

## PROTECT YOUR FUTURE

GREEN

GENERAL

RISK

## INFORMATION

ecoSHIELD™ Eco Nitrile PF 250















- ⇒ Powder-free ambidextrous extra length (250-260 mm / 9.8"-10.2") non-sterile nitrile protective gloves.
- ⇒ Personal Protective Equipment Category III (PPE Complex Design) according to Regulation (EU) 2016/425.
- ⇒ Medical Device Class 1 (MDD) according to the Directive 93/42/EEC.
- ⇒ Fully compliant to the latest EU PPE norms relating to protective gloves against chemicals, micro-organisms and viruses.

DESCRIPTION			
FORMULATION Nitrile synthetic rubber (acrylonitrile butadiene).			
DESIGN	Green (Outer)/ White (Inner), ambidextrous, beaded cuff, textured fingertips.		
PACKAGING	150 gloves per dispenser - 10 dispensers per carton.		

SIZES	6/XS	7/S	8/M	9/L	10/XL
CODES	62 5121	62 5122	62 5123	62 5124	62 5125

STANDARDS	
CE REGISTRATION	PPE Category III (Complex Design) - Regulation (EU) 2016/425.  Notified Body No 0598: SGS Fimko Oy, Helsinki - FINLAND.  MDD Class 1 - Directive 93/42/EEC.
EU PPE NORMS	EN 420:2003+A1:2009, EN 421:2010, ISO 374-1:2016+A1:2018, EN 374-2:2014, ISO 374-4:2013, ISO 374-5:2016, EN 16523-1:2015+A1:2018 and ISO 16604:2004 Procedure B.
EU MDD NORMS	EN 455-1:2000, EN 455-2:2015, EN 455-3:2015 and EN 455-4:2009.
USA STANDARDS	ASTM D3767-03 (2014), ASTM D573-04 (2015), ASTM D412-16.
OTHER STANDARDS	EN1149-1/2/3 & 5, ISO 21171:2006, ISO 10993-10:2010.

QUALITY	
QUALITY ASSURANCE	Production management in accordance with ISO 9001:2015 and ISO 13485:2016.
TECHNOLOGY	twinSHIELD™ double-walled protection to offer a stronger glove and to reduce the risk of pinholes.  Two colours: green to make it easier to select according to the risk, combined with a soft and comfortable white interior.
ECOLOGICAL	50% more products in the same volume to save storage space. Ink on the packaging reduced by 60%. Packaging made from recycled cardboard. Supply chain optimized to reduce CO <sup>2</sup> emissions by more than 15% in the delivery of product.

DOCUMENTATION		
DECLARATION OF CONFORMITY	These documents can be freely downloaded from the product page on our website: www.shieldscientific.com.	
EU TYPE EXAMINATION CERTIFICATE	For an easy access, scan the QR code.	
PRODUCT INSERT		

## PHYSICAL PROPERTIES



NOM	IINAL THICKNESS	mm <sup>1</sup>	mil	Norm
$\Rightarrow$	Finger	0.17	6.7	ASTM D3767-03 (2014)
$\Rightarrow$	Palm	0.10	3.9	
$\Rightarrow$	Cuff	0.08	3.1	

<sup>&</sup>lt;sup>1</sup> Thickness (+/- 0.03 mm)

LE	NGTH	Minimum	Typical	Norm
$\Rightarrow$	From middle finger tip to edge of cuff 6/XS - 9/L	≥ 250 mm / 9.8"	255 mm / 10"	EN 420:2003+A1:2009
$\Rightarrow$	From middle finger tip to edge of cuff 10/XL	≥ 260 mm / 10.2"	265 mm / 10.4"	EN 420.2003+A1.2009

	ENGTH OPERTIES	Force at break (spec.)		Ultimate elongation (spec.)	Force at break (typical)	Norm
$\Rightarrow$	Before aging	≥ 6.0N	14 Mpa	≥ 500%	8.0N	EN 455-2:2015
$\Rightarrow$	After aging	≥ 6.0N	14 Mpa	≥ 400%	7.0N	ASTM D573-04 (2015) & ASTM D412-16

FREEDOM FROM HOLES	Performance	Norm	
⇒ Acceptable Quality Level (AQL)	< 0.25 <sup>2</sup> - Level 3	EN 374-2:2014 EN 455-1:2000	

<sup>&</sup>lt;sup>2</sup> AQL as defined per ISO 2859-1:1999 for sampling by attributes.

## PROTECTION PROPERTIES

RISKS	Description	Norm
MICRO-ORGANISMS	1000 ml water test. Performance level 3, AQL < 0.25 (inspection level G1).	EN 374-2:2014
VIRUSES	Viral penetration test using Phi-X174 bacteriophage according to ISO 16604:2004 Procedure B.	ISO 374-5:2016
CHEMICALS	Performance: Type B (JKPT). Permeation: Extensively tested. Online chemical resistance guide on www.shieldscientific.com. Degradation: Tested for determination of resistance to degradation by chemicals.	ISO 374-1:2016+A1:2018 EN 16523-1:2015+A1:2018 EN 374-4:2013
RADIOACTIVITY	Protection from radioactive contamination.	EN 421:2010
ESD	Tested for electrostatic properties.	EN 1149-1/2/3 & 5

ALLERGIES	
BIO-COMPATIBILITY	Demonstrated by skin irritation and sensitization tests in accordance with ISO 10993-10:2010.
ACCELERATORS	Free of Thiazoles and Thiurams. These chemical accelerators are excluded from the manufacturing process.
CHEMICAL ALLERGENS	Non-detectable levels using aqueous solution extraction (Phosphate buffered solution) and High Performance Liquid Chromatography (HPLC) assay method for quantitative analysis.
RESIDUAL POWDER	Powder-free to minimize the potential consequences of powder-borne dermatitis. Residual powder content is 1.0 mg/glove (typical) with a limit of 2.0 mg/glove (ISO 21171:2006).
LATEX PROTEIN	Latex-free.



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