

Fusion S-BFAP Sterile Disposable Neoprene (Polychloroprene Cleanroom Glove)

Sterile polychloroprene cleanroom gloves, combining tactility and comfort

- Assured comfort and tactility: Easy to don and doff and ideal for double gloving, BioClean[™] Fusion (Sterile) S-BFAP neoprene gloves' optimal thickness levels maintain comfort and tactility with prolonged use
- Allergy protection: As these powderfree disposable gloves are made from latex-free polymers, they eliminate Type I latex allergy risks
- Enhanced cuff design: Their beaded, extended-length cuff design combines a secure fit with additional arm coverage, boosting hand and wrist protection
- Minimized contamination risks: They come in non-particulating EasyTear packaging, reducing contamination risks in the cleanroom environment
- CAUTION: Please contact Ansell Customer Service for specific chemotherapy drug permeation times and recommendations

KEY FEATURES & BENEFITS

- Optimal thickness: Assured comfort and tactile sensitivity
- Latex-free neoprene formulation: Type I allergy risks eliminated
- Extended beaded cuff design: Enhanced hand protection

Industries

- Controlled and Critical Environments
- Production and Manufacturing
- Lab and R&D
- Sterile Processing Department







WWW.ANSELL.COM



Fusion S-BFAP Sterile Disposable Neoprene (Polychloroprene Cleanroom Glove)

TECHNICAL DATA SHEET

Color Gr Shape An Cuff Be	eoprene (Polychloroprene) reen mbidextrous eaded O 14001, Manufacturing QMS Audit Standards ISO 9001, PPE Regulation 2016 425 Module D, NEBB ertified Cleanrooms
Shape An Cuff Be	mbidextrous eaded O 14001, Manufacturing QMS Audit Standards ISO 9001, PPE Regulation 2016 425 Module D, NEBB
Cuff Be	eaded O 14001, Manufacturing QMS Audit Standards ISO 9001, PPE Regulation 2016 425 Module D, NEBB
	O 14001, Manufacturing QMS Audit Standards ISO 9001, PPE Regulation 2016 425 Module D, NEBB
Manufacturing/OMS ISC	
	E 0598, EN ISO 21420:2020, EN 421:2010, EN ISO 374-1:2016, EN ISO 374-5:2016, EN 455 Part 2, ategory III
	ne pair per inner PE wallet; one wallet per sealed EasyTear PE pouch; 10 pouches per sealed outer E bag; 20 outer bags per lined carton (200 pairs)
	eep out of direct sunlight; store in a cool and dry place. Keep away from sources of ozone or nition.
Country of Origin Inc	donesia
Available sizes 5.0	0-5.5, 6-6.5, 7-7.5, 8-8.5, 9.0, 10.0
Powder Content Po	owder-Free
External Glove Surface Te	extured Fingers
Internal Glove Surface Po	olymer Coated
Sterilization Method GA	AMMA irradiation (25 kGy)
Sterilization Minimum Dose 25	5kGy
Sterility Assurance Level)-6
Cleanroom Class Cla	lass 10/ISO Class 4 & EU GMP Grade A/B and other sterile cleanrooms
Shelf Life Fiv	ve (5) years from date of manufacture.
Tested for use with Chemotherapy Drugs	es
Protein Level N/	/A: contains no natural rubber latex
Anti-static Ye	es





Fusion S-BFAP

Sterile Disposable Neoprene (Polychloroprene Cleanroom Glove)

Physical Properties							Testing Method
Sizes	5.0-5.5	6.0-6.5	7.0-7.5	8.0-8.5	9.0	10.0	
Typical Length (mm/in)			EN 420				
Palm Width (mm/in)	76/3	86/3.4	95/3.7	106/4.2	115/4.5	120/4.7	
Freedom from Holes		0.	EN 374-2				
Typical Particle Count ≥0.5µm (counts / cm²)	850						IEST-RP-CC005.4
Target Single Wall Palm Thickness (mm/mil)	0.10 / 3.94						EN 455-2
Target Single Wall Finger Thickness (mm/mil)	0.12 / 4.72					EN 455-2	
Target Single Wall Cuff Thickness (mm/mil)	0.07/2.76					EN 455-2	

IONIC CONTENT

Concentration in µg/cm ²	Typical	Concentration in µg/cm ²	Typical	
Ammonium	0.003	Nitrate	0.884	
Bromide	Not Detected	Nitrite	Not Detected	
Calcium	0.652	Phosphate	Not Detected	
Chloride	0.194	Potassium	O.315	
Fluoride	Not Detected	Sodium	0.099	
Lithium	Not Detected	Sulphate	0.025	
Magnesium	0.007	Zinc	Not Detected	

ORDERING INFORMATION

SIZE	5.0-5.5	6.0-6.5	7.0-7.5	8.0-8.5	9.0	10.0
REORDER NO.	S-BFAP-5055	S-BFAP-6065	S-BFAP-7075	S-BFAP-8085	S-BFAP-9090	S-BFAP-1010

PERFORMANCE STANDARDS AND REGULATORY COMPLIANCE



For additional information visit us at www.ansell.com, or call us at

Europe, Middle East & Africa Region Ansell Healthcare Europe NV T: +32 (0) 2 528 74 00 F: +32 (0) 2 528 74 01

Asia Pacific Region

Ansell Global Trading Center T: +603 8310 6688 F: +603 8310 6699

North America Region Ansell Healthcare Products LLC US T; +1 800 800 0444 US F; +1 800 800 0445 CA T; +1-800-363-8340

Latin America & Caribbean Region Ansell Commercial Mexico S.A. de C.V. T: +52 442 248 1544 / 248 3133 Australia Ansell Limited T: +61 1800 337 041 F: +61 1800 803 578

UΚ Ansell Nitritex T: +44 1638 663338 F: +44 1638 668890

Ansell, [®] and [™] are trademarks owned by Ansell Limited or one of its affiliates. US Patented and US and non-US Patents Pending: www.ansell.com/patentmarking © 2022 Ansell Limited. All Rights Reserved.

Neither this document nor any other statement made herein by or on behalf of Ansell should be construed as a warranty of merchantability or that any Ansell product is fit for a particular purpose. Ansell assumes no responsibility for the suitability or adequacy of an end user's selection of gloves for a specific application.

