

ENCORE® Latex Underglove

SURGICAL GLOVES

GENERAL DESCRIPTION

Material	Natural Rubber Latex
Powder Content	Powder-Free
Color	Green
Shape	Anatomic with Curved Fingers
Internal Glove Surface	Coated with HYDRASOFT™ Technology
External Glove Surface	Smooth with Micro-Textured Finish
Cuff Style	Beaded with SUREFIT™ Technology

PHYSICAL PROPERTIES

	Finger	Palm	Cuff
Thickness Typical Average (mm/mils)	0.19 ± 0.02 / 7.48 ± 0.79	0.19 ± 0.03 / 7.48 ± 1.18	0.16 ± 0.02 / 6.30 ± 0.79
Glove Length Typical (mm/in)	302 / 11.9		
Strength Minimum	Before Aging / Fresh		
	ASTM ¹	EN ²	ISO ³
Elongation At Break (%)	≥ 750	-	≥ 700
Force At Break (N)	-	≥ 9.0	≥ 12.5
Tensile Strength (MPa)	≥ 24	-	-
	After Aging / After Challenge		
	ASTM ¹	EN ²	ISO ³
Elongation At Break (%)	≥ 560	-	≥ 550
Force At Break (N)	-	≥ 9.0	≥ 9.5
Tensile Strength (MPa)	≥ 18	-	-
Comfort Maximum		ASTM ¹	ISO ³
Force at 300% Elongation (N)		-	≤ 2.0
Force at 500% Elongation (MPa)		≤ 5.5	-

ALLERGENIC SAFETY PROFILE

Vulcanization Chemical Accelerators	Zinc Dibutyldithiocarbamate (ZDBC), Zinc Diethyldithiocarbamate (ZDEC)
Primary Skin Irritation	Not considered a primary irritant as per FHSA Regulation guidelines 16 CFR 1500 / ISO 10993-10
Skin Sensitization	No evidence of delayed dermal contact sensitization as per ISO 10993-10
Protein Level	30 µg/g or less of total extractable protein as per EN 455, 50 µg/dm ² or less of total extractable protein as per ASTM D5712
Allergy Prevention	None

SAFETY STANDARDS

Freedom From Holes (AQL)	0.65 AQL
Sterilization Method	GAMMA irradiation (25 kGy)
Tested For Use With Chemotherapy Drugs	Yes, in accordance with ASTM D6978 (Not listed in the US FDA 510k)
Viral Penetration	Passes ASTM F1671 using Phi X 174, Passes ISO 16604 using Phi X 174

STANDARDS AND CERTIFICATIONS

Product Standards	ASTM D3577, ASTM D7160, EN 16523-1, EN 455 1-4, EN ISO 21420:2020, EN ISO 374-1:2016, EN 374-2 and -4, EN ISO 374-5:2016, ISO 10282
Quality / Environmental Standards	EN 556, ISO 11137-Part 1, ISO 13485, ISO 14001, ISO 9001
Medical Registration	BFE = Bacterial Filtration Efficiency, PFE = Submicron Particle Filtration Efficiency
Product Certification	EU: CE Marked to Medical Device Regulation (EU) 2017/745 (Class IIa), PPE Regulation (EU) 2016/425 (Cat. III risks) US: 510K # K051793, Class I / CA: MDALL # 14561, Class 2

¹ ASTM standards refer to ASTM D3577. ² EN standards refer to EN 455-2. ³ ISO standards refer to ISO 10282.

Contact your Ansell representative for ordering or more information.

North America

US: 800 952 9916
CA: 1-844-494-7854

insidesalesus@ansell.com

Central & South America

+52(442) 296 20 50
cslac@ansell.com

Brazil

+55-11-3356-3100
luvas.medicas@ansell.com

Europe, Middle East & Africa

+32 (0)2 528 74 00
info@ansell.eu

Asia Pacific

MY +603 8310 6688
apac.medical@ansell.com
CN: +86 21 38275000
infochina@ansell.com
JP: +813 5549 8151
info.medical.jp@ansell.com

Australia & New Zealand

+61 3 9270 7270
protection@ap.ansell.com



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S U R G I C A L G L O V E S

PACKAGING AND STORAGE

Packaging	50 pairs per box; 4 boxes per carton/case; 200 pairs per carton/case
Shelf Life	3 years
Shared Storage Instructions	Keep out of direct sunlight; store in a cool and dry place. Keep away from sources of ozone or ignition.
Disposal Recommendations	Gloves and pouch should be disposed of as clinical waste. Paper inner wrap, box and carton/case are recyclable but can be disposed of as clinical waste.

ANSELL PRODUCT CODES

Size	Code	Size	Code	Size	Code
5.5	330113055	6	330113060	6.5	330113065
7	330113070	7.5	330113075	8	330113080
8.5	330113085	9	330113090		



EN ISO 374-1:2016
Type B



K P T

EN ISO 374-5



VIRUS

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+52(442) 296 20 50

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Europe, Middle East & Africa

+32 (0)2 528 74 00

info@ansell.eu

Asia Pacific

MY +603 8310 6688

apac.medical@ansell.com

CN: +86 21 38275000

infochina@ansell.com

JP: +813 5549 8151

info.medical.jp@ansell.com

Australia & New Zealand

+61 3 9270 7270

protection@ap.ansell.com

ansell.com

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