4.4.1. Pull three gloves from the production package and weigh to the nearest 0.1 mg.

4.4.2. Record as A.

4.4.3. Cut the 3 gloves with square die (5X5 cm.) by wheel cutter at palm. This will give you six cut-out sections.

4.4.4. Weight the six cut-out sections. Record this as B.

4.4.5. Calculate the surface area of the glove using the following equation :

$$\frac{A \times 5 \times 5 \times 4}{B}$$

5. Calculations

5.1. Calculate counts/cm^2 by channel size using the following equation:

$$\frac{\text{(Sample (counts/mL)-Blank (Counts/mL) x Extraction volume (mL) x DF}}{\text{Surface area (in cm}^2\text{)}}$$

5.2. Total Counts/cm^2 : $= \sum AllChannelSizes$

6. Reporting

6.1. The final report should include the Lot Number, Batch number, Product Description, Part Number, and any other pertinent information about the sample, as well as the final calculated counts/cm^2 by channel size and a total counts/cm^2 greater than 0.5 microns.

6.2. Statistics will be calculated and reported on sample sizes greater than three.

Test Method for Analyzing Extractables

This test method is used to analyze the soluble ionic extractable contaminants from cleanroom gloves.

1. Scope
   1.1. The test method covers the average ionic contamination found on gloves designated for cleanroom applicability.
   1.2. The average contaminant concentration will be reported in one of two ways:
      1.2.1. Micrograms of ionic contaminant per gram of glove weight (ug/g), also described as ppm.
      1.2.2. Micrograms of ionic contaminant per square centimeter of glove area (ug/cm^2)
   1.3. This test method does not cover contaminants that are insoluble in water, or organic macromolecules.
   1.4. The safe and proper use of gloves is beyond the scope of this test method.
   1.5. This test method does not purport to address all of the safety concerns, if any, associated with its use. It is the responsibility of the user of this Test Method to establish appropriate safety and health practices and determine the applicability of regulatory limitations prior to use.

2. Referenced Documents
   2.1. IEST-RP-CC005.2 Recommended Practice for Gloves and Finger Cots Used in Cleanrooms and Other Controlled Environments.
   2.2. Work Instruction WI 10-05-26, Work Instruction for Performing Ion Chromatography Analysis of Gloves

3. Apparatus
   3.1. Analytical Balance, capable of readability and repeatability to 0.1 mg
   3.2. Ion Chromatograph
   3.3. Extraction Containers, 1 liter capacity, HDPE with screw type lids
   3.4. Stainless Steel Forceps, 10" length
   3.5. 500 ml Volumetric Flask
   3.6. High Purity Deionized Water System, capable of producing 18.0 MOhm quality water
   3.7. Point of Use Filter, 0.1 micron size
   3.8. Circular Die, 1.5 inch diameter, calibrated

4. Procedure
   4.1. Test Preparation
      4.1.1. Prior to extraction, all extraction containers will be cleaned using high purity deionized water high purity deionized water filtered to 0.2 microns at point of use.
      4.1.2. All related equipment (forceps, volumetric flasks, etc.) must be rinsed with high purity de-ionized water prior to use.

   4.2. Extraction
      4.2.1. Randomly pull a glove from the package.
      4.2.2. Place glove finger-first into the one liter Erlenmeyer flask and hold open by cuff using the rinsed forceps.
      4.3. Empty into the inside of the glove approximately 250 ml high purity filtered deionized water.
      4.4. Allow the glove to settle into the extraction container.
      4.5. Pour remaining 250 ml high purity filtered deionized water over the glove within the extraction container.
      4.6. Place the lid upon the container and seal tightly.
      4.7. Gently swirl the container to ensure that all surfaces of the glove are wetted.
      4.8. Allow the glove to extract in the deionized water for at least 10 minutes, but no longer than 11 minutes.
      4.9. Remove the glove by the fingers, allowing most of the water trapped in the fingers to drain back in to the extraction container.
      4.10. Dispose of the glove.
      4.11. Repeat extraction two additional times to complete the set.
      4.12. Prepare a sample blank, using all the steps in section 2, without placing the glove in the extraction container.

4.13. Measurement

4.13.1. Follow the guidelines for the Ion Chromatograph for analyzing aqueous solutions.

4.14. Glove weight and surface area

4.14.1. Pull three gloves from the production package and weigh to the nearest 0.1 mg.

4.14.2. Record as A.

4.14.3. Cut the 3 gloves with square die (5X5 cm.) by wheel cutter at palm. This will give you six cut-out sections.

4.14.4. Weight the six cut-out sections. Record this as B.

4.14.5. Calculate the surface area of the glove using the following equation :

$$\text{Surface area} = \frac{A \times 5 \times 5 \times 4}{B}$$

5. Calculations

5.1. Once the data output from the Chromatograph has been reviewed for errors, calculate the following:

5.1.1. ug/g (ppm) contamination: $= \dfrac{(AnalyteConc.) * (500ml)}{GloveWeight}$

5.1.2. ug/cm^2 contamination: $= \dfrac{(AnalyteConc.) * (500ml)}{SurfaceArea}$

6. Reporting

6.1. The final report should include the Lot number, Batch number, Product description, Part number, and any other pertinent information about the sample, as well as the final calculated contaminant concentration in ug/g and ug/cm^2.

Case Label

G3 Sterile Latex Gloves

6.5

12" (30.5cm) 20 x 10 = 200

(EN) G3 Sterile Latex Gloves
(FR) G3 Gants en latex stériles
(ES) Guantes estériles de látex G3
(DE) G3 Sterile Latexhandschuhe
(NL) G3 steriele latex handschoenen
(IT) G3 Guanti sterili in lattice
(RU) G3 Стерильные латексные перчатки
(UA) Рукавички стерильні латексні G3
(PT) Luvas de látex estéreis G3
(KO) G3 멸균 라텍스 장갑
(ZH) G3 无菌乳胶手套
(JA) G3滅菌ラテックス手袋

EN ISO 374-1:2016+ A1:2018/Type C

EN ISO 374-5:2016 VIRUS

56844 **40**

LOT XXXXXX-XXXXXXXXXXX
Lot Number
Номер партии
製造番号

MM-YYYY
Date of Manufacturing
Дата производства
製造年月

YYYY-MM
Expiration Date
Использовать до
使用期限

CE 0123 EAC TP TC 019/2011
UK CA 0168
LATEX AQL 1.5
LM5684440OL-00

1 00 36000 56844 5

KIMTECH

Sterile Pair Pouch

KDF Artwork


KIMTECH
G3 Sterile Latex Gloves
56844 40
KIMTECH
KIMTECH
KIMTECH

KIMTECH™

G3 Sterile Latex Gloves
12" / 30.5cm - Hand Specific Pairs

(EN) G3 Sterile Latex Gloves
- 12"/30.5cm Length
- Hand Specific Pairs
- Textured
- For the Sterile Cleanroom Environment
- For Industrial Use Only

NOTICE: THIS INSERT SHOULD BE FURNISHED OR MADE AVAILABLE TO THE USERS OF THESE GLOVES AS A SAFETY PRECAUTION.
This is a Category III PPE certified according to Regulation (EU) 2016/425 and Regulation (EU) 2016/425 as brought into UK law and amended. Risk: Gloves offer protection against chemicals (Splash) and micro-organisms.
This information does not reflect the actual duration of protection in the workplace and the differentiation between mixtures and pure chemicals. Tested for Microorganism Hazards / not tested against viruses. The chemical resistance has been assessed under laboratory conditions from samples taken from the palm only and relates only to the chemical tested. It can be different if the chemical is used in a mixture. 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 temperature, abrasion and degradation. Degradation results indicate puncture resistance of glove after chemical exposure. When used, protective gloves may provide less resistance to the dangerous chemical due to changes in the 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. The penetration resistance has been assessed under laboratory conditions and relates only to the tested specimen. Before usage, inspect the gloves for any defect or imperfections. Discard any gloves presenting a defect. Refer to enclosed donning and doffing instructions. Store in a cool dry place. Dispose of according to local regulations. A list of substances known to cause allergies can be supplied on demand.
CONTACT US: If you have any questions about this product, call the manufacturer at (US) 1-800-255-6401 (EU) +44(0) 1737 736000 (AP) +603 7807 8210

(FR) G3 Gants en latex stériles
- Longueur 12"/30,5cm
- Paires s'adaptant à la main
- Texturés
- Pour les environnements de salles blanches stériles
- Pour usage industriel uniquement

AVIS : PAR MESURE DE SÉCURITÉ, CET ENCART DOIT ÊTRE FOURNI AUX UTILISATEURS DE CES GANTS OU ÊTRE À LEUR DISPOSITION.
Il s'agit d'un EPI de catégorie III certifié conformément à la réglementation (UE) 2016/425. Risque : Les gants offrent une protection contre les produits chimiques (éclaboussures) et les micro-organismes.
Ces informations ne reflètent pas nécessairement la durée réelle de protection en milieu de travail ni la différence entre les mélanges et les produits chimiques purs. Protection contre les micro-organismes / non testés pour les virus. La résistance aux produits chimiques a été évaluée en laboratoire avec des échantillons prélevés dans la paume seulement et ne concerne que le produit chimique testé. Les résultats peuvent être différents si le produit chimique est utilisé dans un mélange. Il est recommandé de s'assurer que les gants conviennent à l'usage prévu en conditions réelles car celles-ci peuvent différer de celles du test standard en fonction de la température, de l'abrasion et de la dégradation. Des résultats montrant une dégradation indiquent que les gants résistent aux perforations lors d'une exposition aux produits chimiques. Lorsqu'ils sont utilisés, les gants peuvent fournir moins de résistance aux produits chimiques dangereux en raison de changements dans les propriétés physiques. Les mouvements, déchirures, frottements et dégradations engendrés lors du contact avec les produits chimiques, etc. peuvent réduire la durée réelle d'utilisation de façon significative. Dans le cas des produits chimiques corrosifs, la dégradation peut être le facteur le plus important à considérer lors du choix de gants résistants aux produits chimiques. La résistance à la pénétration a été évaluée en laboratoire et ne concerne que l'échantillon testé. Inspecter les gants avant l'utilisation pour vérifier qu'ils ne comportent pas de défauts ou d'imperfections. Jeter les gants présentant un défaut. Consulter les instructions ci-jointes pour enfiler et retirer les gants. Ranger dans un endroit frais et sec. Mettre au rebut conformément aux règlements municipaux. Une liste des substances connues pour causer des allergies peut être fournie sur demande.
NOUS CONTACTER : Pour tout renseignement concernant ce produit, appeler le fabricant au (États-Unis) 1-800-255-6401
(Europe) +44(0) 1737 736000 (Asie-Pacifique) +603 7807 8210

(DE) G3 Sterile Latexhandschuhe
- 12"/30.5 cm Länge
- Handspezifische Paare
- Texturiert
- Für die sterilkritische Reinraumumgebung
- Nur für die industrielle Verwendung

HINWEIS: DIESE PACKUNGSBEILAGE SOLLTE ANWENDERN ALS SICHERHEITSVORKEHRUNG AUSGEHÄNDIGT ODER ZUR VERFÜGUNG GESTELLT WERDEN.
Dies ist ein PSA-Produkt der Kategorie III, das nach der Verordnung (EU) 2016/425 zertifiziert ist. Risiko: Handschuhe bieten Schutz gegen Chemikalien (Spritzer) und Mikroorganismen.
Diese Informationen spiegeln nicht die tatsächliche Schutzdauer am Arbeitsplatz und die Differenzierung zwischen Mischungen und reinen Chemikalien wider. Geprüft auf Gefährdung durch Mikroorganismen / nicht auf Viren geprüft. Die Chemikalienbeständigkeit wurde unter Laborbedingungen durch ausschließlich an der Handfläche entnommene Proben bestimmt und bezieht sich nur auf die geprüfte Chemikalie. Die Beständigkeit kann unterschiedlich sein, wenn die Chemikalie in einer Mischung verwendet wird. Es wird empfohlen, die Eignung der Handschuhe für den vorgesehenen Verwendungszweck zu prüfen, da sich die Bedingungen am Arbeitsplatz von den Prüfbedingungen hinsichtlich Temperatur, Abnutzung und Zersetzung unterscheiden können. Ergebnisse der Degradationsprüfung zeigen Stichfestigkeit auch nach Kontakt mit Chemikalien. Schutzhandschuhe können bei der Verwendung aufgrund von Veränderungen der physikalischen Eigenschaften eine geringere Beständigkeit gegen die gefährliche Chemikalie aufweisen. Bewegungen, Verhakung, Reibung, durch den Kontakt mit Chemikalien verursachte Zersetzung usw. können die tatsächliche Verwendungszeit erheblich verringern. Bei korrosiven Chemikalien kann Zersetzung der wichtigste Faktor sein, der bei der Auswahl von chemikalienbeständigen Handschuhen zu berücksichtigen ist. Der Penetrationswiderstand wurde unter Laborbedingungen geprüft und bezieht sich nur auf die geprüfte Probe. Die Handschuhe vor der Verwendung auf Mängel oder Fehler prüfen. Handschuhe mit Mängeln sind zu entsorgen. Siehe beigefügte Anweisungen zum Anziehen und Ausziehen. An einem kühlen, trockenen Ort lagern. Gemäß den lokalen Bedingungen entsorgen. Eine Liste der Stoffe, die bekanntermaßen Allergien auslösen, kann auf Anfrage geliefert werden.
SO KONTAKTIEREN SIE UNS: Bei Fragen zu diesem Produkt rufen Sie bitte den Hersteller an unter der Nummer (US) 1-800-255-6401;
(EU) +44(0) 1737 736000; (AP) +603 7807 8210

(NL) G3 steriele latex handschoenen
- 30.5cm/12 inch lang
- Handspecifieke paren
- Getextureerd
- Voor steriele schone ruimtes
- Alleen voor industrieel gebruik

WAARSCHUWING: DEZE BIJSLUITER DIENT ALS VEILIGHEIDSMAATREGEL GEGEVEN TE WORDEN AAN OF TER BESCHIKKING GESTELD TE WORDEN VAN DE GEBRUIKERS VAN DEZE HANDSCHOENEN.
Dit is een persoonlijk beschermingsmiddel van categorie III volgens Verordening (EU) 2016/425/EEG. Risico: Handschoenen bieden bescherming tegen chemische stoffen (spatten) en micro-organismen.
Deze informatie is geen weerspiegeling van de werkelijke beschermingsduur in de werkomgeving en de differentiatie tussen mengsels en zuivere chemicaliën. Getest op gevaren door micro-organismen / niet getest voor virussen. De chemische weerstand is onder laboratoriumomstandigheden beoordeeld op grond van monsters genomen van alleen de palm en heeft alleen betrekking op het geteste chemische product. Het kan anders zijn als het chemische product in een mengsel wordt gebruikt. Het wordt aanbevolen te controleren of de handschoenen geschikt zijn voor het beoogde gebruik omdat de omstandigheden in de werkomgeving kunnen verschillen van de typetest afhankelijk van temperatuur, schuring en afbraak. Bij een afbraakproef bleek dat de handschoen na chemische blootstelling bestand is tegen prikken. Bij het gebruik kunnen beschermende handschoenen minder weerstand bieden tegen het gevaarlijke chemische product vanwege veranderingen in de fysische eigenschappen. Bewegingen, blijven haken, wrijven, afbraak veroorzaakt door contact met het chemische product etc. kunnen de werkelijke gebruiksduur aanzienlijk verminderen. Bij corrosieve chemische producten kan afbraak de belangrijkste factor zijn waarmee rekening moet worden gehouden bij de keuze van chemisch bestendige handschoenen. De weerstand tegen indringen is beoordeeld onder laboratoriumomstandigheden en heeft alleen betrekking op het geteste specimen. Controleer de handschoenen vóór gebruik op beschadiging of onvolkomenheden. Gooi handschoenen met een beschadiging weg. Raadpleeg de bijgevoegde instructies voor aan- en uittrekken. Op een koele, droge plaats bewaren. Afvoeren volgens de plaatselijke voorschriften. Een lijst van stoffen waarvan bekend is dat ze allergieën veroorzaken, is op aanvraag verkrijgbaar.
CONTACT MET ONS OPNEMEN: Als u vragen hebt over dit product, kunt u de fabrikant bereiken op nr.: (Verenigde Staten) +1-800-255-6401
(Europa) +44(0) 1737 736000 (Azië-Pacific) +603 7807 8210.

(IT) G3 Guanti sterili in lattice
- Lunghezza 12"/30.5 cm
- Paia destri e sinistri
- Ruvidi
- Per camera bianca sterile
- Solo per uso industriale

AVVISO - QUESTO INSERTO DEVE ESSERE FORNITO O RESO DISPONIBILE COME MISURA DI SICUREZZA A COLORO CHE UTILIZZANO QUESTI GUANTI.
Questo prodotto è certificato come DPI di categoria III secondo il Regolamento (UE) 2016/425. Rischio: i guanti offrono protezione contro sostanze chimiche (schizzi) e microrganismi.
Queste informazioni non riflettono la durata effettiva della protezione sul luogo di lavoro e la distinzione tra prodotti chimici miscelati e puri. Testato per rischi da microrganismi/non testato contro i virus. La resistenza chimica è stata misurata in condizioni di laboratorio su campioni presi solo dal palmo della mano e si riferisce solo al prodotto chimico testato. Può essere diverso se il prodotto chimico viene utilizzato in una miscela. Si consiglia di controllare che i guanti siano idonei per l'uso previsto poiché le condizioni del luogo di lavoro possono differire dal tipo di test a seconda della temperatura, abrasione e degradazione. I risultati di degradazione indicano la resistenza dei guanti alle perforazioni dopo l'esposizione a prodotti chimici. Quando utilizzati, i guanti di protezione possono fornire meno resistenza ai prodotti chimici pericolosi a causa di cambiamenti delle proprietà fisiche. Il tempo effettivo di utilizzo può essere ridotto significativamente a causa di movimenti, sfilacciamento, strofinamento o degradazione dovuti al contatto con prodotti chimici, ecc. In caso di contatto con prodotti chimici corrosivi, il fattore più determinante da considerare nella scelta di guanti resistenti ai prodotti chimici è la resistenza alla degradazione. La resistenza alla penetrazione è stata misurata in condizioni di laboratorio e riguarda solo il campione testato. Prima dell'uso, ispezionare i guanti per verificare l'assenza di difetti o imperfezioni. Smaltire adeguatamente qualsiasi guanto che presenti difetti. Consultare le istruzioni allegate per indossare e togliere il prodotto. Conservare in un luogo asciutto e fresco. Smaltire in conformità alle disposizioni locali. Un elenco di sostanze note come causa di allergie può essere fornito su richiesta.
PER CONTATTARCI - Per chiarimenti circa questo prodotto rivolgersi al produttore al numero 1-800-255-6401 (USA), +44(0) 1737 736000 (Europa), +603 7807 8210 (Asia Pacifico).

(ES) Guantes estériles de látex G3
- 12 pulg./30,5 cm de largo
- Pares específicos para cada mano
- Texturizados
- Para entornos de sala blanca de esterilización
- Sólo para uso industrial

AVISO: COMO MEDIDA DE SEGURIDAD, ESTE ENCARTE SE DEBE ENTREGAR O PONER A DISPOSICIÓN DE LOS USUARIOS DE ESTOS GUANTES
Este es un producto de Categoría III PPE certificado según el Reglamento (UE) 2016/425. Riesgo: Estos guantes ofrecen protección frente a químicos (salpicaduras) y microorganismos.

REF **G3 Sterile Latex -**
6.0 = 56843
6.5 = 56844
7.0 = 56845
7.5 = 56846
8.0 = 56847
8.5 = 56848
9.0 = 56849
10.0 = 56842

CE 0123 EAC TP TC 019/2011
UKCA 0168

AQL 1.5

EN ISO 374-1:2018+A1/ Type C
- (EN) Tested for Watertightness, Chemical Permeation and Chemical Degradation
- (FR) Testés pour l'imperméabilité, la perméation de produits chimiques et la dégradation chimique
- (ES) Sometidos a pruebas de estanqueidad, permeación química y degradación química
- (DE) Geprüft auf Wasserdichtigkeit, Permeation von chemischen Substanzen und chemische Abbaubarkeit
- (RU) Прошли испытания на водонепроницаемость, проницаемость для химических веществ и химическое разрушение
- (UA) Пройшли випробування на водонепроникність і захист від проникнення та стійкість до хімічних речовин
- (JA) 水密性、化学物質の浸透、化学的劣化は試験済み

EN ISO 374-5:2016 VIRUS
- (EN) Tested for Microorganism Hazards
- (FR) Testé contre les risques de microorganismes
- (ES) Sometido a pruebas de peligros presentados por microorganismos
- (DE) Geprüft für Gefahren durch Mikroorganismen
- (RU) Испытано на наличие опасных микроорганизмов
- (UA) Перевірено на наявність небезпечних мікроорганізмів
- (JA) 微生物学的危険性で検査済み

- (EN) Single Use Only.
- (FR) Usage unique seulement
- (ES) Usese una sola vez
- (DE) Nur zur einmaligen Verwendung
- (RU) Только для одноразового применения
- (UA) Виключно для одноразового застосування
- (JA) 再使用禁止

- (EN) Protect from Heat and Radioactive Sources
- (FR) À protéger contre les sources de chaleur et radioactives
- (ES) Proteger contra fuentes de calor y radiactividad
- (DE) Vor Hitze und radioaktiven Strahlen schützen
- (RU) Беречь от нагрева и источников радиоактивного излучения
- (UA) Оберігати від нагрівання і джерел радіоактивного випромінювання
- (JA) 熱遮へい及び放射線防護

- (EN) Keep Dry
- (FR) Conserver au sec
- (ES) Mantener secos
- (DE) Trocken halten
- (RU) Хранить в сухом месте
- (UA) Зберігати в сухому місці
- (JA) 湿気厳禁

LATEX

G3 Sterile Latex Gloves

	Permeation Test EN ISO 374-1:2016+A1:2018			Degradation Test EN ISO 374-4:2019
(EN) Chemical	Breakthrough Time(min.)		Performance Level	Performance Level %
Sodium Hydroxide, 40% (K)	>480		Class 6	-58.8

EN ISO 21420:2020 Dexterity Classification = 5

Certificates available from www.kimtech.com/certificates
EU/UK Declarations of Conformity available at: www.kimtech.eu

προερχόμενα μόνο από την παλάμη και σχετίζεται μόνο με τη χημική ουσία που ελέγχθηκε. Μπορεί να διαφέρει, αν η χημική ουσία χρησιμοποιείται σε μείγμα. Συνιστάται ο έλεγχος της συμβατότητας των γαντιών με την προβλεπόμενη χρήση, επειδή οι συνθήκες στο χώρο εργασίας ενδέχεται να διαφέρουν από τον έλεγχο τύπου ανάλογα με τη θερμοκρασία, την τριβή και την αποδόμηση. Τα αποτελέσματα αποδόμησης υποδεικνύουν αντοχή σε διάτρηση του γαντιού μετά από την έκθεση σε χημικά. Κατά τη χρήση των προστατευτικών γαντιών ενδέχεται να παρέχεται λιγότερη αντίσταση στην επικίνδυνη χημική ουσία εξαιτίας αλλαγών στις φυσικές ιδιότητες. Κινήσεις, σχισίματα, τριβή, αποδόμηση που προκλήθηκε από επαφή με τη χημική ουσία κ.λπ. ενδέχεται να μειώσουν σημαντικά τον πραγματικό χρόνο χρήσης. Όσον αφορά στις διαβρωτικές χημικές ουσίες, η αποδόμηση είναι ο πιο σημαντικός παράγοντας που πρέπει να λάβει κανείς υπόψη του κατά την επιλογή γαντιών ανθεκτικών στις χημικές ουσίες. Η αντίσταση διείσδυσης έχει υπολογιστεί σε εργαστηριακές συνθήκες και σχετίζεται μόνο με το ελεγχόμενο είδος. Πριν από τη χρήση ελέγξτε τα γάντια για ελαττώματα ή ατέλειες. Απορρίψτε τυχόν γάντια που παρουσιάζουν ελάττωμα. Ανατρέξτε στις συνημμένες οδηγίες φορέματος και αλλάγματος. Φυλάσσεται σε δροσερό και ξηρό χώρο. Απορρίψτε σύμφωνα με τους τοπικούς κανονισμούς. Ένας κατάλογος ουσιών που είναι γνωστό ότι προκαλούν αλλεργίες, μπορεί να παρασχεθεί κατ'αίτηση.
ΕΠΙΚΟΙΝΩΝΙΑ: Αν έχετε ερωτήσεις σχετικά με το παρόν προϊόν, καλέστε τον κατασκευαστή στον αριθμό (Η.Π.Α.) 1-800-255-6401 (Ε.Ε.)
+44(0) 1737 736000 (Ασία-Ειρηνικός) +603 7807 8210

(TR) G3 Steril Lateks Eldiven
- 12"/30.5 cm Uzunluğunda
- Ele Özel Çift
- Dokulu
- Steril Temiz Oda Ortamı İçin
- Yalnızca Endüstriyel Kullanım İçindir

ÖNEMLİ: BU BİLGİLENDİRME EKİ GÜVENLİK ÖNLEMİ OLARAK KULLANICIYA ELDEN VERİLMELİ YA DA KULLANICININ ERİŞİMİNE SUNULMALIDIR.
Bu, (EU) 2016/425 sayılı Yönetmeliğe göre onaylanmış bir Kategori III KKD ürünüdür. Risk: Eldivenler kimyasallara (Sıçrama) ve mikroorganizmalara karşı koruma sağlar.
Bu bilgi, çalışma yerlerindeki gerçek koruma süresini ve karışımlar ile saf kimyasallar arasındaki farklılığı yansıtmamaktadır. Mikroorganizma Tehlikesi Testi Yapılmıştır / virüslere karşı test yapılmamıştır. Kimyasal direnç, laboratuvar koşulları altında yalnızca avuç içinden alınan numuneler ile ölçülmüştür ve sadece test edilen kimyasala ilişkindir. Kimyasal bir karışımda kullanılıyorsa, farklı olabilir. Eldivenlerin istenen kullanım için uygun olup olmadığını kontrol etmeniz önerilir, çünkü çalışma yerindeki koşullar sıcaklık, aşınma ve bozunmaya bağlı olarak tip testinden farklı olabilir. Bozunum sonuçları, kimyasal maruziyet sonrasında eldivenin delinme direncini göstermektedir. Koruyucu eldivenler kullanıldığında, fiziksel özelliklerde meydana gelen değişiklikler nedeniyle tehlikeli kimyasallara karşı daha az direnç gösterebilir. Hareketler, delinmeler, sürtünme, kimyasal temastan kaynaklanan bozunmalar gerçek kullanım süresini büyük ölçüde azaltabilir. Aşındırıcı kimyasallar için kimyasallara dirençli eldiven seçiminde bozunma en önemli faktördür. Nüfuz etmeye karşı direnç laboratuvar koşulları altında ölçülmüştür ve yalnızca test edilen numuneye ilişkindir. Kullanmadan önce eldivenlerde herhangi bir kusur veya eksiklik olup olmadığını kontrol edin. Kusurlu eldivenleri atın. Ekli takma ve çıkarma talimatlarına bakın. Serin ve kuru ortamda saklayın. Yerel düzenlemelere göre bertaraf edin. Alerjiye neden olduğu bilinen maddelerin listesi talep üzerine temin edilebilir.
BİZİ ARAYIN: Bu ürün hakkındaki her türlü sorunuz için 1-800-255-6401 (ABD), +44(0) 1737 736000 (Avrupa), +603 7807 8210 (Asya) numaralı telefondan imalatçıya ulaşabilirsiniz.

(ZH) G3 无菌乳胶手套
- 12"/30.5 cm 长
- 左右手特定成套
- 有纹理
- 针对无菌洁净室环境
- 仅适于工业用途

注释: 应该为将该手套作为安全防护措施的用户提供本说明书。
本产品属于类别 III PPE 产品，获得了法规 (EU) 2016/425 认证。风险提示：手套可以起到对化学品（溅出）和微生物的保护作用。
本信息未反应工作场所的实际防护持续时间以及混合物与纯化学品之间的区别。 进行微生物危害检测/非针对病毒进行检测。 仅在实验条件下通过手掌测量耐化学性，仅与被测试的化学品相关。 化学品用于混合物时，情况有所不同。 由于温度、磨损、降解等原因，工作场所的的条件可能与典型试验的条件有所不同，因此建议检查手套是否适用于预期用途。 降解结果表明手套在接触化学品后的耐穿刺性。 使用时，由于物理性质的变化，防护手套对危险化学品的抵抗性可能有所降低。 运动、障碍、摩擦、化学接触引起的降解等可能会严重缩短实际使用时间。 对于腐蚀性化学品，在选择防化手套时，降解可能是要考虑的最重要的因素。 已在实验条件下测量抗渗透性，仅与检测的样本相关。 使用前检查手套有无缺陷。 丢弃任何有缺陷的手套 请参阅随附的穿脱说明。 储存在阴凉干燥处。 按照当地法规处理。
可以根据需要提供已知会导致过敏的物质清单。
联系我们：如果您对本品有任何疑问，请致电制造商（美国）：(US) 1-800-255-6401 (EU) +44(0) 1737 736000 (AP) +603 7807 8210

(KR) G3 멸균 라텍스 장갑
- 12"/30.5cm 길이
- 한손형 한 켤레
- 엠보싱 처리
- 멸균 클린룸 환경용
- 공업용

참고: 이 인서트는 이 장갑 사용자가 안전 예방책으로 사용할 수 있도록 가공 또는 제조되었습니다.
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- ความยาว 12 นิ้ว/30.5 ซม.
- คู่ของมือที่ถนัด
- ปั๊มลาย
- สำหรับสภาพแวดล้อมภายในห้องปลอดเชื้อที่ปราศจากเชื้อ
- สำหรับใช้ในอุตสาหกรรมเท่านั้น

หมายเหตุ: เอกสารเพิ่มเติมนี้ควรได้รับการจัดทำหรือจัดเตรียมให้แก่ผู้ใช้ถุงมือเหล่านี้สำหรับใช้เป็นข้อควรระวังเพื่อความปลอดภัย
ผลิตภัณฑ์ PPE นี้ได้รับการจัดหมวดหมู่เป็นประเภท III ตามกฎระเบียบ (EU) 2016/425
ความเสี่ยง: ถุงมือช่วยป้องกันสารเคมี (กระเด็น) และจุลินทรีย์
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เนื่องจากสภาวะในสถานที่ทำงานอาจแตกต่างจากประเภททั่วไปของถุงมือที่ทดสอบ ทั้งนี้ขึ้นอยู่กับอุณหภูมิ การขัดข่วน และการเสื่อมสภาพ
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เนื่องจากการเปลี่ยนแปลงในคุณสมบัติทางกายภาพ การเคลื่อนไหว การเฉือน การถู ความเสื่อมสภาพที่เกิดจากการสัมผัสสารเคมี เป็นต้น อาจลดทอนเวลา สำหรับสารเคมีกัดกร่อน ความเสื่อมสภาพอาจเป็นปัจจัยที่สำคัญที่สุดในการพิจารณาเลือกถุงมือที่ทนต่อสารเคมี ความทนต่อการเจาะได้รับการประเมินภายใต้สภาพห้องปฏิบัติการ และสัมพันธ์กับตัวอย่างที่ทดสอบเท่านั้น ก่อนการใช้งาน ให้ตรวจสอบตำหนิหรือข้อบกพร่องต่าง ๆ ทิ้งถุงมือใดๆ ที่มีข้อบกพร่อง โปรดอ่านคำแนะนำที่แนบมาด้วยเกี่ยวกับการสวมและถอดถุงมือ เก็บในที่แห้งและเย็น ก่อนตามระเบียบข้อบังคับท้องถิ่น สามารถจัดหารายชื่อสารที่ทราบว่าอาจก่อให้เกิดอาการแพ้ให้ตามความต้องการ
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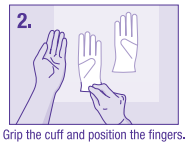
(JA) G3滅菌ラテックス手袋
- 長さ30.5 cm
- 左右別ペア
- テクスチャー加工
- 滅菌クリーンルーム環境対応
- 工業用途のみ

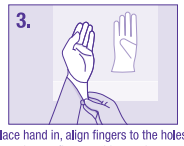
注意事項：本添付文書は、安全上の注意事項として、手袋の使用者に渡すか、使用者が参照できるようにしてください。
これは PPE 規則 (EU) 2016/425 に基づいてカテゴリ III 製品の認定を受けています。 リスク：手袋は、化学物質（液体に含まれる）や微生物から使用者を保護します。
この情報は、実際の作業場での保護期間、および、化学物質が混合物か純粋なものかを保障するものではありません。 微生物の有害性に対する保護性能は試験済み／ウイルスに対する保護性能は未試験。 耐薬品性とは、手のひらのみから採取したサンプルを、実験室条件下で特定の化学品のみに対して試験し、得られた結果を指します。 これは、化学物質が混合物の一部として使用される場合、異なることもあります。

実際の作業場の状態は、温度、磨耗および劣化など、タイプテストと異なる場合があるため、手袋が意図した用途に適していることを確認することが推奨されています。 劣化の結果は、化学物質への暴露後の手袋の対破壊性を示します。 保護手袋は、使用中に起こる物理的特性の変化により、危険な化学物質に対する抵抗力が低下することがあります。 動作、かぎ裂き、摩擦、化学物質との接触による劣化により、実際の耐用期間が大きく短縮される可能性があります。 腐食性化学物質の場合、劣化は耐薬品性手袋の選択において最も重要な要素となります。 耐貫通性とは、実験室条件下で評価され、使用検体で得られた結果のみを指します。 使用前に、手袋に欠陥や不完全な点ががないか点検してください。 欠陥のある手袋は廃棄すること。 添付の着用および着脱手順を参照してください。 涼しく乾燥した場所に保管してください。 地域の規制に従って廃棄してください。 アレルギーを引き起こすことが知られている物質のリストは、オンデマンドで供給可能です。
お問い合わせ先：本品についてご不明な点がございましたら、製造業者へ電話(1-800-255-6401 (米国)、+44 (0)1737 736000(ヨーロッパ)、(+603) 7807 8210 (アジア)でお問い合わせください。

Donning Gloves

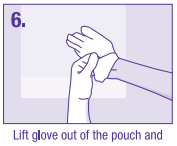

1. Check seal is undamaged. Open wallet on a clean surface. Open the pair pouch.

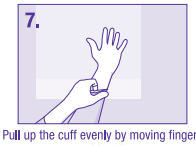

2. Grip the cuff and position the fingers.

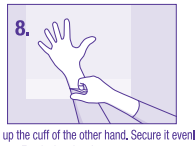

3. Place hand in, align fingers to the holes. Open and close fingers whilst pulling glove up.


4. Once first glove is on, leave the cuff unfolded.


5. Start with the second glove. Place the fingers in-between fold and pouch. Lift fingers to reveal opening.


6. Lift glove out of the pouch and repeat step 3.


7. Pull up the cuff evenly by moving fingers round beading.


8. Pull up the cuff of the other hand. Secure it evenly up the arm. For double donning restart process at step 1.

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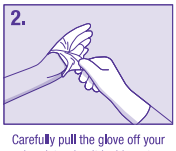
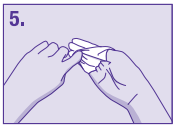
Doffing Gloves


1. Grasp the exterior of one glove with your other gloved hand.


2. Carefully pull the glove off your hand, turning it inside-out.


3. Ball the glove up and hold in your other gloved hand.


4. Slide your ungloved finger into the opening of the other glove.


5. Carefully pull the glove off your hand, turning it inside out again.

6. Discard appropriately.

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