

pure[11]-Nr.: 1105211, Marke:

Eigenschaften

Empfohlene Reinraumklassen

ISO

GMP

we care

pure[11] GmbH |
Bavariafilmplatz 7 | D-82031 Grünwald
T +49 89 5589434 0
F +49 89 5589434 77
www.pure11.de
info@pure11.de

Material

-

Verpackung

- Box

Produktvarianten

pure¹¹-Nr.: 1105211,

pure¹¹-Nr.: 1105211WH10, Halyard Nitril-Handschuhe PUREZERO HG3 weiß #CLN323210

Farbe: Weiß; Größe: 10 / VE: 200PAAR

pure¹¹-Nr.: 1105211WH6, Halyard Nitril-Handschuhe PUREZERO HG3 weiß #CLN323260

Farbe: Weiß; Größe: 6 / VE: 200PAAR

pure¹¹-Nr.: 1105211WH6_5, Halyard Nitril-Handschuhe PUREZERO HG3 weiß #CLN323265

Farbe: Weiß; Größe: 6,5 / VE: 200PAAR

pure¹¹-Nr.: 1105211WH7, Halyard Nitril-Handschuhe PUREZERO HG3 weiß #CLN323270

Farbe: Weiß; Größe: 7 / VE: 200PAAR

pure¹¹-Nr.: 1105211WH7_5, Halyard Nitril-Handschuhe PUREZERO HG3 weiß #CLN323275

Farbe: Weiß; Größe: 7,5 / VE: 200PAAR

pure¹¹-Nr.: 1105211WH8, Halyard Nitril-Handschuhe PUREZERO HG3 weiß #CLN323280

Farbe: Weiß; Größe: 8 / VE: 200PAAR

pure¹¹-Nr.: 1105211WH8_5, Halyard Nitril-Handschuhe PUREZERO HG3 weiß #CLN323285

Farbe: Weiß; Größe: 8,5 / VE: 200PAAR

pure¹¹-Nr.: 1105211WH9, Halyard Nitril-Handschuhe PUREZERO HG3 weiß #CLN323290

Farbe: Weiß; Größe: 9 / VE: 200PAAR

HALYARD

HALYARD* **PURE**ZERO* HG3 WHITE STERILE CLEANROOM GLOVES

DATA PACK



For the Cleanroom Environment
For Industrial Use Only

HALYARD* **PURE**ZERO* HG3 WHITE STERILE CLEANROOM GLOVES DATA PACK

Some documents depicted here are FOR EXAMPLE ONLY. For specific documents related to lot numbers, please visit **www.halyardhealth.com/information**, or ask your Distributor or HALYARD* Representative.

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TECHNICAL DATA SHEET



HALYARD*

HALYARD* **PURE**ZERO* HG3 White Sterile Nitrile Gloves

For the Critical Cleanroom Environment
For Industrial Use Only

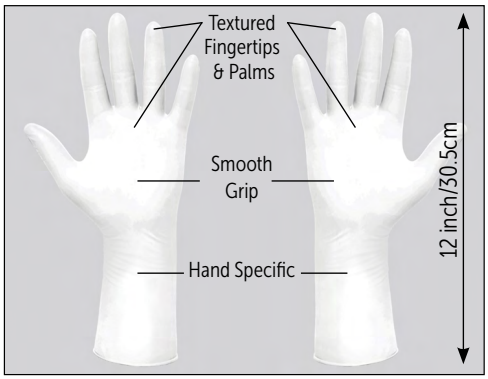
TECHNICAL DATA SHEET

Meets USP <797> Guidelines.

Meets USP <800> Guidelines.

Description

HALYARD* **PURE**ZERO* HG3 White Sterile Nitrile Cleanroom Gloves are designed for critical cleanroom environments such as pharmaceutical and biotechnology cleanroom manufacturing as well as sterile compounding cleanroom applications. These hand specific gloves are clean processed (washed repeatedly in deionized water) to ensure consistent control of low particles, extractables and endotoxin levels, and are recommended for use in ISO Class 3 or higher and Grade A/B cleanrooms. Because HALYARD* **PURE**ZERO* HG3 Sterile Cleanroom Gloves are made with an **accelerator-free**[1] nitrile polymer, there is a reduced risk of allergies and skin irritation associated with accelerator chemicals in other nitrile gloves.



Manufactured at OUR OWN Safeskin Facility in Thailand.

Cleanliness Properties

Max Particle Count (>0.5 µm)	<950 Particles/cm^2	IEST RP-CC005
Max Endotoxin Level	<20 EU	
Ionic Content (Extractable ions)	**Max Level (ug/g)**	IEST RP-CC005
Calcium	50	
Chloride	35	
Magnesium	5	
Nitrate	20	
Potassium	5	
Sodium	10	
Sulfate	10	
Zinc	7	
Ammonium	5	

Physical Properties

AQL	0.65
Sterile	✓
Hand Specific Pairs	✓
Smooth Grip	✓
Textured Fingertips and Palms	✓
Accelerator-Free[1]	✓
Latex-Free	✓
Powder-Free	✓
Silicone-Free	✓
Contains No Nickel	✓
Static Dissipative in Use	✓
Tensile Strength[2]	35 MPa (Target)
Ultimate Elongation[2]	600%
Sterility Assurance Level (SAL)	10^{-6}
Shelf Life	3 Years

HALYARD*

HALYARD* **PURE**ZERO* HG3 White Sterile Nitrile Gloves

For the Critical Cleanroom Environment
For Industrial Use Only

TECHNICAL DATA SHEET

Glove Dimensions

	6.0	6.5	7.0	7.5	8.0	8.5	9.0	10.0
Glove Length (inch/cm)	12"/30.5	12"/30.5	12"/30.5	12"/30.5	12"/30.5	12"/30.5	12"/30.5	12"/30.5
Width of Palm (mm)	80	87	94	98	109	114	120	128
Middle finger length (mm)	70.8	72.4	80.6	83.9	86.0	90.1	92.7	96.2
Finger Tip Thickness	0.16 mm (6.3 mil)	0.16 mm (6.3 mil)	0.16 mm (6.3 mil)	0.16 mm (6.3 mil)	0.16 mm (6.3 mil)	0.16 mm (6.3 mil)	0.16 mm (6.3 mil)	0.16 mm (6.3 mil)
Palm Thickness	0.13 mm (5.1 mil)	0.13 mm (5.1 mil)	0.13 mm (5.1 mil)	0.13 mm (5.1 mil)	0.13 mm (5.1 mil)	0.13 mm (5.1 mil)	0.13 mm (5.1 mil)	0.13 mm (5.1 mil)
Cuff Thickness	0.10 mm (3.9 mil)	0.10 mm (3.9 mil)	0.10 mm (3.9 mil)	0.10 mm (3.9 mil)	0.10 mm (3.9 mil)	0.10 mm (3.9 mil)	0.10 mm (3.9 mil)	0.10 mm (3.9 mil)

Packaging Data

Triple layer packaging (poly pouch and poly bag plus case liner)

200 hand specific pairs per case: 2 gloves/poly wallet & pouch x 20 sealed pouches per bag x 10 PE bags per lined carton

Packaged in ISO Class 5 Cleanroom

The PUREZERO* HG3 Sterile Nitrile Cleanroom glove sterile pouch is made from PET/PE film pouch and is Plastic Recycling Number/Category 7.

Quality & Regulatory Standards

Compliant to these regulatory standards:
- ISO 9001
- ISO 13485
- ISO 14001

Compliant to these food handling regulatory standards:
- FDA 21 CFR 177-2600
- FDA 21 CFR 180.22
- Commission Regulation (EU) No 10/2011

CE 2797 PPE Category III according to Regulation (EU) 2016/425 EEC
- EN 16523-1:2015+A1:2018
- EN ISO 374-2:2019
- EN ISO 374-4:2019
- EN ISO 374-5:2016 Bacteria, Fungi, and Virus Protection
- EN ISO 374-1:2016+A1:2018/ Type B
- EN ISO 21420:2020 Dexterity Classification: Level 5

Compliant with the REACH regulation
- RoHS 2011/65/EU Annex II

Static Dissipative in Use
- Tested per ANSI/ ESD SP 15.1

Meets USP <800> Guidelines
Meets USP <797> Guidelines

CE 2797
(PPE Cat. III)

ISO 374-5:2016
VIRUS

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

Additional Glove Information

Recommended for use in ISO Class 3 or higher and Grade A/B cleanrooms.

Tested against 29 chemicals, 14 chemotherapy drugs and Fentanyl.

Made in Thailand

Declaration of Conformity (DoC), Certificates of Analysis (COA) and Certificates of Processing (COP) for every production lot available online at **halyardhealth.com/information**

Storage Instructions

HALYARD* **PURE**ZERO* Nitrile Gloves shall be stored in conditions where the product is kept dry (away from moisture), away from direct sunlight, away from sources of heat, and away from radioactive sources.

Ordering Information

HALYARD* PUREZERO*** **HG3 WHITE STERILE NITRILE GLOVES, HAND SPECIFIC, SMOOTH GRIP, TEXTURED FINGERTIPS AND PALMS**

Size	Code	Size	Code
6.0	CLN323260	8.0	CLN323280
6.5	CLN323265	8.5	CLN323285
7.0	CLN323270	9.0	CLN323290
7.5	CLN323275	10.0	CLN323210

For additional information or samples, contact your local distributor or visit **www.purezerogloves.com**

1 Not formulated with these commonly used vulcanizing chemicals: Sulfur, Thiurams, Thiazoles, Guanidines and Carbamates.
2 Tested per ASTM D412
This fact sheet has been created using the most recent information. In the interest of continuous improvement, the characteristics of the products may change without prior notice.

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Owens & Minor

HALYARD* is an Owens & Minor brand

Owens & Minor | HALYARD

Safeskin Medical & Scientific (Thailand) Ltd. 200 Moo 8 Kanchanavarich Road Tambol Prik, Amphur Sadao Songkhla 90120 Thailand

CERTIFICATE OF ANALYSIS

Product Description : HALYARD* PUREZERO* HG3 Sterile White Nitrile Gloves - Pairs

Catalog Numbers : CLN323260, CLN323265, CLN323270, CLN323275, CLN323280, CLN323285, CLN323290, CLN323210

OMH Code : 48192-82, 48193-82, 48194-82, 48195-82, 48196-82, 48197-82, 48198-82, 48199-82

Batches : SM503225X to SM505925X

Total Cases per Lot : 532
Date of Manufacture : Feb-25
Expiration Date : 2028-01-30

Physical Test Data

		Visual Defects			Tensile (MPa)	Elongation (%)	Median Force at Break (N)
	Watertight	Minor	Negligible	Dimensions	Pre Aging	Pre Aging	Pre Aging
Sample Size :	330	330	330	91	39	39	39
AQL Level :	0.65	2.5	4.0	4.0	4.0	4.0	4.0
Failures Allowed per AQL :	2	13	19	7	3	3	3
Failures :	0	0	0	0	0	0	0
Inspection Results :	Accept	Accept	Accept	Accept	Accept	Accept	Accept
				Averages:	41.19	590	14.63

Test Methods : Water tight test ASTM D 5151, EN 455-1, Elongation and Tensile ASTM D 412, ASTM D 6319, Median Force at Break EN 455-2, Dimension ASTM D 6319, EN 455-2

Particle Test Data

Particle	Total Particles/cm²	Inspection Result
#/cm2:	852	Accept

Particulate Level (≥0.5 microns) #/cm²

Test Method : IEST-RP-CC005.4

Extractable Ion Test Data

	Anions Results			Cations Results				
	Chloride	Nitrate	Sulfate	Sodium	Ammonium	Potassium	Magnesium	Calcium
Units :	Cl^-	NO_3^-	SO_4^{-2}	Na^+	NH_4^+	K^+	Mg^{+2}	Ca^{+2}
µg/g glove	28.183	4.575	2.105	3.607	1.980	1.533	1.129	16.688
Inspection Result :	Accept	Accept	Accept	Accept	Accept	Accept	Accept	Accept

Trace Element Results

	Zinc
Units :	Zn
µg/g glove	6.29
Inspection Result :	Accept

Test Method : IEST-RP-CC005.4

Endotoxin Test Data

Test Result : 0.4169 **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.

Review By : 12 Mar 2025

(QA Sr. Manager)

FORM-28014/8

STERIS: Certificate Of Processing

Prepared For SAFESKIN MEDICAL & SCIENTIFIC (THAILAND) LTD (10230)

Gamma Process Run ID 3023-5024A

Product Code	Lot Number	Quantity	UOM
HALYARD*PUREZERO*HG3 STERILE WHITE NITRILE GLOVES-PAIRS	N/A	69	Case

Validation Reference Number: 0.0969 Rev01

Processing Run Start Date 02-Apr-2025 11:27 PM

Processing Run End Date 03-Apr-2025 9:06 AM

Specified Dose Range (kGy)	**25.0 - 50.0**	**Calculated Min Dose (kGy)**	**26.7**
Reference Dose Range (kGy)	**30.0 - 45.6**	**Calculated Max Dose (kGy)**	**38.0**

PO Number: 4027025838

Product meets Customer specifications; zero nonconformities occurred during this irradiation run.

Other Information

Refer Attachment 1

Signature Manifest

Reviewed and E-Signed By: **Nursyuhada Shoib (Quality Analyst)**

DateTime Esigned 03-Apr-2025 5:56 PM

Document Content Revision 1

Operating facility is in compliance with applicable regulations providing services under a certified quality system which meets the requirements of FDA QSR, EN/ISO 13485, ISO 9001 current version and is in alignment with EN/ISO 11137 current version.

Processing Location

Synergy Sterilisation (M) Sdn Bhd
Plot 203, Kuala Ketil Industrial Park
09300 Kuala Ketil
Kedah Malaysia
Phone: (60)44152111

STERIS

Applied Sterilization Technologies

Attachment 1

Run ID Process :3023-5024A

HALYARD* PUREZERO* HG3 Sterile White Nitrile Gloves - Pairs

Catalog Numbers : CLN323260, CLN323265, CLN323270, CLN323275, CLN323280, CLN323285, CLN323290,CLN323210.

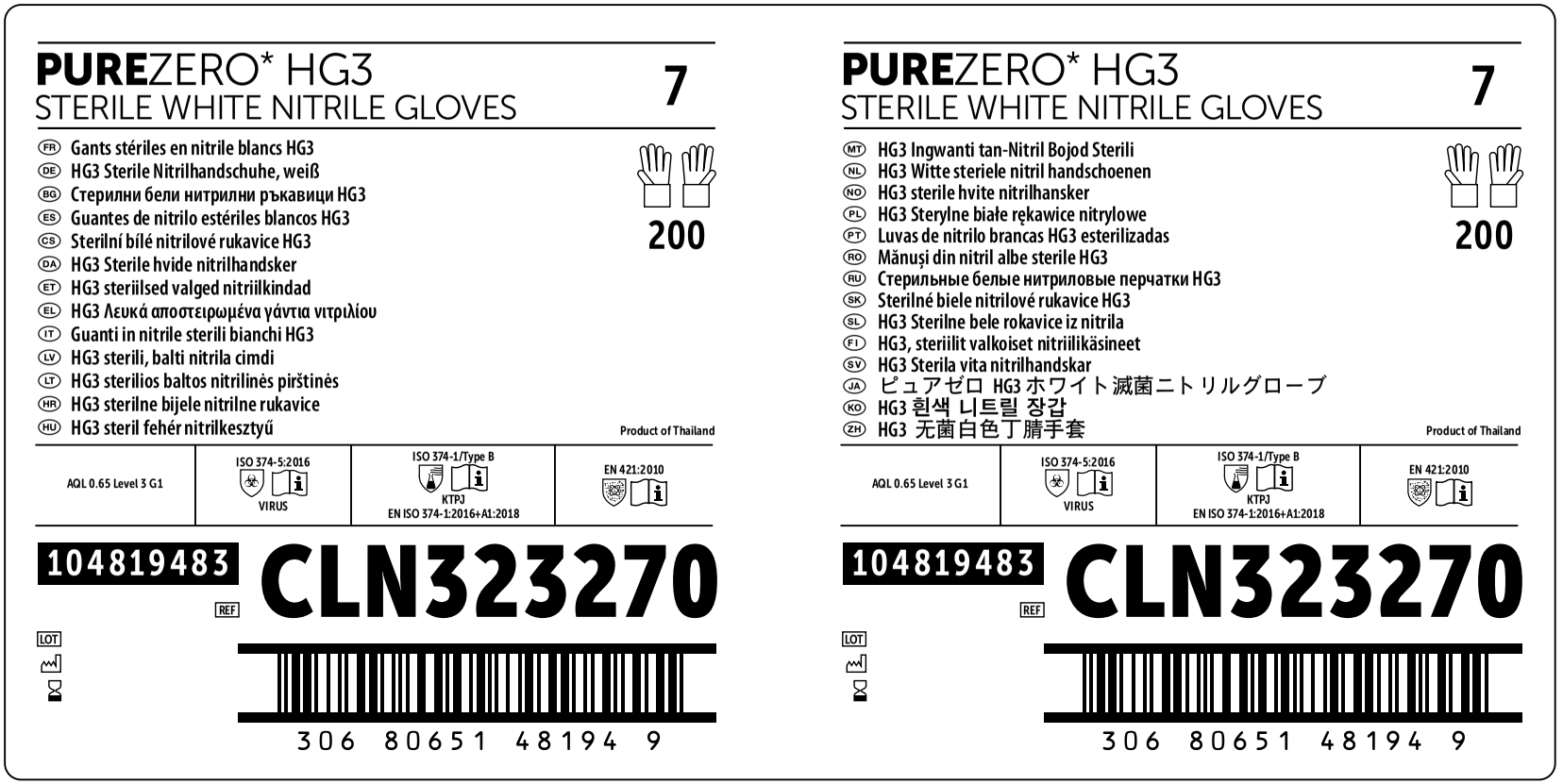
OMH Code : 48192-82, 48193-82, 48194-82, 48195-82, 48196-82, 48197-82, 48198-82, 48199-82

Catalog Number(s)	Lot No./Batch No.	Quantity
CLN323275 48195-82	NIL/SM506225X	51
CLN323290 48198-82	NIL/SM506225X	18

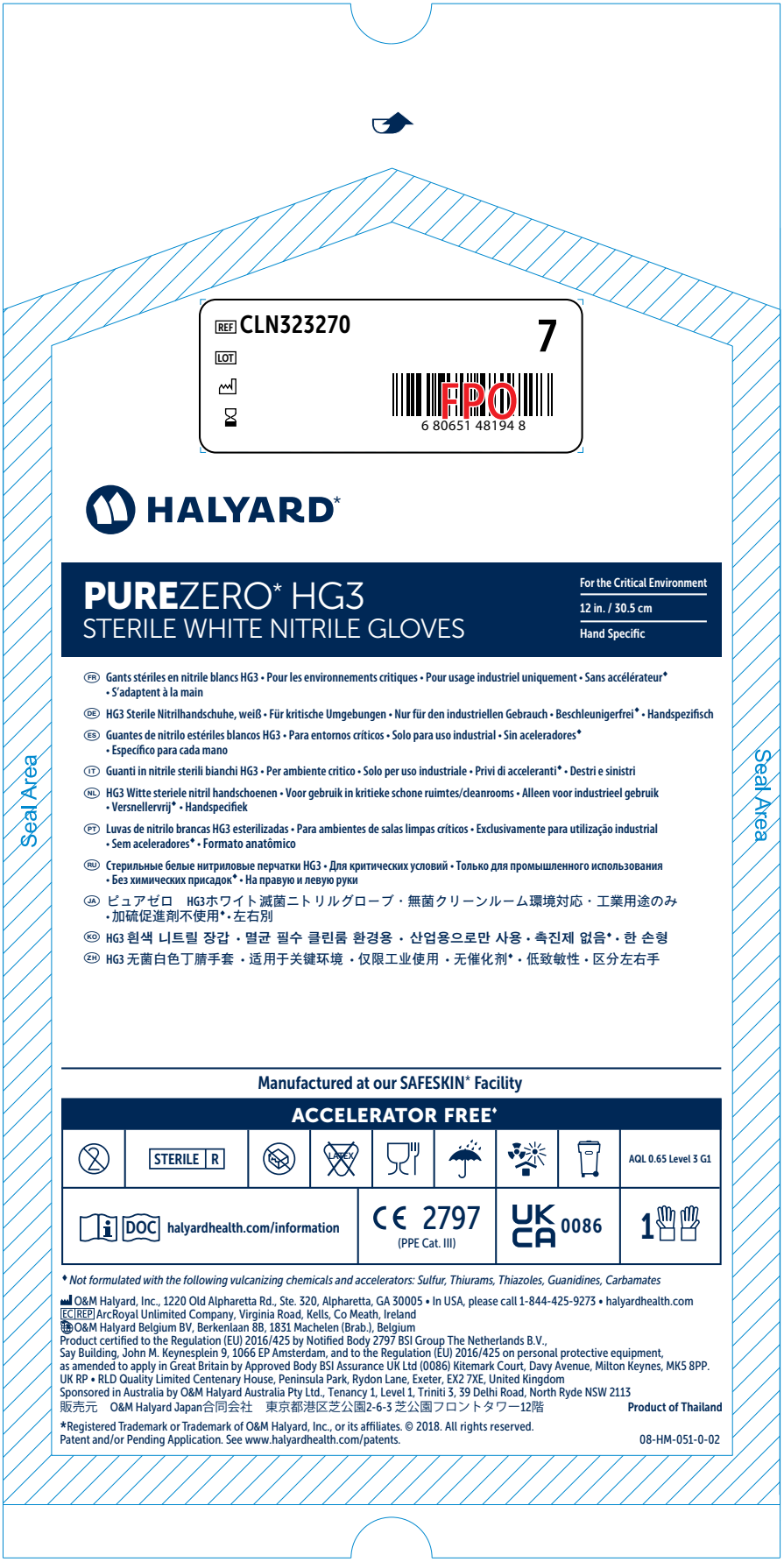
PACKAGE LABELS



CASE LABEL

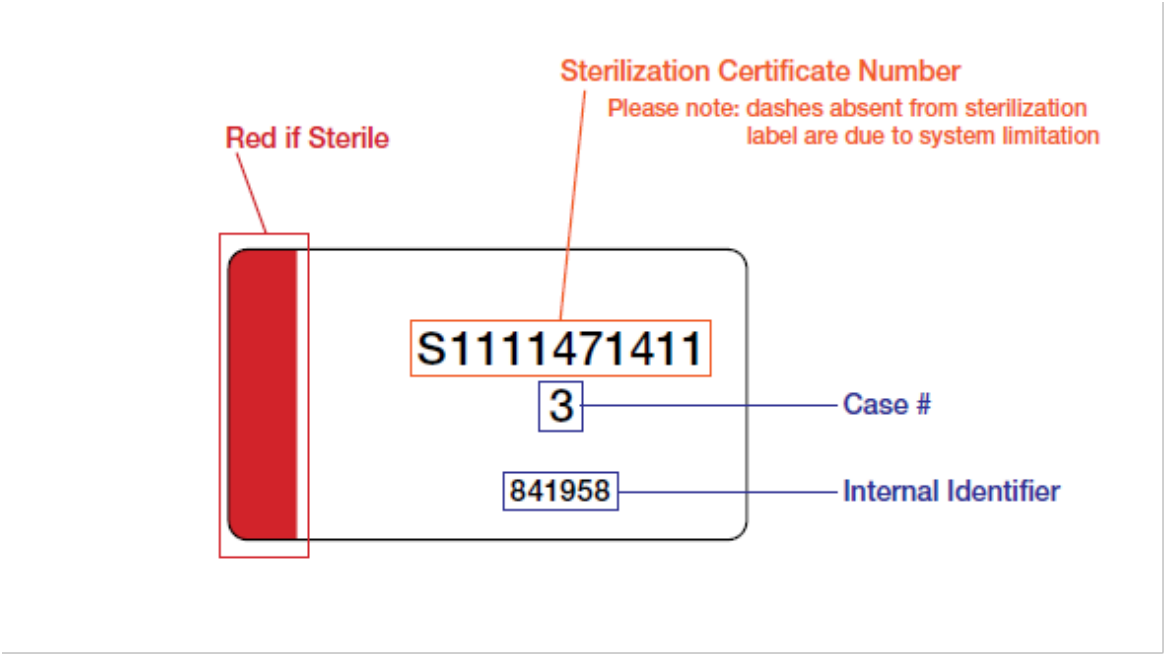

PUREZERO* HG3
STERILE WHITE NITRILE GLOVES
7
FR Gants stériles en nitrile blancs HG3
DE HG3 Sterile Nitrilhandschuhe, weiß
BG Стерилни бели нитрилни ръкавици HG3
ES Guantes de nitrilo estériles blancos HG3
CS Sterilní bílé nitrilové rukavice HG3
DA HG3 Sterile hvide nitrilhandsker
ET HG3 steriilsed valged nitriilkindad
EL HG3 Λευκά αποστειρωμένα γάντια νιτριλίου
IT Guanti in nitrile sterili bianchi HG3
LV HG3 sterili, balti nitrila cimdi
LT HG3 sterilios baltos nitrilinės pirštinės
HR HG3 sterilne bijele nitrilne rukavice
HU HG3 steril fehér nitrilkesztyű
200
Product of Thailand
AQL 0.65 Level 3 G1
ISO 374-5:2016
VIRUS
ISO 374-1/Type B
KTPJ
EN ISO 374-1:2016+A1:2018
EN 421:2010
104819483
REF
CLN323270
LOT
306 80651 48194 9
PUREZERO* HG3
STERILE WHITE NITRILE GLOVES
7
MT HG3 Ingwanti tan-Nitril Bojod Sterili
NL HG3 Witte steriele nitril handschoenen
NO HG3 sterile hvite nitrilhansker
PL HG3 Sterylne białe rękawice nitrylowe
PT Luvas de nitrilo brancas HG3 esterilizadas
RO Mănuși din nitril albe sterile HG3
RU Стерильные белые нитриловые перчатки HG3
SK Sterilné biele nitrilové rukavice HG3
SL HG3 Sterilne bele rokavice iz nitrila
FI HG3, steriilit valkoiset nitriilikäsineet
SV HG3 Sterila vita nitrilhandskar
JA ピュアゼロ HG3 ホワイト滅菌ニトリルグローブ
KO HG3 흰색 니트릴 장갑
ZH HG3 无菌白色丁腈手套
200
Product of Thailand
AQL 0.65 Level 3 G1
ISO 374-5:2016
VIRUS
ISO 374-1/Type B
KTPJ
EN ISO 374-1:2016+A1:2018
EN 421:2010
104819483
REF
CLN323270
LOT
306 80651 48194 9

POUCH LABEL


REF CLN323270
7
LOT
FPO
6 80651 48194 8
HALYARD*
PUREZERO* HG3
STERILE WHITE NITRILE GLOVES
For the Critical Environment
12 in. / 30.5 cm
Hand Specific
FR Gants stériles en nitrile blancs HG3 • Pour les environnements critiques • Pour usage industriel uniquement • Sans accélérateur♦ • S'adaptent à la main
DE HG3 Sterile Nitrilhandschuhe, weiß • Für kritische Umgebungen • Nur für den industriellen Gebrauch • Beschleunigerfrei♦ • Handspezifisch
ES Guantes de nitrilo estériles blancos HG3 • Para entornos críticos • Solo para uso industrial • Sin aceleradores♦ • Específico para cada mano
IT Guanti in nitrile sterili bianchi HG3 • Per ambiente critico • Solo per uso industriale • Privi di acceleranti♦ • Destri e sinistri
NL HG3 Witte steriele nitril handschoenen • Voor gebruik in kritieke schone ruimtes/cleanrooms • Alleen voor industrieel gebruik • Versnellervrij♦ • Handspecifiek
PT Luvas de nitrilo brancas HG3 esterilizadas • Para ambientes de salas limpas críticos • Exclusivamente para utilização industrial • Sem aceleradores♦ • Formato anatômico
RU Стерильные белые нитриловые перчатки HG3 • Для критических условий • Только для промышленного использования • Без химических присадок♦ • На правую и левую руки
JA ピュアゼロ HG3ホワイト滅菌ニトリルグローブ・無菌クリーンルーム環境対応・工業用途のみ・加硫促進剤不使用♦・左右別
KO HG3 흰색 니트릴 장갑 · 멸균 필수 클린룸 환경용 · 산업용으로만 사용 · 촉진제 없음♦ · 한 손형
ZH HG3 无菌白色丁腈手套 · 适用于关键环境 · 仅限工业使用 · 无催化剂♦ · 低致敏性 · 区分左右手
Seal Area
Seal Area
Manufactured at our SAFESKIN* Facility
ACCELERATOR FREE♦
STERILE R
LATEX
AQL 0.65 Level 3 G1
DOC halyardhealth.com/information
CE 2797
(PPE Cat. III)
UKCA 0086
1
♦ Not formulated with the following vulcanizing chemicals and accelerators: Sulfur, Thiurams, Thiazoles, Guanidines, Carbamates
O&M Halyard, Inc., 1220 Old Alpharetta Rd., Ste. 320, Alpharetta, GA 30005 • In USA, please call 1-844-425-9273 • halyardhealth.com
EC REP ArcRoyal Unlimited Company, Virginia Road, Kells, Co Meath, Ireland
O&M Halyard Belgium BV, Berkenlaan 8B, 1831 Machelen (Brab.), Belgium
Product certified to the Regulation (EU) 2016/425 by Notified Body 2797 BSI Group The Netherlands B.V., Say Building, John M. Keynesplein 9, 1066 EP Amsterdam, and to the Regulation (EU) 2016/425 on personal protective equipment, as amended to apply in Great Britain by Approved Body BSI Assurance UK Ltd (0086) Kitemark Court, Davy Avenue, Milton Keynes, MK5 8PP. UK RP • RLD Quality Limited Centenary House, Peninsula Park, Rydon Lane, Exeter, EX2 7XE, United Kingdom
Sponsored in Australia by O&M Halyard Australia Pty Ltd., Tenancy 1, Level 1, Triniti 3, 39 Delhi Road, North Ryde NSW 2113
販売元 O&M Halyard Japan合同会社 東京都港区芝公園2-6-3 芝公園フロントタワー12階
Product of Thailand
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Patent and/or Pending Application. See www.halyardhealth.com/patents.
08-HM-051-0-02

STERILIZATION/IRRADIATION LABEL

Irradiation Label Format – HALYARD* PUREZERO**HG3 Sterile Nitrile Gloves**



PACKAGING CERTIFICATES



POUCH

TIP CORPORATION SDN BHD Reg. No: 198901010228 (187530-A)
Lot 2288, Jalan P4/8, Bandar Teknologi Kajang,
Semenyih, 43500 Selangor, Malaysia.
Tel: +603-8723 7889
E-mail: sales@tipcorp.com Website: www.tipcorp.com
SST No: W24-1808-21017734

FORM QC-21

LPC Test Report

Item:	C/R EASY PEEL-PRT	Lot No:	K34514(Outgoing A)
Testing Date:	26.10.2020	Size:	152.4 X 304.8 X 0.07MM
Customer :	SAFESKIN	Method:	BY HAND
		Machine:	BS-5

Blank Particle Counts (Cumulative)

	Blank Cts 1 Counts	Blank Cts 2 Counts	Blank Ct 3 Counts
0.5 micron	2.0	4.0	3.0
1.0 micron	0.0	0.0	0.0
2.0 micron	0.0	0.0	0.0
5.0 micron	0.0	0.0	0.0

Average Blank (A) Counts: 3.0 / 0.0 / 0.0 / 0.0 Total counts 3.0

Sample Particle Counts (Cumulative)

	Washed Cts 1 Counts	Washed Cts 2 Counts	Washed Cts 3 Counts
0.5 micron	16.0	13.0	10.0
1.0 micron	3.0	4.0	2.0
2.0 micron	0.0	1.0	0.0
5.0 micron	0.0	0.0	0.0

Average Washed (B) Counts: 13.0 / 3.0 / 0.3 Total counts 16.3 / 0.0

(C) Wet Particle Counts

Ave.Washed - Ave.Blank (C = B - A)

0.5 micron	10.0
1.0 micron	3.0
2.0 micron	0.3
5.0 micron	0.0

(D) Total Vol. for Washed Sample

Washed Volume (ml) 400 ml

(E) Sampling size 5 ml

(F) Total Area of Plastic

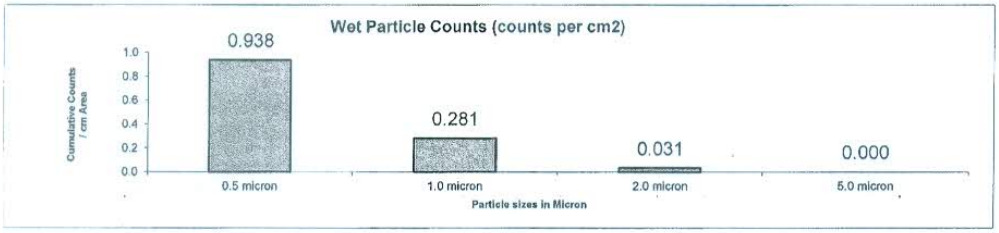
Total Washed Area 853 cm^2

(G) Particle counts per ml (G = C / E)

0.5 micron	2.000
1.0 micron	0.600
2.0 micron	0.067
5.0 micron	0.000

(H) Total Wet Part Cts / cm^2 (H = G X D / F)

0.5 micron	0.938
1.0 micron	0.281
2.0 micron	0.031
5.0 micron	0.000



Cts/cm2

Total Particle (≥0.5) : 1.250

SPEC: ≥ 0.5u ≤100 COUNTS/CM^2

Tested By : SUHAIDA
Date : 26.10.2020

Verified By : RUDY
Date : 26.10.2020

TIP CORPORATION SDN BHD QA RUDY

POUCH

TIP CORPORATION SDN BHD Reg. No: 198901010228 (187530-A)
Lot 2288, Jalan P4/8, Bandar Teknologi Kajang,
Semenyih, 43500 Selangor, Malaysia.
Tel: +603-8723 7889
E-mail: sales@tipcorp.com Website: www.tipcorp.com
SST No: W24-1808-21017734

FORM QC-21

LPC Test Report

Item:	C/R EASY PEEL-PRT	Lot No:	K34514(Outgoing B)
Testing Date:	26.10.2020	Size:	152.4 X 304.8 X 0.07MM
Customer :	SAFESKIN	Method:	BY HAND
		Machine:	BS-5

Blank Particle Counts (Cumulative)

	Blank Cts 1 Counts	Blank Cts 2 Counts	Blank Ct 3 Counts	Average Blank (A) Counts	
0.5 micron	5.0	2.0	4.0	3.7	
1.0 micron	0.0	0.0	0.0	0.0	
2.0 micron	0.0	0.0	0.0	0.0 Total counts	3.7
5.0 micron	0.0	0.0	0.0	0.0	

Washed Sample Particle Counts (Cumulative)

	Washed Cts 1 Counts	Washed Cts 2 Counts	Washed Cts 3 Counts	Average Washed (B) Counts	
0.5 micron	19.0	15.0	17.0	17.0	
1.0 micron	4.0	3.0	2.0	3.0	
2.0 micron	1.0	1.0	1.0	1.0 Total counts	21.7
5.0 micron	1.0	1.0	0.0	0.7	

(C) Wet Particle Counts

Ave.Washed - Ave.Blank (C = B - A)	
0.5 micron	13.3
1.0 micron	3.0
2.0 micron	1.0
5.0 micron	0.7

(D) Total Vol. for Washed Sample

Washed Volume (ml)	400 ml

(E) Sampling size 5 ml

(F) Total Area of Plastic

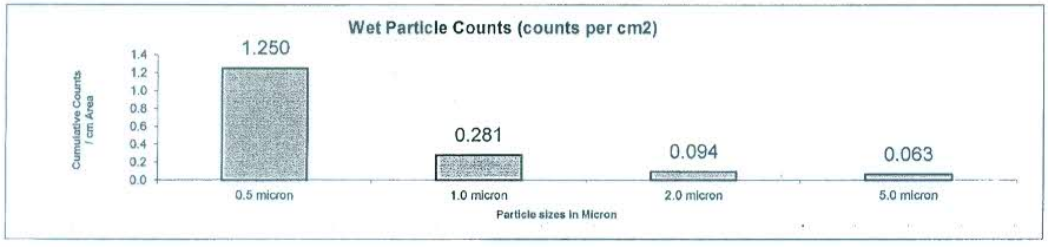
Total Washed Area	853 cm^2

(G) Particle counts per ml (G = C / E)

0.5 micron	2.667
1.0 micron	0.600
2.0 micron	0.200
5.0 micron	0.133

(H) Total Wet Part Cts / cm^2 (H = G X D / F)

0.5 micron	1.250
1.0 micron	0.281
2.0 micron	0.094
5.0 micron	0.063



Cts/cm2

Total Particle (≥0.5) : 1.688

SPEC: ≥ 0.5u ≤100 COUNTS/CM^2

Tested By : SUHAIDA
Date : 26.10.2020

Verified By : RUDY
Date : 26.10.2020

TIP CORPORATION SDN. BHD. QA RUDY

POUCH

TIP CORPORATION SDN BHD Reg. No: 198901010228 (187530-A)

Lot 2288, Jalan P4/8, Bandar Teknologi Kajang,
Semenyih, 43500 Selangor, Malaysia.
Tel: +603-8723 7889
E-mail: sales@tipcorp.com Website: www.tipcorp.com
SST No: W24-1808-21017734

FORM QC-21

LPC Test Report

Item: C/R EASY PEEL-PRT
Testing Date: 26.10.2020
Customer : SAFESKIN

Lot No: K34514(Outgoing C)
Size: 152.4 X 304.8 X 0.07MM
Method: BY HAND
Machine: BS-5

Blank Particle Counts (Cumulative)

	Blank Cts 1 Counts	Blank Cts 2 Counts	Blank Ct 3 Counts
0.5 micron	4.0	3.0	3.0
1.0 micron	0.0	0.0	0.0
2.0 micron	0.0	0.0	0.0
5.0 micron	0.0	0.0	0.0

Average Blank (A) Counts: 3.3, 0.0, 0.0 Total counts 3.3, 0.0

Washed Sample Particle Counts (Cumulative)

	Washed Cts 1 Counts	Washed Cts 2 Counts	Washed Cts 3 Counts
0.5 micron	20.0	11.0	18.0
1.0 micron	8.0	3.0	3.0
2.0 micron	3.0	1.0	0.0
5.0 micron	0.0	0.0	0.0

Average Washed (B) Counts: 16.3, 4.7, 1.3 Total counts 22.3, 0.0

(C) Wet Particle Counts

Ave.Washed - Ave.Blank (C = B - A)

0.5 micron	13.0
1.0 micron	4.7
2.0 micron	1.3
5.0 micron	0.0

(D) Total Vol. for Washed Sample
Washed Volume (ml) 400 ml

(E) Sampling size 5 ml

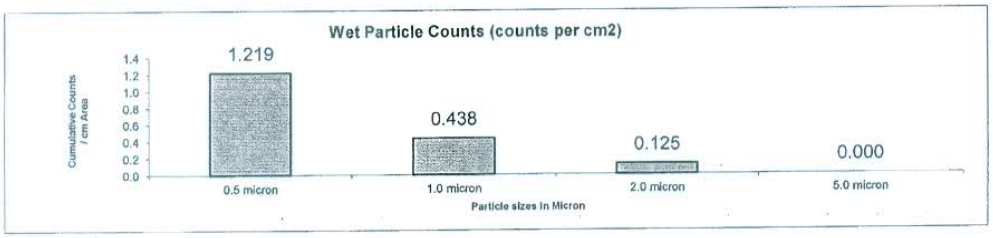
(F) Total Area of Plastic
Total Washed Area 853 cm^2

(G) Particle counts per ml (G = C / E)

0.5 micron	2.600
1.0 micron	0.933
2.0 micron	0.267
5.0 micron	0.000

(H) Total Wet Part Cts / cm^2 (H = G X D / F)

0.5 micron	1.219
1.0 micron	0.438
2.0 micron	0.125
5.0 micron	0.000



Cts/cm2

Total Particle (≥0.5) : 1.782

SPEC: ≥ 0.5u ≤100 COUNTS/CM^2

Tested By : SUHAIDA
Date : 26.10.2020

Verified By : RUDY
Date : 26.10.2020

TIP CORPORATION SDN BHD QA RUDY

WALLET

TIP CORPORATION SDN BHD Reg. No: 198901010228 (187530-A)

Lot 2288, Jalan P4/8, Bandar Teknologi Kajang,
Semenyih, 43500 Selangor, Malaysia.
Tel: +603-8723 7889
E-mail: sales@tipcorp.com Website: www.tipcorp.com
SST No: W24-1808-21017734

FORM QC-21

LPC Test Report

Item:	C/R LDPE WALLET	Lot No:	K55532
Testing Date:	09.11.2020	Size:	220X340X0.06MM
Customer :	SAFESKIN	Method:	BY HAND
			KCCS-17

Blank Particle Counts (Cumulative)

	Blank Cts 1 Counts	Blank Cts 2 Counts	Blank Ct 3 Counts	Average Blank (A) Counts	
0.5 micron	1.0	0.0	1.0	0.7	
1.0 micron	0.0	0.0	0.0	0.0	
2.0 micron	0.0	0.0	0.0	0.0	
0.5 micron	0.0	0.0	0.0	0.0 Total counts	0.7

ash Sample Particle Counts (Cumulative)

	Washed Cts 1 Counts	Washed Cts 2 Counts	Washed Cts 3 Counts	Average Washed (B) Counts	
0.5 micron	60.0	58.0	52.0	56.7	
1.0 micron	13.0	21.0	18.0	17.3	
2.0 micron	4.0	5.0	6.0	5.0	
0.5 micron	2.0	0.0	3.0	1.7 Total counts	80.7

(C) Wet Particle Counts

Ave.Washed - Ave.Blank (C = B - A)

0.5 micron	56.0
1.0 micron	17.3
2.0 micron	5.0
0.5 micron	1.7

(D) Total Vol. for Washed Sample

Washed Volume (ml) 400 ml

(E) Sampling size 5 ml

(F) Total Area of Plastic

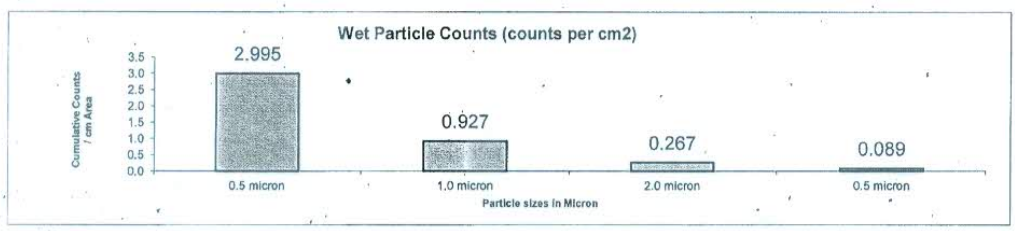
Total Washed Area 1496 cm^2

(G) Particle counts per ml (G = C / E)

.nicron	11.200
1.0 micron	3.467
2.0 micron	1.000
0.5 micron	0.333

(H) Total Wet Part Cts / cm^2 (H = G X D / F)

0.5 micron	2.995
1.0 micron	0.927
2.0 micron	0.267
0.5 micron	0.089



Cts/cm2

Total Particle (≥0.5) : **4.278**

SPEC: ≥ 0.5u ≤100 COUNTS/CM^2

Tested By : SUHAIDA
Date : 09.11.2020

Verified By : RUDY
Date : **09.11.2020**

WALLET

TIP CORPORATION SDN BHD Reg. No: 198901010228 (187530-A)

Lot 2288, Jalan P4/8, Bandar Teknologi Kajang,
Semenyih, 43500 Selangor, Malaysia.
Tel: +603-8723 7889
E-mail: sales@tipcorp.com Website: www.tipcorp.com
SST No: W24-1808-21017734

FORM QC-21

LPC Test Report

Item:	C/R LDPE WALLET	Lot No:	K55753
Testing Date:	09.11.2020	Size:	220X340X0.06MM
Customer :	SAFESKIN	Method:	BY HAND
			KCCS-17

Blank Particle Counts (Cumulative)

	Blank Cts 1 Counts	Blank Cts 2 Counts	Blank Ct 3 Counts	Average Blank (A) Counts	
0.5 micron	6.0	0.0	1.0	2.3	
1.0 micron	2.0	0.0	0.0	0.7	
2.0 micron	0.0	0.0	0.0	0.0	
5.0 micron	0.0	0.0	0.0	0.0 Total counts	3.0

ash Sample Particle Counts (Cumulative)

	Washed Cts 1 Counts	Washed Cts 2 Counts	Washed Cts 3 Counts	Average Washed (B) Counts	
0.5 micron	56.0	45.0	72.0	57.7	
1.0 micron	16.0	23.0	32.0	23.7	
2.0 micron	4.0	6.0	9.0	6.3	
5.0 micron	2.0	0.0	2.0	1.3 Total counts	89.0

(C) Wet Particle Counts

Ave.Washed - Ave.Blank (C = B - A)

0.5 micron	55.3
1.0 micron	23.0
2.0 micron	6.3
5.0 micron	1.3

(D) Total Vol. for Washed Sample

Washed Volume (ml) 400 ml

(E) Sampling size 5 ml

(F) Total Area of Plastic

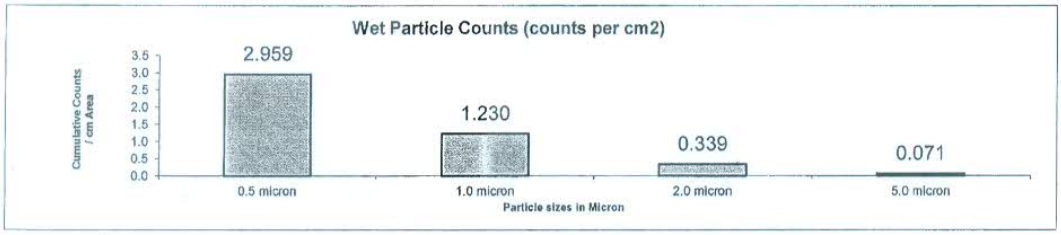
Total Washed Area 1496 cm^2

(G) Particle counts per ml (G = C / E)

micron	11.067
1.0 micron	4.600
2.0 micron	1.267
5.0 micron	0.267

(H) Total Wet Part Cts / cm^2 (H = G X D / F)

0.5 micron	2.959
1.0 micron	1.230
2.0 micron	0.339
5.0 micron	0.071



Cts/cm2

Total Particle (≥0.5) : 4.599

SPEC: ≥ 0.5u ≤100 COUNTS/CM^2

Tested By : SUHAIDA
Date : 09.11.2020

Verified By : RUDY
Date : 09.11.2020

TIP CORPORATION SDN BHD QA RUDY

WALLET

TIP CORPORATION SDN BHD Reg. No. 198901010228 (187530-A)
Lot 2288, Jalan P4/8, Bandar Teknologi Kajang,
Semenyih, 43500 Selangor, Malaysia.
Tel: +603-8723 7889
E-mail: sales@tipcorp.com Website: www.tipcorp.com
SST No: W24-1808-21017734

FORM QC-21

LPC Test Report

Item:	C/R LDPE WALLET	Lot No:	K55752
Testing Date:	09.11.2020	Size:	220X340X0.06MM
Customer :	SAFESKIN	Method:	BY HAND
			KCCS-17

Blank Particle Counts (Cumulative)

	Blank Cts 1 Counts	Blank Cts 2 Counts	Blank Ct 3 Counts	Average Blank (A) Counts	
0.5 micron	9.0	6.0	5.0	6.7	
1.0 micron	0.0	0.0	1.0	0.3	
2.0 micron	0.0	0.0	0.0	0.0	
5.0 micron	0.0	0.0	0.0	0.0 Total counts	7.0

Wash Sample Particle Counts (Cumulative)

	Washed Cts 1 Counts	Washed Cts 2 Counts	Washed Cts 3 Counts	Average Washed (B) Counts	
0.5 micron	65.0	96.0	98.0	86.3	
1.0 micron	22.0	25.0	28.0	25.0	
2.0 micron	8.0	6.0	6.0	6.7	
5.0 micron	2.0	3.0	0.0	1.7 Total counts	119.7

(C) Wet Particle Counts

Ave.Washed - Ave.Blank (C = B - A)	
0.5 micron	79.7
1.0 micron	24.7
2.0 micron	6.7
5.0 micron	1.7

(D) Total Vol. for Washed Sample

Washed Volume (ml) 400 ml

(E) Sampling size 5 ml

(F) Total Area of Plastic

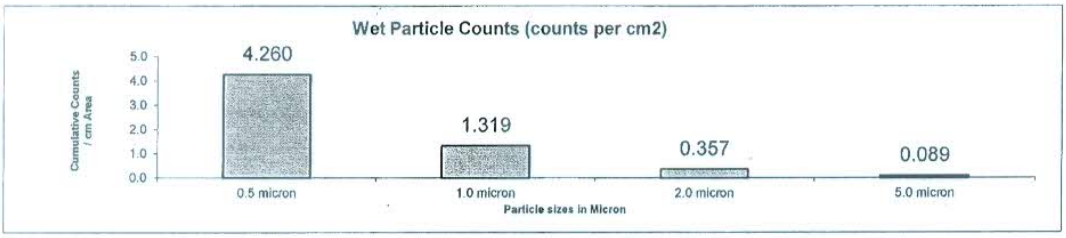
Total Washed Area 1496 cm^2

(G) Particle counts per ml (G = C / E)

0.5 micron	15.933
1.0 micron	4.933
2.0 micron	1.333
5.0 micron	0.333

(H) Total Wet Part Cts / cm^2 (H = G X D / F)

0.5 micron	4.260
1.0 micron	1.319
2.0 micron	0.357
5.0 micron	0.089



Cts/cm2

Total Particle (≥0.5) : 6.025

SPEC: ≥ 0.5u ≤100 COUNTS/CM^2

Tested By : SUHAIDA
Date : 09.11.2020

Verified By : RUDY
Date : 09.11.2020

TIP CORPORATION SDN BHD SUHAIDA RUDY

WALLET

TIP CORPORATION SDN BHD Reg. No: 198901010228 (187530-A)
Lot 2288, Jalan P4/8, Bandar Teknologi Kajang,
Semenyih, 43500 Selangor, Malaysia.
Tel: +603-8723 7889
E-mail: sales@tipcorp.com Website: www.tipcorp.com
SST No: W24-1808-21017734

FORM QC-21

LPC Test Report

Item:	C/R LDPE WALLET	Lot No:	K55751
Testing Date:	09.11.2020	Size:	18"X0.06MM(18"X18")
Customer :	SAFESKIN	Method:	BY SHAKER
		Machine:	KL-2

Blank Particle Counts (Cumulative)

	Blank Cts 1 Counts	Blank Cts 2 Counts	Blank Ct 3 Counts	Average Blank (A) Counts	
0.5 micron	2.0	1.0	0.0	1.0	
1.0 micron	0.0	1.0	0.0	0.3	
2.0 micron	0.0	0.0	0.0	0.0	
5.0 micron	0.0	0.0	0.0	0.0 Total counts	1.3

Wash Sample Particle Counts (Cumulative)

	Washed Cts 1 Counts	Washed Cts 2 Counts	Washed Cts 3 Counts	Average Washed (B) Counts	
0.5 micron	289.0	258.0	286.0	277.7	
1.0 micron	120.0	92.0	111.0	107.7	
2.0 micron	39.0	51.0	33.0	41.0	
5.0 micron	9.0	5.0	6.0	6.7 Total counts	433.0

(C) Wet Particle Counts

Ave.Washed - Ave.Blank (C = B - A)	
0.5 micron	276.7
1.0 micron	107.3
2.0 micron	41.0
5.0 micron	6.7

(D) Total Vol. for Washed Sample

Washed Volume (ml) 1000 ml

(E) Sampling size 5 ml

(F) Total Area of Plastic

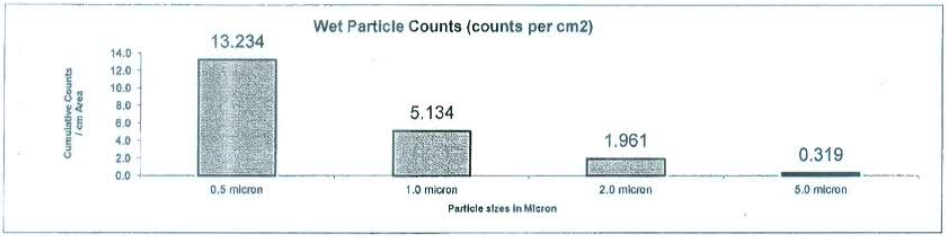
Total Washed Area 4181 cm^2

(G) Particle counts per ml (G = C / E)

0.5 micron	55.333
1.0 micron	21.467
2.0 micron	8.200
5.0 micron	1.333

(H) Total Wet Part Cts / cm^2 (H = G X D / F)

0.5 micron	13.234
1.0 micron	5.134
2.0 micron	1.961
5.0 micron	0.319



Cts/cm2

Total Particle (≥0.5) : 20.649

SPEC: ≥ 0.5u ≤30 COUNTS/CM^2
SPEC: ≥ 0.5u ≤10 COUNTS/CM^2

Tested By : SUHAIDA
Date : 09.11.2020

Verified By : RUDY
Date : 09.11.2020

TIP CORPORATION SDN BHD QA RUDY

WALLET

TIP CORPORATION SDN BHD Reg. No: 198901010228 (187530-A)
Lot 2288, Jalan P4/8, Bandar Teknologi Kajang,
Semenyih, 43500 Selangor, Malaysia.
Tel: +603-8723 7889
E-mail: sales@tipcorp.com Website: www.tipcorp.com
SST No: W24-1808-21017734

FORM QC-21

LPC Test Report

Item:	C/R LDPE WALLET	Lot No:	K55749
Testing Date:	09.11.2020	Size:	220X340X0.06MM
Customer :	SAFESKIN	Method:	BY HAND
			KCCS-17

Blank Particle Counts (Cumulative)

	Blank Cts 1 Counts	Blank Cts 2 Counts	Blank Ct 3 Counts	Average Blank (A) Counts	
0.5 micron	3.0	3.0	2.0	2.7	
1.0 micron	0.0	0.0	0.0	0.0	
2.0 micron	0.0	0.0	0.0	0.0	
0.5 micron	0.0	0.0	0.0	0.0 Total counts	2.7

ash Sample Particle Counts (Cumulative)

	Washed Cts 1 Counts	Washed Cts 2 Counts	Washed Cts 3 Counts	Average Washed (B) Counts	
0.5 micron	48.0	58.0	72.0	59.3	
1.0 micron	17.0	18.0	14.0	16.3	
2.0 micron	9.0	10.0	3.0	7.3	
0.5 micron	0.0	2.0	0.0	0.7 Total counts	83.7

(C) Wet Particle Counts

Ave.Washed - Ave.Blank (C = B - A)	
0.5 micron	56.7
1.0 micron	16.3
2.0 micron	7.3
0.5 micron	0.7

(D) Total Vol. for Washed Sample

Washed Volume (ml) 400 ml

(E) Sampling size 5 ml

(F) Total Area of Plastic

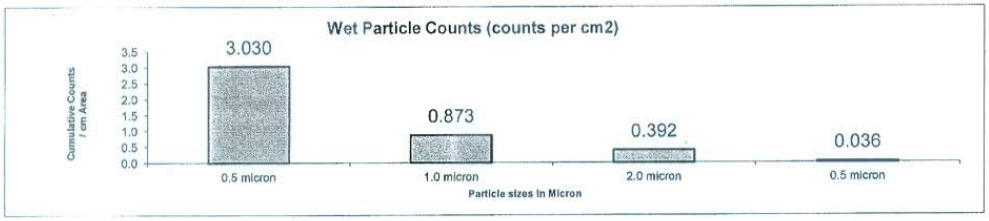
Total Washed Area 1496 cm^2

(G) Particle counts per ml (G = C / E)

micron	11.333
1.0 micron	3.267
2.0 micron	1.467
0.5 micron	0.133

(H) Total Wet Part Cts / cm^2 (H = G X D / F)

0.5 micron	3.030
1.0 micron	0.873
2.0 micron	0.392
0.5 micron	0.036



Cts/cm2

Total Particle (≥0.5) : 4.332

SPEC: ≥ 0.5u ≤100 COUNTS/CM^2

Tested By : SUHAIDA
Date : 09.11.2020

Verified By : RUDY
Date : 09.11.2020

TIP CORPORATION SDN BHD QA RUDY

DOSE MAPPING REPORT



STERIS
Applied Sterilization Technologies

Safeskin Medical & Scientific (Thailand) Ltd

Please find enclosed with our compliments, the completed process validation report in accordance with EN ISO 11137-1:2015 for the products as below:

Product name:	**HALYARD*PUREZERO* HG3 Sterile White Nitrile Gloves - Pairs**
The validation ref no:	**0.0969 Rev 01**

We would appreciate you quoting the product name as above in your PO/DO/reference documents to STERIS, this will aid in identifying your products prior to processing.

Your sincerely,

07 May 2021

Yow Hueh Koon
Malaysia Radiation Validation Manager
STERIS Applied Sterilization Technologies
Tel: +603 3101 3366
Email: huehkoon_yow@steris.com

Customer Acknowledge,

Name :
Designation :
(Please sign this section and scan a copy to huehkoon_yow@steris.com)

STERIS
Applied Sterilization Technologies

DOSE MAPPING REPORT
Safeskin Medical & Scientific (Thailand) Ltd
For

HALYARD*PUREZERO* HG3
Sterile White Nitrile Gloves - Pairs

Validation Report Number : 0.0969 Rev 01

Summary of Performance Qualification:

Customer Name	:	**Safeskin Medical & Scientific (Thailand) Ltd**
Report Ref.	:	**0.0969 Rev 01**
Issue Date	:	**29.04.2021**
Product Description:	:	**HALYARD*PUREZERO* HG3 Sterile White Nitrile Gloves - Pairs**
Type of package:	:	**Corrugated carton**
Carton dimension :	:	**345 mm (L) x 285 mm (W) x 390 mm (H) (15 ctns/tote)**
Method	:	**ISO/ASTM 52303: 2015 (E)**
Reference Standard	:	**EN ISO 11137-1: 2015**
Reference Procedure	:	**QWI 24 Rev 12 Dose Mapping**

Dose Specification:

Minimum dose (kGy)	:	**25.0**
Maximum dose (kGy)	:	**50.0**

Requirement	Minimum specification	Maximum specification
Dose at DRef	30.1	45.6
Dwell time(s)	137	202
Number of Xs	36	53
Correction Factor (Rmin/Rmax)	0.832	1.096

Conclusion

The delivered dose in the product presentation illustrated on page 4 and 5 achieves the requested dose specification of minimum and maximum specification. In order to meet this specification during routine processing the recorded dose at Dref must be as above specification. This incorporates an estimation of uncertainty associated with the measurement system.

Authorisation

	Position	Signature	Date
Prepared by	Malaysia Radiation Validation Manager		06 May 2021
Reviewed by	Operation Management or designee		07/05/2021
Approved by	QA Management or designee		07/05/2021

Note:
It is the responsibility if the Customer to routinely provide product in the presentation and orienation outlined in this report. This performance qualification validates the delivery of radiation doses only. Validation of the sterilisation effect and radiation induced material effects, if any, are not addressed by this qualification.

Product description: **HALYARD*PUREZERO* HG3 Sterile White Nitrile Gloves - Pairs**

Qualification data is obtained by placing 4034 PW Red dosimeters in a defined pattern throughout a STERIS tote loaded with product.

The performance qualification is conducted in triplicate with the dosimeters placed throughout the product to assess both the dose distribution and determine the variability of the absorbed dose measurements, when processed under the same processing conditions.

Following processing the relationship between D_{ref} / D_{min} and D_{ref} / D_{max} are calculated to determine an acceptable D_{ref} processing range.

D_{ref} processing range is calculated by multiplying the R_{min} by the Customer minimum specification and the R_{max} by the customer maximum specification .
During routine processing if the D_{ref} value falls within this range then processing is deemed as meeting the required specification:

D_{ref} Minimum = Expected value of R_{min} x Minimum Dose Required
D_{ref} Maximum = Expected value of R_{max} x Maximum Dose Required

Uncertainty

The specification for D_{ref} incorporates an estimation of the uncertainty associated with the measurement system. Uncertainty has been calculated using the method outlined in ASTM 52303:2015 (E). This method provides a confidence level of 95%

Definitions

D_{ref}	- Reference Dose
D_{min}	- Minimum Dose
D_{max}	- Maximum Dose
R_{min}	- D_{ref} / D_{min} ratio
R_{max}	- D_{ref} / D_{max} ratio
CV%	- Coefficient of Variance
C060	- Cobalt 60

Performance Qualification STERIS Kuala Ketil

Validation Ref : 0.0969 Rev 01

Product: HALYARD*PUREZERO* HG3 Sterile White Nitrile Gloves - Pairs **LOW DENSITY**

Analysis for the Calculation of Release Specification Incorporating Uncertainties

Position	PQ1		PQ2		PQ3		Mean	Stdev	CV	Sum of Squared Differences
A11	32.3		32.3		32.2		32.3	0.06	0.19	0.01
A21	30.9		32.2		31.8		31.6	0.67	2.12	0.89
A31	30.2		30.3		30.8		30.4	0.32	1.05	0.21
A41	32.5		33.2		32.4		32.7	0.44	1.35	0.38
A51	33.2		33.0		33.7		33.3	0.36	1.08	0.26
A15	33.3		34.9		34.7		34.3	0.87	2.54	1.52
A25	32.3		33.2		33.5		33.0	0.62	1.88	0.78
A35	34.4		34.7		34.3		34.5	0.21	0.61	0.09
A45	34.7		34.9		34.8		34.8	0.10	0.29	0.02
A55	35.6		35.4		35.0		35.3	0.31	0.88	0.19
A19	33.1		33.4		**35.1**	√-MAX	33.9	1.08	3.19	2.33
A29	34.2		35.2		33.9		34.4	0.68	1.98	0.93
A39	34.7		35.7		34.8		35.1	0.55	1.57	0.61
A49	**36.1**	√-MAX	35.3		34.9		**35.4**	0.61	1.72	0.75
A59	35.5		**36.1**	√-MAX	34.2		35.3	0.97	2.75	1.89
B11	28.6		28.8		28.9		28.8	0.15	0.52	0.05
B21	**27.8**	√-MIN	**28.7**	√-MIN	**28.4**	√-MIN	**28.3**	0.46	1.63	0.42
B31	28.5		28.8		**28.4**	√-MIN	28.6	0.21	0.73	0.09
B41	31.9		32.0		32.0		32.0	0.06	0.19	0.01
B51	32.6		32.4		32.8		32.6	0.20	0.61	0.08
B15	30.4		31.3		30.9		30.9	0.45	1.46	0.41
B25	29.9		29.8		30.5		30.1	0.38	1.26	0.29
B35	29.8		29.7		28.8		29.4	0.55	1.87	0.61
B45	30.2		30.7		31.5		30.8	0.66	2.14	0.86
B55	31.2		32.2		31.2		31.5	0.58	1.84	0.67
B19	30.1		30.1		30.9		30.4	0.46	1.51	0.43
B29	29.3		29.4		30.1		29.6	0.44	1.49	0.38
B39	29.6		29.3		28.8		29.2	0.40	1.37	0.33
B49	32.9		33.7		33.2		33.3	0.40	1.20	0.33
B59	34.4		33.8		33.0		33.7	0.70	2.08	0.99

Pooled variance (s^2overall) **0.28**
Minimum detectable difference (б) **0.72**
Mean Minimum dose (DMin) **28.3 B21**
Mean Maximum dose (DMax) **35.4 A49**

Expected value of Rmin (Dref/Dmin) **1.142** *
Expected value of Rmax (Dref/Dmax) **0.912** *

DRef Ratio :	
1/Rmin	**0.876**
1/Rmax	**1.096**

Remarks: * incorporates an estimation of the uncertainty
Customer Spec Min: **25.0** Max: **50.0**
D reference Min: **28.6** Max: **45.6**
Dose Uniformity Ratio (DUR): **1.25**

Performance Qualification STERIS Kuala Ketil

Validation Ref : 0.0969 Rev 01

Product: HALYARD*PUREZERO* HG3 Sterile White Nitrile Gloves - Pairs **HIGH DENSITY**

Analysis for the Calculation of Release Specification Incorporating Uncertainties

Position	PQ1		PQ2		PQ3		Mean	Stdev	CV	Sum of Squared Difference
A11	31.5		32.3		32.8		32.2	0.66	2.05	0.86
A21	31.5		31.7		31.6		31.6	0.10	0.32	0.02
A31	29.9		29.4		30.2		29.8	0.40	1.34	0.33
A41	31.7		31.4		31.4		31.5	0.17	0.54	0.06
A51	34.5		32.9		32.9		33.4	0.92	2.75	1.71
A15	34.0		33.9		34.2		34.0	0.15	0.44	0.05
A25	32.3		31.9		31.7		32.0	0.31	0.97	0.19
A35	34.1		33.7		34.0		33.9	0.21	0.62	0.09
A45	34.8		32.9		33.7		33.8	0.95	2.81	1.82
A55	31.8		33.8		34.5		33.4	1.40	4.19	3.93
A19	33.6		34.1		33.7		33.8	0.26	0.77	0.14
A29	33.9		35.0		33.9		34.3	0.64	1.87	0.81
A39	**35.2**	√-MAX	34.4		34.8		34.8	0.40	1.15	0.32
A49	34.5		34.2		34.3		34.3	0.15	0.44	0.05
A59	34.9		**35.4**	√-MAX	**35.1**	√-MAX	**35.1**	0.25	0.71	0.13
B11	27.8		28.6		28.6		28.3	0.46	1.63	0.43
B21	27.6		28.0		28.1		27.9	0.26	0.93	0.14
B31	**26.8**	√-MIN	**26.7**	√-MIN	**26.8**	√-MIN	**26.8**	0.06	0.22	0.01
B41	31.3		30.6		31.3		31.1	0.40	1.29	0.33
B51	32.9		32.2		32.0		32.4	0.47	1.45	0.45
B15	30.0		29.7		30.6		30.1	0.46	1.53	0.42
B25	28.3		28.7		28.7		28.6	0.23	0.80	0.11
B35	27.4		27.6		27.9		27.6	0.25	0.91	0.13
B45	29.5		29.0		30.6		29.7	0.82	2.76	1.34
B55	31.2		28.5		31.7		30.5	1.72	5.64	5.93
B19	29.9		31.9		30.4		30.7	1.04	3.39	2.17
B29	28.3		30.8		28.9		29.3	1.31	4.47	3.41
B39	28.6		31.4		28.4		29.5	1.68	5.69	5.63
B49	33.3		32.7		32.5		32.8	0.42	1.28	0.35
B59	33.6		34.3		33.4		33.8	0.47	1.39	0.45

Pooled variance (s^2overall) **0.53**

Minimum detectable difference (б) **0.99**

Mean Minimum dose (DMin) **26.8 B31**

Mean Maximum dose (DMax) **35.1 A59**

Expected value of Rmin (Dref/Dmin) **1.202** *

Expected value of Rmax (Dref/Dmax) **0.917** *

Dref Ratio :	
1/Rmin	**0.832**
1/Rmax	**1.091**

Remarks: * incorporates an estimation of the uncertainty

Customer Spec Min: **25.0** Max: **50.0**

D reference Min: **30.1** Max: **45.9**

Dose Uniformity Ratio (DUR): **1.31**

Performance Qualification STERIS Kuala Ketil

Product Detail **Validation Ref** : 0.0969 Rev 01

Customer Name : **Safeskin Medical & Scientific (Thailand) Ltd**

Issue Date : **29.04.2021**

Product Description : **HALYARD*PUREZERO* HG3 Sterile White Nitrile Gloves - Pairs**

Plant Batch No : **S12562828-1-1 (HD - high density, size 10.0)**
S12562828-1-2 (LD - low density, size 6.0)

				Weight (LD)	:	**6.45** kg
Current Cobalt Loading	:	**4,591,067**	Curries	Weight (HD)	:	**8.20** kg
Standard Plant Dwell Time	:	**146**	Sec	Length	:	**345** mm
Actual Dwell Time	:	**146**	Sec	Width	:	**285** mm
Number of Xs	:	**38.0**		Height	:	**390** mm
Value of X	:	**3.83**		Density (LD)	:	**0.168** g/cm^3
Type of package	:	**Corrugated carton**		Density (HD)	:	**0.214** g/cm^3
Dose Range Specification (kGy)	:	**25**	Min	Tote Fit	:	**15** ctn/tote
		50	Max			

Low Density Dwell Time Control

Minimum dwell time to achieve	Minimum dose		Maximum dose	
First tote	131	√	202	√
Second tote	127		202	√
Third tote	129		208	

High Density Dwell Time Control

Minimum dwell time to achieve	Minimum dose		Maximum dose	
First tote	136		207	
Second tote	137	√	206	√
Third tote	136		208	

Reference Procedures: : QWI 24 Rev 12 Dose Mapping
Method : ISO/ASTM 52303: 2015 (E)
Reference Standard : EN ISO 11137-1: 2015

Performance Qualification STERIS Kuala Ketil

Validation Ref : 0.0969 Rev 01

Customer Name : **Safeskin Medical & Scientific (Thailand) Ltd**

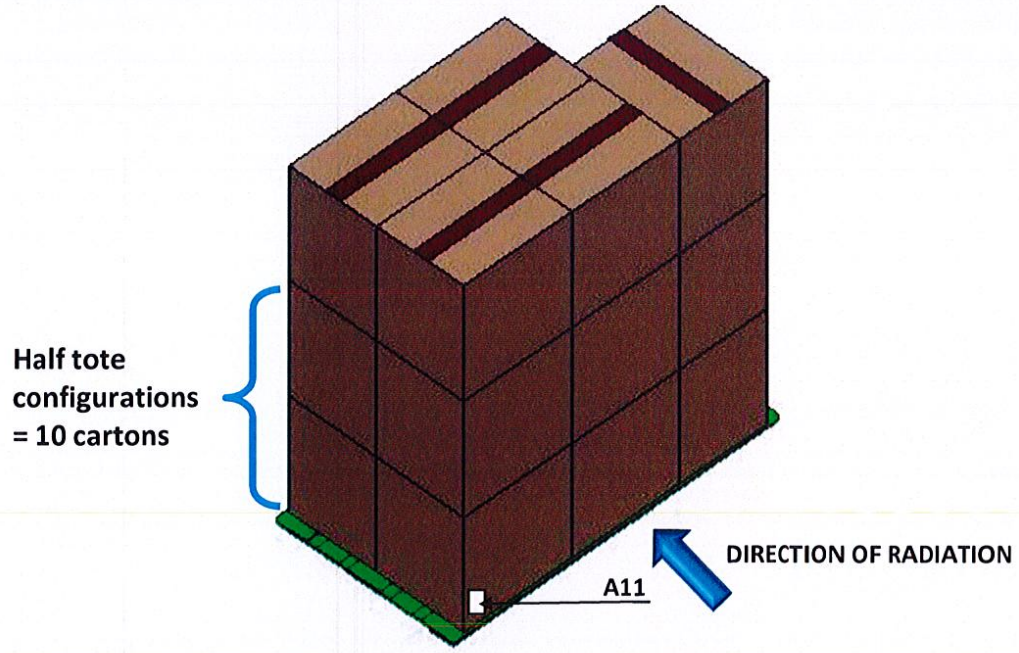
Type of Package : **Corrugated carton**

Product Description : **HALYARD*PUREZERO* HG3 Sterile White Nitrile Gloves - Pairs**

Issue Date : **29.04.2021**

This performance qualification relates only to the above product loaded in the configuration outlined below.

Carton dimension : **345 mm (L) x 285 mm (W) x 390 mm (H) (15 ctns/tote)**



Remark : A11 is the routine process monitoring

Authorised By:	Signature	Date
Radiation Technology Management or designee		06 May 2021

Customer Name : **Safeskin Medical & Scientific (Thailand) Ltd**

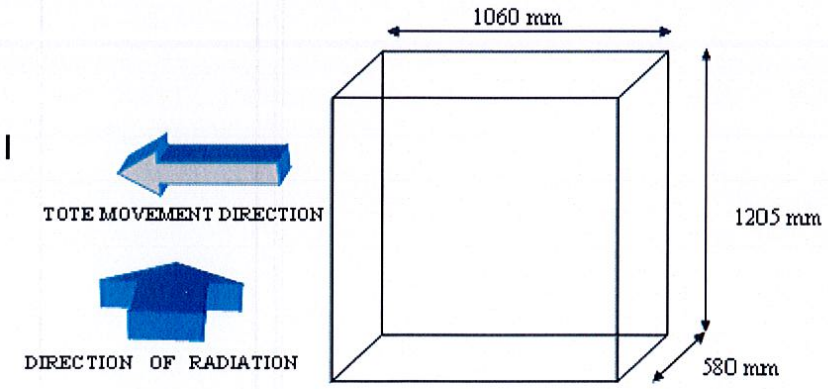
Type of Package : **Corrugated carton**

Product Description : **HALYARD*PUREZERO* HG3 Sterile White Nitrile Gloves - Pairs**

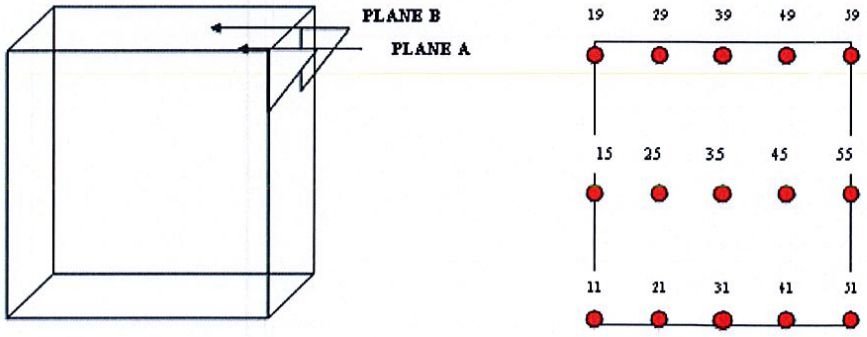
Issue Date : **29.04.2021**

This performance qualification relates only to the above product loaded in the configuration outlined below.

TOTE CHARACTERISTICS AND ORIENTATION



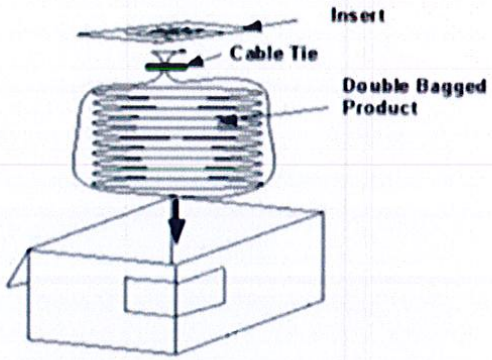
LOCATION OF DOSIMETERS



As per our Operational Qualification Report, from latest source loading replenishment, the dose distribution in the front plane (plane A) is the same as in the back plane (plane C).

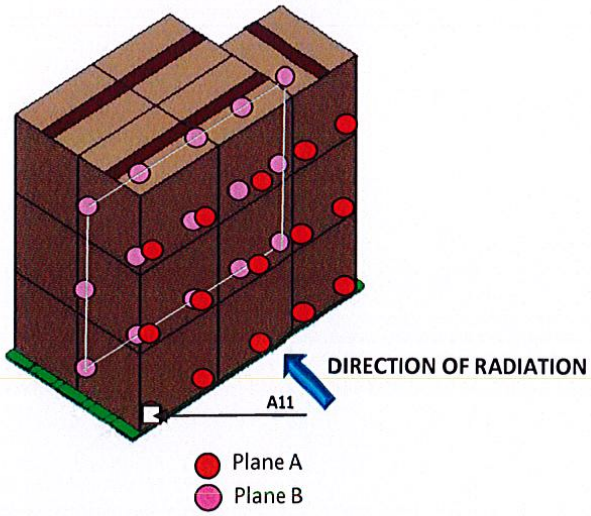
Customer Name: **Safeskin Medical & Scientific (Thailand) Ltd**

Orientation of product within the package



Dosimeter position





Remark : A11 is the routine process monitoring

Authorised By:	Signature	Date
Radiation Technology Management or designee		06 May 2021

SAFETY DATA SHEET



1. IDENTIFICATION

Product Name: ***HALYARD* PUREZERO* HG3 Nitrile Gloves***

Product Number: Various

Date Prepared:	6/24/2021
Manufacturer:	O&M Halyard, Inc. 1 Edison Dr. Alpharetta, Georgia 30005
Telephone: MEDICAL EMERGENCY: TRANSPORTATION EMERGENCY: INFORMATION:	 Call 911 or your local Emergency Room or Poison Control Services Chemtrec 1-800-424-9300 or 1-703-527-3887 (collect calls accepted) 1-844-HALYARD

Product Use: Protective gloves.

2. HAZARDS IDENTIFICATION

GHS Classification: Not Hazardous. These products are manufactured articles or articles as this term is defined in the OSHA Hazard Communication Standard (20CFR 1910.1200), EU REACH, Canadian WHMIS, Australia WHS and the GHS. These products are out of the scope of the GHS and no SDS or labeling are required.

GHS Labeling: No GHS labeling required.

3. COMPOSITION/INFORMATION ON INGREDIENTS

Chemical Name	CAS-No./ EINECS-No	*%*	*GHS Classification:*
Non-Hazardous Ingredients	Mixture	100	Not Hazardous

4. FIRST AID MEASURES

Inhalation: No first aid needed.

Skin contact: No first aid needed. These products are intended to be used in contact with the skin.

Eye contact: No first aid needed.

Ingestion: Not a relevant route of exposure.

Most Important symptoms and effects, both acute and delayed: No adverse effects.

Indication of any immediate medical attention and special treatment needed: None required.

5. FIRE-FIGHTING MEASURES

Suitable (and unsuitable) Extinguishing Media: Use media suitable for surrounding materials.

Specific Hazards Arising from the Chemical: These products can melt in a fire. Molten material may flow or drip and spread the fire.

Hazardous Combustion Products: Combustion may produce toxic and irritating gases and vapors.

Special Protective Equipment and Precautions for Fire-fighters: None required.

6. ACCIDENTAL RELEASE MEASURES

Personal Precautions, Protective Equipment, and Emergency Procedures: None required.

Environmental Precautions: None required.

Methods and Materials for Containment and Cleaning Up: No special precautions required. Pick up and dispose of in an appropriate container.

7. HANDLING AND STORAGE

Precautions for Safe Handling: No special handling is required.

Conditions for Safe Storage, including any Incompatibilities: Store away from heat and flames.

8. EXPOSURE CONTROLS / PERSONAL PROTECTION

Exposure limit(s)

Chemical Name	Exposure limit(s)
Non-Hazardous Ingredients	None established

Appropriate Engineering Controls: None required.

Individual Protection Measures, such as Personal Protective Equipment:

Respiratory Protection: None required.

Protective Gloves: None required.

Eye Protection: None required.

9. PHYSICAL AND CHEMICAL PROPERTIES

Appearance and Odor	Solid Gloves, blue or white
Odor Threshold:	Not applicable
Boiling point/range	Not applicable
Melting point/range	Not determined
Relative Density (water=1)	Not determined
Vapor pressure	Not applicable
Vapor density (air=1)	Not applicable
Solubility	Insoluble
pH	Not applicable
Partition coefficient (n-octanol/water):	Not applicable
Evaporation Rate (Butyl acetate=1)	Not applicable
Viscosity:	Not applicable
Volatile Organic Carbon Compounds (VOC) (g/L)	Not applicable
Flashpoint:	None.
Flammable Limits in Air % by Volume:	LEL (Lower):Not applicable UEL (Upper): Not applicable
Autoignition temperature:	Not determined.
Decomposition temperature:	Not determined
Flammability (solid, gas):	Not a flammable solid

10. STABILITY AND REACTIVITY

Reactivity: Not reactive.

Chemical Stability: Stable at ambient temperature and pressure.

Possibility of Hazardous Reactions: None known.

Conditions to Avoid: Avoid heat and flames.

Incompatible Materials: None known.

Hazardous Decomposition Products: Thermal decomposition may produce carbon and nitrogen oxides and unidentified hydrocarbon fragments.

11. TOXICOLOGICAL INFORMATION

Symptoms/Effects of Overexposure: These gloves are manufactured articles. Their use will not result in exposure to any hazardous materials.

Inhalation: Not applicable.

Skin contact: None. Intended for skin contact. These products contain no natural latex.

Eye contact: None expected.

Ingestion: Not applicable.

Chronic toxicity: None expected.

Carcinogenicity Data: None of the components of this product is listed as a carcinogen by IARC, NTP, US OSHA or the EU CLP Annex VI.

Reproductive Toxicity: None known.

Numerical Measures of Toxicity:
Not acutely toxic.

12. ECOLOGICAL INFORMATION

Ecotoxicity: These gloves are manufactured articles. Their use will not result in adverse environmental effects.

Persistence and Degradability: Not degradable.

Bioaccumulative Potential: Not bioaccumulative.

Mobility in Soil: Not mobile.

Other Adverse Effects: None currently known.

13. DISPOSAL CONSIDERATIONS

Dispose in accordance with all state, local and federal regulations. If unused, the product may be disposed of as a non-hazardous material.

14. TRANSPORT INFORMATION

US DOT Shipping Description: Not regulated

IATA Shipping Description (Air): Not regulated

IMDG Shipping Description (Vessel): Not regulated

15. REGULATORY INFORMATION

US Regulations

EPA Toxic Substances Control Act (TSCA): These gloves are articles and exempt from the TSCA regulations.

SARA 302 Listed Chemicals: None.

SARA 311/312 Hazard Categories: Not applicable to articles

SARA 313 This Product Contains the Following Chemicals Subject to Annual Release Reporting Requirements Under the SARA Section 313 (40 CFR 372): Not applicable to articles.

16. OTHER INFORMATION

Date Prepared: 6/24/2021

HMIS Rating: Health 0 Fire 1 Physical Hazard 0

The information contained herein is true and correct to the best of O&M Halyard, Inc's knowledge. However, no warranty, expressed or implied, is made. Nothing herein should be interpreted as a recommendation to infringe existing patents or violate any laws or regulations. Final determination of the suitability of the material is the sole responsibility of the user.

July 11, 2023

RE: HALYARD* PUREZERO* **HG3 and HG5 Cleanroom Nitrile Gloves**

O&M Halyard, Inc. confirms that our Halyard **PURE**ZERO* HG3 and Halyard **PURE**ZERO HG5 Nitrile Cleanroom Glove families are fully compliant with the related requirements of the European Union Regulation (EC) 1907/2006 concerning Registration, Evaluation, Authorization, and Restriction of Chemicals (REACH).

O&M Halyard, Inc. hereby confirms that our Halyard **PURE**ZERO* HG3 and Halyard **PURE**ZERO HG5 Nitrile Cleanroom Gloves either do not contain the presence of greater than 0.1% of substances identified on the SVHC Candidate list according to Article 33 of the REACH Regulation and/or do not contain any of the listed substances.

Thank you for your interest in Halyard products. If you have any questions or need additional information, please do not hesitate to contact us at PIQ@hyh.com or call us directly at (844) 425-9273.

Sincerely,

Caitlin Senter
Director
Global Regulatory Affairs
O&M Halyard, Inc.

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bsi.

By Royal Charter

Certificate of Registration

QUALITY MANAGEMENT SYSTEM - ISO 9001:2015

This is to certify that:

Safeskin Medical & Scientific
(Thailand) Ltd.
200 Moo 8, Kanchanavanich Road,
Tambol Prik, Amphur Sadao,
Songkhla
90120
Thailand

Holds Certificate Number: **FM 697757**

and operates a Quality Management System which complies with the requirements of ISO 9001:2015 for the following scope:

Design and Development, Production and Distribution of Industrial Gloves-Sterile and Non-Sterile Examination Gloves.

For and on behalf of BSI:

Michael Lam - Managing Director Assurance, APAC

Original Registration Date: 2018-08-12
Effective Date: 2022-09-01
Latest Revision Date: 2022-09-01
Expiry Date: 2025-08-31

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This certificate was issued electronically and remains the property of BSI and is bound by the conditions of contract.
An electronic certificate can be authenticated online.
Printed copies can be validated at www.bsi-global.com/ClientDirectory or telephone +66(2) 2944889-92.
Further clarifications regarding the scope of this certificate and the applicability of ISO 9001:2015 requirements may be obtained by consulting the organization.
This certificate is valid only if provided original copies are in complete set.

ISO 9001:2015 (continued)

Certificate No: **FM 697757**

Location	Registered Activities
Safeskin Medical & Scientific (Thailand) Ltd. 200 Moo 8, Kanchanavanich Road, Tambol Prik, Amphur Sadao, Songkhla 90120 Thailand	Design and Development, Production and Distribution of Industrial Gloves-Sterile and Non-Sterile Examination Gloves. Including the following support function: Owens & Minor Halyard Incorporation. 5405 Windward Parkway, Alpharetta, GA, USA : Design and Development of Industrial Gloves-Sterile and Non-Sterile Examination Gloves.

Original Registration Date: 2018-08-12
Latest Revision Date: 2022-09-01

Effective Date: 2022-09-01
Expiry Date: 2025-08-31

This certificate was issued electronically and remains the property of BSI and is bound by the conditions of contract.
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Further clarifications regarding the scope of this certificate and the applicability of ISO 9001:2015 requirements may be obtained by consulting the organization. This certificate is valid only if provided original copies are in complete set.

Information and Contact: BSI, Kitemark Court, Davy Avenue, Knowlhill, Milton Keynes MK5 8PP. Tel: + 44 345 080 9000
BSI Assurance UK Limited, registered in England under number 7805321 at 389 Chiswick High Road, London W4 4AL, UK.

ISO 13485:2016

bsi.

By Royal Charter

Certificate of Registration

QUALITY MANAGEMENT SYSTEM - ISO 13485:2016 & EN ISO 13485:2016

This is to certify that:

O & M Halyard, Inc.
9120 Lockwood Blvd
Mechanicsville
Virginia
23116
USA

Holds Certificate Number: **MD 735497**

and operates a Quality Management System which complies with the requirements of ISO 13485:2016 & EN ISO 13485:2016 for the following scope:

Design, development, manufacture and distribution of sterile and non-sterile medical examination gloves.

For and on behalf of BSI:

Graeme Tunbridge, Senior Vice President Medical Devices

Original Registration Date: 2020-11-25
Latest Revision Date: 2023-01-05

Effective Date: 2023-01-09
Expiry Date: 2026-01-08

Page: 1 of 2

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ISO 13485:2016 (continued)

Certificate No: **MD 735497**

Location	Registered Activities
O & M Halyard, Inc. 9120 Lockwood Blvd Mechanicsville Virginia 23116 USA	Headquarter management activities.
O & M Halyard, Inc. 1 Edison Drive Alpharetta Georgia 30005 USA	The design and development of sterile and non-sterile medical examination gloves.
Safeskin Medical & Scientific (Thailand) Ltd. 200 Moo 8, Kanchanavanich Road, Tambol Prik, Amphur Sadao, Songkhla 90120 Thailand	Design, development, manufacture and distribution of sterile and non-sterile medical examination gloves.

Original Registration Date: 2020-11-25
Latest Revision Date: 2023-01-05

Effective Date: 2023-01-09
Expiry Date: 2026-01-08

This certificate was issued electronically and remains the property of BSI and is bound by the conditions of contract.
An electronic certificate can be authenticated online.

ISO 14001:2015

bsi.

By Royal Charter

Certificate of Registration

ENVIRONMENTAL MANAGEMENT SYSTEM - ISO 14001:2015

This is to certify that:

Safeskin Medical & Scientific
(Thailand) Ltd.
200 Moo 8, Kanchanavanich Road,
Tambol Prik, Amphur Sadao,
Songkhla
90120
Thailand

Holds Certificate Number: **EMS 772160**

and operates an Environmental Management System which complies with the requirements of ISO 14001:2015 for the following scope:

The Manufacturing of Medical and Scientific Gloves.

For and on behalf of BSI:

Michael Lam - Managing Director Assurance, APAC

Original Registration Date: 2022-10-27
Effective Date: 2022-10-27
Latest Revision Date: 2022-10-27
Expiry Date: 2025-10-26

Page: 1 of 1

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Further clarifications regarding the scope of this certificate and the applicability of ISO 14001:2015 requirements may be obtained by consulting the organization.
This certificate is valid only if provided original copies are in complete set.

Information and Contact: BSI, Kitemark Court, Davy Avenue, Knowlhill, Milton Keynes MK5 8PP. Tel: + 44 345 080 9000
BSI Assurance UK Limited, registered in England under number 7805321 at 389 Chiswick High Road, London W4 4AL, UK.
A Member of the BSI Group of Companies.

May 4, 2023

RE: **HALYARD* PURE**ZERO**HG3 and HG5 Nitrile Gloves**

Dear Valued Customer,

This letter is in response to your recent accelerator-free inquiry for the **HALYARD* PURE**ZERO**HG3 and HG5 Nitrile Gloves.**

The following gloves are accelerator-free and are not formulated with the following vulcanizing chemicals and accelerators: Sulfur, Thiurams, Thiaxoles, Guanidines, Carbamates.†

HALYARD* PUREZERO**HG3 White Nitrile Gloves**
HALYARD* PUREZERO**HG3 Light Blue Nitrile Gloves**
HALYARD* PUREZERO**HG3 White SGX* Nitrile Gloves**
HALYARD* PUREZERO**HG3 White Sterile Nitrile Gloves**
HALYARD* PUREZERO**HG3 Light Blue Sterile Nitrile Gloves**

HALYARD* PUREZERO**HG5 White Nitrile Gloves**
HALYARD* PUREZERO**HG5 Blue SGX* Nitrile Gloves**

Thank you for your interest in Halyard products. If you have any questions or need additional information, please do not hesitate to contact us at PIQ@hyh.com or call us directly at (844) 425-9273.

Sincerely,

Shannon Smith

Shannon Smith
Associate Product Manager
O&M Halyard, Inc.

† The chemicals and accelerators shown is not a comprehensive list.

*Registered Trademark or Trademark of O&M Halyard or its affiliates.

COPY-04632

PUREZERO*

HG3 Nitrile Cleanroom Gloves

ADDED PROTECTION FROM THE RISK OF CHEMICAL EXPOSURE

It's critical to protect staff from exposure to potentially hazardous chemicals and chemotherapy drugs. In addition to providing a barrier to chemical splash, microorganisms and viruses, **PURE**ZERO* HG3 Cleanroom Gloves are **now tested against 29 chemicals, 14 chemotherapy drugs and Fentanyl.**

CHEMOTHERAPY DRUG RESISTANCE GUIDE[4]

Chemotherapy Agent (Concentration in mg/ml)	HALYARD* **PURE**ZERO* HG3 White Sterile Nitrile Gloves	
	Minimum Breakthrough Detection Time (minutes)	Average Steady State Perm. Rate (µg/cm²/min)
Carmustine (BCNU) (3.3)	158.5	0.06
Cisplatin (1.0)	>240	N/A
Cyclophasphamide (20.0)	>240	N/A
Dacarbazine (10.0)	>240	N/A
Doxorubicin HCL (2.0)	>240	N/A
Etoposide (20.0)	>240	N/A
Fluorouracil (50.0)	>240	N/A
Ifosfamide (50.0)	>240	N/A
Methotrexate (25.0)	>240	N/A
Mitomycin C (0.5)	>240	N/A
Mitoxantrone (2.0)	>240	N/A
Paclitaxel (6.0)	>240	N/A
Thiotepa (10.0)	>240	N/A
Vincristine (1.0)	>240	N/A
Additional Tests		
Fentanyl Citrate Injection, 100 mcg/2mL	>240	N/A

Use the rating system below to determine the chemotherapy compatibility for exposure:

<10	Not recommended for use - breakthrough can occur in less than 10 minutes.
11 - 239	Use with caution - breakthrough can occur between 11 and 239 minutes.
>240	Recommended for protection - no breakthrough up to 240 minutes.
N/A	The chemotherapy drug did not reach the minimum permeation rate (0.01 µg/cm²/min) as defined within ASTM D6978.

CAUTION: It's the user's responsibility to determine the applicability of these gloves for their intended use with chemotherapy drugs.

DEFINITION OF TERMS

Breakthrough time: The time required for the test chemical to be detected on the inside of the glove. Essentially, this is the amount of time that the glove can resist a chemical when the glove is fully immersed in the chemical.

Permeation: The process where chemicals, such as liquids, gases and vapors can pass through a glove film (or other PPE interfaces) without penetrating directly through a pinhole, tear or other visible opening.

Permeation rate: The flowrate of the chemical after the chemical breaks through the inside of the glove. It is measured in amount per surface area of the glove per time (µg/cm²/min).

Contact your Sales Representative for additional chemical and chemotherapy drug testing information.

PUREZERO*

HG3 Nitrile Cleanroom Gloves

CHEMICAL RESISTANCE GUIDE[5]

Chemical (Concentration %)	HALYARD* **PURE**ZERO* HG3 White Sterile Nitrile Gloves	
	Minimum Breakthrough Time (minutes)	Permeation Rate (µg/cm²/min)
1-Butanol (99)	85.6	4.0
Acetic Acid (30)	>480	<1.0
Acrylamide (40)	>480	<1.0
Benzalkonium Chloride (1)	>480	<1.0
Chloroform (70)	0	N/A
Citric Acid (30)	>480	<1.0
Citric Acid (70)	>480	<1.0
Cyclohexane (99)	147.8	2.7
Didecyldimethylammonium Chloride (0.04)	>480	<1.0
Dimethylformamide (99)	0	N/A
Dimethyl Sulfoxide (99)	10.7	N/A
Ethanol (70)	19.1	1.2E+01
Ethanol (99)	16.3	2.5E+01
Ethidium Bromide (1)	>480	<1.0
Formaldehyde (37)	>480	<1.0
Glycolic Acid (10)	>480	<1.0
Glutaraldehyde (50)	>480	<1.0
Heptane (99)	>480	<1.0
Hydrazine Monohydrate (55)	>480	<1.0
Hydrochloric Acid (30)	>480	<1.0
Hydrogen Peroxide (30)	310.0	1.0
Isopropyl Alcohol (70)	189.5	1.0
Isopropyl Alcohol (99)	105.3	3.5
Methanol (99)	7.7	1.9E+01
Nitric Acid (65)	12.0	1.7E+02
Peracetic Acid (5)	>480	<1.0
Phosphoric Acid (70)	>480	<1.0
Sodium Hydroxide (50)	>480	<1.0
Sodium Hypochlorite (10-13)	>480	<1.0
Spor-Klenz (N/A)	>480	<1.0
Sulfuric Acid (50)	>480	<1.0
Sulfuric Acid (99)	11.0	N/A

Use the rating system below to determine the chemical compatibility for exposure:

<10	Not recommended for use - breakthrough can occur in less than 10 minutes.
11 - 479	Use with caution - breakthrough can occur between 11 and 479 minutes.
>480	Permeation breakthrough is excellent. Permeation does not occur during the test (8 hours).
N/A	The chemical did not reach the minimum permeation rate (1 µg/cm²/min) as defined within EN 16523-1. There is a possibility for trace amounts of the chemical to permeate through the glove.
–	The permeation rate was beyond the range of the detection instruments. The permeation of the chemical through the glove film may be too high for the detector to reach a steady-state reading.

CAUTION: It's the user's responsibility to determine the applicability of these gloves for their intended use.

November 13th, 2023

RE: **ESD Testing for HALYARD* PURE**ZERO*** Nitrile Gloves**

Dear Valued Customer,

This letter is in response to your recent inquiry regarding ESD testing of **HALYARD* PURE**ZERO*** Nitrile Gloves.**

The following **HALYARD* PURE**ZERO*** Nitrile Glove** families have been tested against ANSI/ESD STM15.1:2019 "In-Use Resistance Testing of Gloves and Finger Cots":

HALYARD* PUREZERO* HG3 Cleanroom Nitrile Gloves

Glove Description	Code
HALYARD* **PURE**ZERO* HG3 White Nitrile Gloves	CLN3031XS
HALYARD* **PURE**ZERO* HG3 White Nitrile Gloves	CLN3031SM
HALYARD* **PURE**ZERO* HG3 White Nitrile Gloves	CLN3031MD
HALYARD* **PURE**ZERO* HG3 White Nitrile Gloves	CLN3031LG
HALYARD* **PURE**ZERO* HG3 White Nitrile Gloves	CLN3031XL
HALYARD* **PURE**ZERO* HG3 Blue Nitrile Gloves	CLN9031XS
HALYARD* **PURE**ZERO* HG3 Blue Nitrile Gloves	CLN9031SM
HALYARD* **PURE**ZERO* HG3 Blue Nitrile Gloves	CLN9031MD
HALYARD* **PURE**ZERO* HG3 Blue Nitrile Gloves	CLN9031LG
HALYARD* **PURE**ZERO* HG3 Blue Nitrile Gloves	CLN9031XL
HALYARD* **PURE**ZERO* HG3 White SGX* Nitrile Gloves	CLN3231XS
HALYARD* **PURE**ZERO* HG3 White SGX* Nitrile Gloves	CLN3231SM
HALYARD* **PURE**ZERO* HG3 White SGX* Nitrile Gloves	CLN3231MD
HALYARD* **PURE**ZERO* HG3 White SGX* Nitrile Gloves	CLN3231LG
HALYARD* **PURE**ZERO* HG3 White SGX* Nitrile Gloves	CLN3231XL
HALYARD* **PURE**ZERO* HG3 White Sterile Nitrile Gloves	CLN323260
HALYARD* **PURE**ZERO* HG3 White Sterile Nitrile Gloves	CLN323265
HALYARD* **PURE**ZERO* HG3 White Sterile Nitrile Gloves	CLN323270
HALYARD* **PURE**ZERO* HG3 White Sterile Nitrile Gloves	CLN323275
HALYARD* **PURE**ZERO* HG3 White Sterile Nitrile Gloves	CLN323280
HALYARD* **PURE**ZERO* HG3 White Sterile Nitrile Gloves	CLN323285
HALYARD* **PURE**ZERO* HG3 White Sterile Nitrile Gloves	CLN323290
HALYARD* **PURE**ZERO* HG3 White Sterile Nitrile Gloves	CLN323210

HALYARD* **PURE**ZERO* HG3 Blue Sterile Nitrile Gloves	CLN923260
HALYARD* **PURE**ZERO* HG3 Blue Sterile Nitrile Gloves	CLN923265
HALYARD* **PURE**ZERO* HG3 Blue Sterile Nitrile Gloves	CLN923270
HALYARD* **PURE**ZERO* HG3 Blue Sterile Nitrile Gloves	CLN923275
HALYARD* **PURE**ZERO* HG3 Blue Sterile Nitrile Gloves	CLN923280
HALYARD* **PURE**ZERO* HG3 Blue Sterile Nitrile Gloves	CLN923285
HALYARD* **PURE**ZERO* HG3 Blue Sterile Nitrile Gloves	CLN923290
HALYARD* PUREZERO* HG3 Blue Sterile Nitrile Gloves	CLN923210

HALYARD* PUREZERO* HG5 Cleanroom Nitrile Gloves

Glove Description	Code
HALYARD* **PURE**ZERO* HG5 White Nitrile Gloves	CLN30512XS
HALYARD* **PURE**ZERO* HG5 White Nitrile Gloves	CLN30512SM
HALYARD* **PURE**ZERO* HG5 White Nitrile Gloves	CLN30512MD
HALYARD* **PURE**ZERO* HG5 White Nitrile Gloves	CLN30512LG
HALYARD* **PURE**ZERO* HG5 White Nitrile Gloves	CLN30512XL
HALYARD* **PURE**ZERO* HG5 Blue SGX* Nitrile Gloves	CLN92512XS
HALYARD* **PURE**ZERO* HG5 Blue SGX* Nitrile Gloves	CLN92512SM
HALYARD* **PURE**ZERO* HG5 Blue SGX* Nitrile Gloves	CLN92512MD
HALYARD* **PURE**ZERO* HG5 Blue SGX* Nitrile Gloves	CLN92512LG
HALYARD* **PURE**ZERO* HG5 Blue SGX* Nitrile Gloves	CLN92512XL

HALYARD* PUREZERO* Nitrile Exam Gloves

Glove Description	Code
HALYARD* **PURE**ZERO* MARIN* Nitrile Exam Gloves	LFS111XS
HALYARD* **PURE**ZERO* MARIN* Nitrile Exam Gloves	LFS111SM
HALYARD* **PURE**ZERO* MARIN* Nitrile Exam Gloves	LFS111MD
HALYARD* **PURE**ZERO* MARIN* Nitrile Exam Gloves	LFS111LG
HALYARD* **PURE**ZERO* MARIN* Nitrile Exam Gloves	LFS111XL
HALYARD* **PURE**ZERO* MARIN-XTRA* Nitrile Exam Gloves	LFS121XS
HALYARD* **PURE**ZERO* MARIN-XTRA* Nitrile Exam Gloves	LFS121SM
HALYARD* **PURE**ZERO* MARIN-XTRA* Nitrile Exam Gloves	LFS121MD
HALYARD* **PURE**ZERO* MARIN-XTRA* Nitrile Exam Gloves	LFS121LG
HALYARD* **PURE**ZERO* MARIN-XTRA* Nitrile Exam Gloves	LFS121XL
HALYARD* **PURE**ZERO* LIMON* Nitrile Exam Gloves	LFS311XS
HALYARD* **PURE**ZERO* LIMON* Nitrile Exam Gloves	LFS311SM
HALYARD* **PURE**ZERO* LIMON* Nitrile Exam Gloves	LFS311MD

HALYARD* **PURE**ZERO* LIMON* Nitrile Exam Gloves	LFS311LG
HALYARD* **PURE**ZERO* LIMON* Nitrile Exam Gloves	LFS311XL
HALYARD* **PURE**ZERO* LIMON-XTRA* Nitrile Exam Gloves	LFS321XS
HALYARD* **PURE**ZERO* LIMON-XTRA* Nitrile Exam Gloves	LFS321SM
HALYARD* **PURE**ZERO* LIMON-XTRA* Nitrile Exam Gloves	LFS321MD
HALYARD* **PURE**ZERO* LIMON-XTRA* Nitrile Exam Gloves	LFS321LG
HALYARD* **PURE**ZERO* LIMON-XTRA* Nitrile Exam Gloves	LFS321XL
HALYARD* **PURE**ZERO* ULTRA VIOLET* Nitrile Exam Gloves	LFS511XS
HALYARD* **PURE**ZERO* ULTRA VIOLET* Nitrile Exam Gloves	LFS511SM
HALYARD* **PURE**ZERO* ULTRA VIOLET* Nitrile Exam Gloves	LFS511MD
HALYARD* **PURE**ZERO* ULTRA VIOLET* Nitrile Exam Gloves	LFS511LG
HALYARD* **PURE**ZERO* ULTRA VIOLET* Nitrile Exam Gloves	LFS511XL

HALYARD* PUREZERO* **Nitrile Gloves**

Glove Description	Code
HALYARD* **PURE**ZERO* MARIN* Nitrile Gloves	LFS1117XS
HALYARD* **PURE**ZERO* MARIN* Nitrile Gloves	LFS1117SM
HALYARD* **PURE**ZERO* MARIN* Nitrile Gloves	LFS1117MD
HALYARD* **PURE**ZERO* MARIN* Nitrile Gloves	LFS1117LG
HALYARD* **PURE**ZERO* MARIN* Nitrile Gloves	LFS1117XL
HALYARD* **PURE**ZERO* LIMON* Nitrile Gloves	LFS3117XS
HALYARD* **PURE**ZERO* LIMON* Nitrile Gloves	LFS3117SM
HALYARD* **PURE**ZERO* LIMON* Nitrile Gloves	LFS3117MD
HALYARD* **PURE**ZERO* LIMON* Nitrile Gloves	LFS3117LG
HALYARD* **PURE**ZERO* LIMON* Nitrile Gloves	LFS3117XL
HALYARD* **PURE**ZERO* ULTRA VIOLET* Nitrile Gloves	LFS5117XS
HALYARD* **PURE**ZERO* ULTRA VIOLET* Nitrile Gloves	LFS5117SM
HALYARD* **PURE**ZERO* ULTRA VIOLET* Nitrile Gloves	LFS5117MD
HALYARD* **PURE**ZERO* ULTRA VIOLET* Nitrile Gloves	LFS5117LG
HALYARD* **PURE**ZERO* ULTRA VIOLET* Nitrile Gloves	LFS5117XL

Based on the ANSI/ESD STM15.1:2019 test results from an external laboratory at conditions of 12% RH and 23°C, all **HALYARD* PURE**ZERO*** Nitrile Glove** product families listed above are within the dissipative classification range (Resistance = 10^4 to 10^{11} ohms) as defined by ANSI/ESD standards.

Therefore, **HALYARD* PURE**ZERO*** Nitrile Gloves** are considered to be static dissipative in-use. Please note that this is based on a standardized test method in a laboratory environment, and the resistance test results and classification do not fully characterize the glove for all electrostatic properties. Other electrostatic properties, such as charge accumulation, and specific process conditions may need to be assessed based on your particular application prior to use.

Thank you for your interest in Halyard products. If you have any questions or need additional information, please do not hesitate to contact us at PIQ@hyh.com or call us directly at (844) 425-9273.

Sincerely,

Caitlin Senter
Director
Global Regulatory Affairs
O&M Halyard, Inc.

Ryan Solan
Manager
Global Research and Development
O&M Halyard, Inc.

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COPY-04825

RE: Compliance statement – HALYARD* **PURE**ZERO* Nitrile Gloves

Please be advised that to the best of O&M Halyard knowledge, the **PURE**ZERO* HG3 Nitrile Gloves and HALYARD* **PURE**ZERO* HG5 Nitrile Gloves are produced entirely from materials of synthetic origin and therefore are free from human or any other animal derived materials including bovine products (Bovine Spongiform Encephalopathy (BSE) and Transmissible Spongiform Encephalopathy (TSE)).

- *Under various environmental protection and product safety regulations, directives, standards, and initiatives "intentionally introduced" means "deliberately utilized during the manufacture of the components and/or for the formulation of a material or component where its continued presence is desired in the final product to provide a specific property, appearance or quality."*

This information is based on O&M Halyard, Inc.'s knowledge of these products at the time of publication.

If you have any questions or need additional information, please do not hesitate to contact us at PIQ@hyh.com or call us directly at (844) 425-9273.

–This statement is valid unsigned –

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March 14th, 2023

RE: **HALYARD* PURE**ZERO*** Nitrile Gloves**

Dear Valued Customer,

This letter is in response to your recent inquiry regarding storage conditions of **HALYARD* PURE**ZERO*** Nitrile Gloves.**

As displayed by the indications on the packaging labeling, **HALYARD* PURE**ZERO*** Nitrile Gloves** shall be stored in conditions where the product is kept dry (away from moisture), away from direct sunlight, away from sources of heat, and away from radioactive sources. To reduce the potential of accelerated aging, it is recommended to store the product at standard room temperature conditions, when possible.

Thank you for your interest in HALYARD* products. If you have any questions or need additional information, please do not hesitate to contact us at PIQ@hyh.com or call us directly at (844) 425-9273.

Sincerely,

Ryan Solan

Ryan Solan
R&D Senior Engineer
Global Research and Development
O&M Halyard, Inc.

Theresa McCoy

Theresa McCoy
Global Product Manager
Global Strategic Marketing
O&M Halyard, Inc.

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COPY-04606

September 24, 2023

RE: HALYARD* PUREZERO* HG3 and HG5 Nitrile Gloves

Dear Valued Customer,

This letter is in response to your recent inquiry regarding the presence of silicone for the PUREZERO* HG3 White Nitrile Gloves, PUREZERO* HG3 White SGX* Nitrile Gloves, PUREZERO* HG3 Light Blue Nitrile Gloves, PUREZERO* HG3 Sterile Light Blue Nitrile Gloves, PUREZERO* HG3 Sterile White Nitrile Gloves, HALYARD* PUREZERO* HG5 White Nitrile Gloves, and HALYARD* PUREZERO* HG5 Blue SGX* Nitrile Gloves.

According to ASTM E1252 Standard Practice for General Techniques for Obtaining Spectra for Qualitative Analysis, the results below reflect the detectable limits of silicone covering the spectral range from 4000cm^{-1} to 50cm^{-1}- via FT-IR (Fourier Transform Infrared Anaysis). The results below indicate that the HALYARD* PUREZERO* HG3 NITRILE GLOVES and HALYARD* PUREZERO* HG5 NITRILE GLOVES have no silicone detectable on the outside surface and inside surface of either gloves sampled per 3rd party testing.

The following product codes are applicable: PUREZERO* HG3 White Nitrile Gloves: CLN92512XS, CLN92512SM, CLN92512MD, CLN92512LG, CLN92512XL; PUREZERO* HG3 Light Blue Nitrile Gloves: CLN90511XS, CLN90511SM, CLN90511MD, CLN90511LG, CLN90511XL; PUREZERO* HG3 White SGX* Nitrile Gloves: CLN3031XS, CLN3031SM, CLN3031MD, CLN3031LG, CLN3031XL; PUREZERO* HG3 Sterile White Nitrile Gloves: CLN323260, CLN323265, CLN323270, CLN323275, CLN323280, CLN323290, CLN323210; PUREZERO* HG3 Sterile Light Blue Nitrile Gloves: CLN923260, CLN923265, CLN923270, CLN923275, CLN923280, CLN923285, CLN923290, CLN923210; HALYARD* PUREZERO* HG5 White Nitrile Gloves: CLN30512XS, CLN30512SM, CLN30512MD, CLN30512LG, and CLN30512XL; HALYARD* PUREZERO* HG5 Blue SGX* Nitrile Gloves: CLN92512XS, CLN92512SM, CLN92512MD, CLN92512LG, and CLN92512XL.

Fourier Transform Infrared Anaysis (FT-IR) Per ASTM E1252-98(21)	
Gloves	Results
PUREZERO* HG3 and HG5 Nitrile Gloves	No silicon detected

Thank you for your interest in Halyard products. If you have any questions or need additional information, please do not hesitate to contact us at PIQ@hyh.com or call us directly at (844) 425-9273.

Sincerely

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COPY-04662

Caitlin Senter
Director
Global Regulatory Affairs
O&M Halyard, Inc.

Nimia Maya
R&D Engineer II
Global Research and Development
O&M Halyard, Inc.

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COPY-04662

October 2023

TITLE: HALYARD* **PURE**ZERO***** HG3 & HG5 Nitrile Gloves Packaging

UPDATE: Alternate packaging supplier for HALYARD* **PURE**ZERO***** HG3 and HG5 Nitrile Gloves

WHAT YOU NEED TO KNOW:

There will be an alternate supplier for the packaging (poly bag, poly pouch and poly wallet) used for **HALYARD* PURE**ZERO*** HG3 and HG5 Nitrile Gloves**

WHAT YOU CAN EXPECT:

There is an alternate supplier being qualified to supply the poly bag, poly pouch and poly wallet used to package **HALYARD* PURE**ZERO*** HG3 and HG5 Nitrile Gloves.** The alternate supplier, TAKO Astatic Technology SDN BHD, will supply the same packaging as our current supplier, Tip Corporation SDN BHD. All specifications for the packaging, including the material, composition, dimensions, thickness, and ink, will be identical to the current packaging. There will be no change to the appearance of the packaging or printing. There will be no change to the sterilization process or the sterility assurance level of the products. Additionally, there will be no changes to the product codes or product labeling.

TIMING OF CHANGE: There will be a rolling change for the alternate supplier's packaging that will enter our inventory as early as December 2023. Both packaging vendors could be used based on supply chain availability.

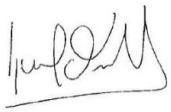
PRODUCTS IMPACTED BY THIS CHANGE:

PUREZERO* HG3 White Nitrile Gloves
CODE
CLN3031XS
CLN3031SM
CLN3031MD
CLN3031LG
CLN3031XL
PUREZERO* HG3 Light Blue Nitrile Gloves
CODE
CLN9031XS
CLN9031SM
CLN9031MD
CLN9031LG
CLN9031MD
PUREZERO* HG3 White SGX* Nitrile Gloves
CODE
CLN3231XS
CLN3231SM
CLN3231MD
CLN3231LG
CLN3231XL
PUREZERO* HG3 White Sterile Nitrile Gloves
CODE
CLN323260
CLN323265
CLN323270
CLN323275
CLN323280
CLN323285
CLN323290
CLN323210
PUREZERO* HG3 Light Blue Sterile Nitrile Gloves
CODE
CLN923260
CLN923265
CLN923270
CLN923275
CLN923280
CLN923285
CLN923290
CLN923210

PUREZERO* HG5 White Nitrile Gloves
CODE
CLN30512XS
CLN30512SM
CLN30512MD
CLN30512LG
CLN30512XL
PUREZERO* HG5 Blue SGX* Nitrile Gloves
CODE
CLN92512XS
CLN92512SM
CLN92512MD
CLN92512LG
CLN92512XL

Thank you for your interest in HALYARD* products. If you have any questions or need additional information including example certifications, please do not hesitate to contact us at PIQ@hyh.com, call us directly at 1-844-425-9273 or reach out to your O&M Halyard commercial representative or distributor.

Sincerely,



Gustavo Fiallos
Director Global Quality Operations, Owens & Minor

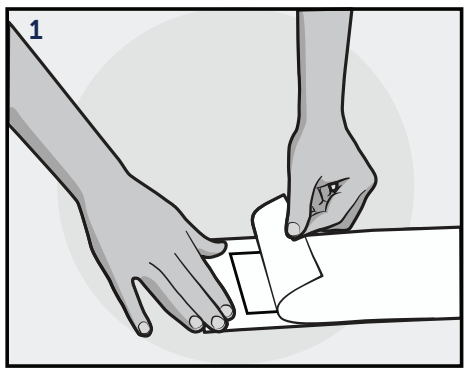
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PUREZERO*
Cleanroom Nitrile Gloves

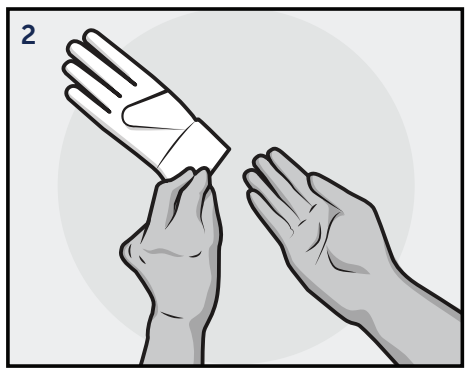
HALYARD*

GUIDE TO DONNING AND DOFFING OF STERILE CLEANROOM GLOVES

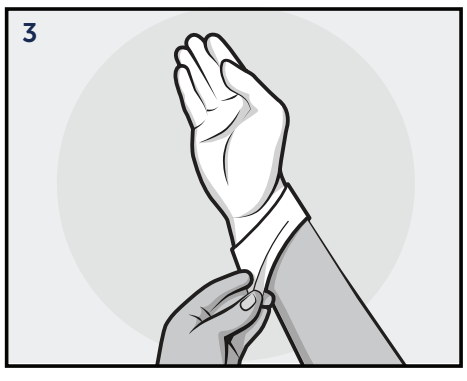
DONNING GLOVES



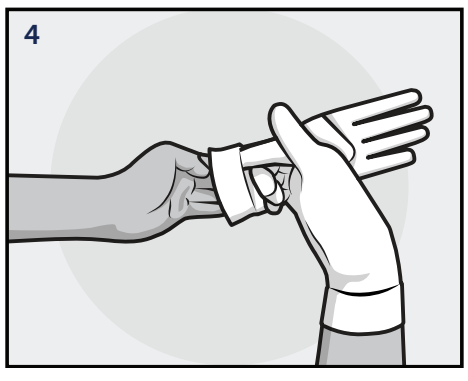
Peel open the pouch which keeps the product sterile. Open clear wallet.



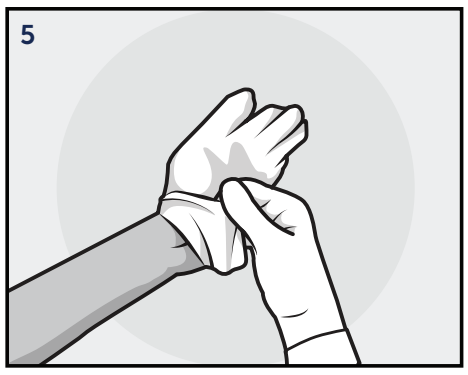
Grasp the interior of one glove and carefully pull over the opposite hand.



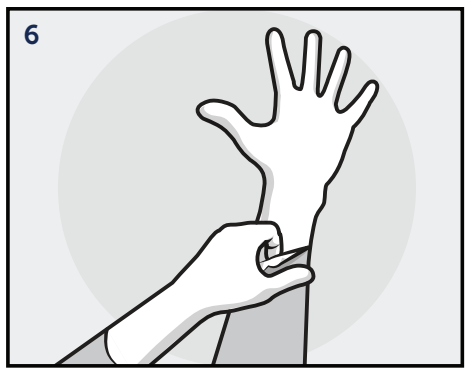
Gently ease the glove up over the hand and wrist, leaving the cuff folded.



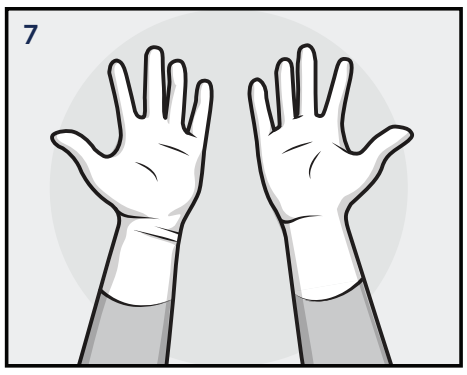
Lift the second glove under the folded cuff using gloved fingers.



Slide your second hand into the glove by holding it under the folded cuff, then unfold the cuff over the wrist and arm.

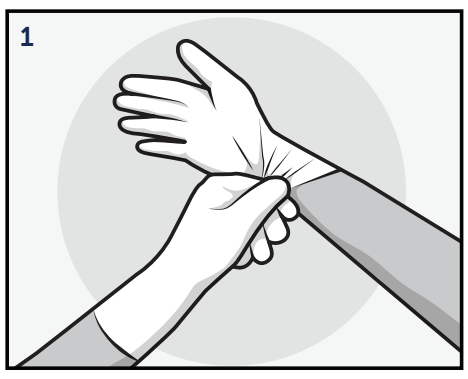


Using gloved fingers, unfold the second cuff over your wrist and arm.

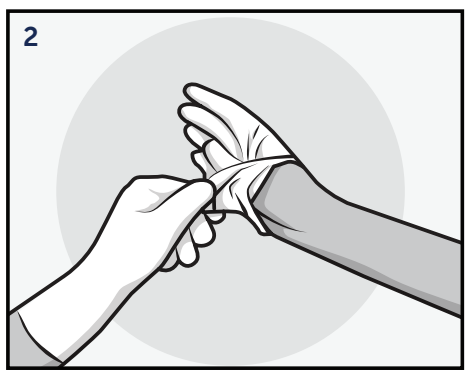


To remain sterile, do not touch anything with your gloved hands.

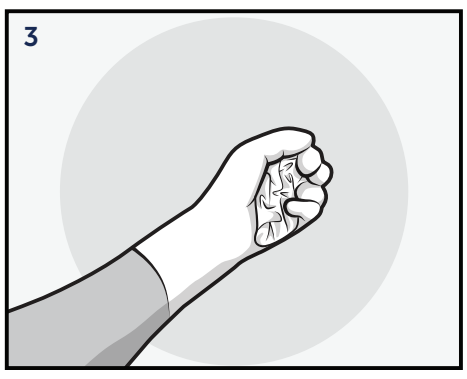
DOFFING GLOVES



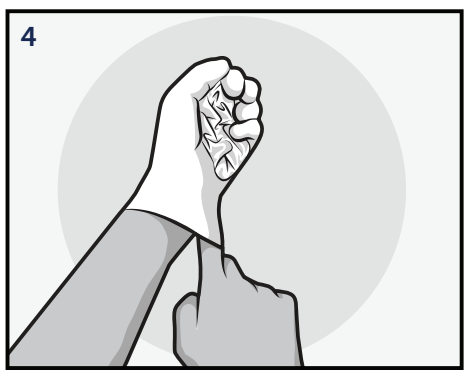
Grasp the exterior of one glove with your other gloved hand.



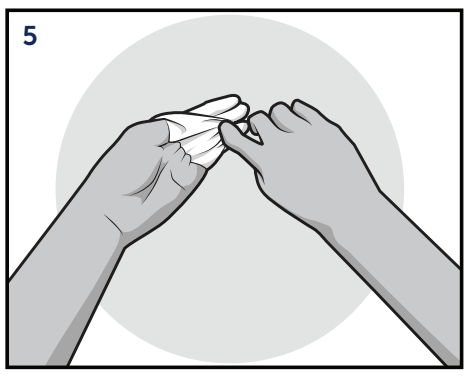
Carefully pull the glove off your hand, turning it inside-out.



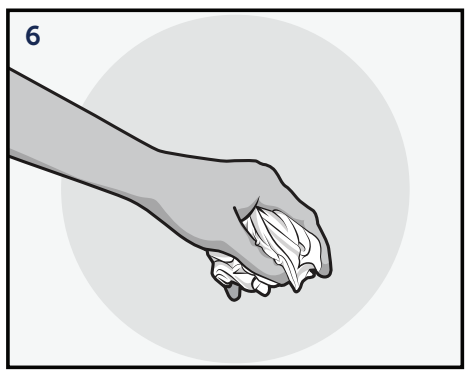
Ball the glove up and hold in your other gloved hand.



Slide your ungloved fingers into the opening of the other glove.



Carefully pull the glove off your hand, turning it inside-out and wrapping it around the doffed glove in your palm.



Discard appropriately.

HALYARD* recommends double donning/gloving. Don the gloves so that the sleeve cuffs are between the two pairs of gloves.

For more information, contact your HALYARD* representative or distributor, or visit **www.purezerogloves.com**

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