

Vasco® Sensitive

NON STERILE EXAMINATION AND PROTECTIVE GLOVES | DATA SHEET



B. Braun Melsungen AG confirms that

Vasco® Sensitive gloves comply with the following standards and regulations:

EC CERTIFICATES AND APPLIED STANDARDS

Medical Device Class I according to Medical Device Regulation (EU) 2017/745

EN 455 1-4, ASTM D3578

Personal Protective Equipment Category III according to Personal Protective Equipment Regulation (EU) 2016/425

EN 420, EN 374, EN 16523, ISO 16604, ASTM F1671

QUALITY CERTIFICATES

ISO 9001, ISO 13485

PERSONAL PROTECTIVE EQUIPMENT

Information and Declaration of Conformity according to PPER (EU) 2016/425:



www.bbraun.com/gloves-declarations-of-conformity

https://www.sritranggloves.com/en/update/document

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Vasco[®] Sensitive

NON STERILE EXAMINATION AND PROTECTIVE GLOVES | REGULATORY INFORMATION

MEDICAL DEVICE **INFORMATION**

MDR (EU) 2017/745 (CLASS I), EN 455







Conformity for food contact according to 1935/2004/EEC





FOOD COMPLIANCE

PERSONAL PROTECTIVE **EQUIPMENT INFORMATION**

Tested in accordance with: ISO 374-1/Type B





 (ϵ) 2777

PPE Regulation (EU) 2016/425 (Cat. III); EN 420:2003+A1:2009

Code letter	Test chemical	EN 374-1:2016 Permeation level	EN 374-4:2013 Mean degradation		
K	Sodium hydroxide 40%	Level 6	-18,2%		
Р	Hydrogen peroxide 30%	Level 3	3,3%		
Т	Formaldehyde 37 %	l evel 5	-28.2 %		

Tested acc. to EN 16523-1:2015

Resistance to virus

Performance levels acc. EN 374-1:2016 +A1:2018	1	2	3	4	5	6
Measured breakthrough times (mins)	> 10	>30	>60	> 120	> 240	>480

Degradation levels indicate the change in puncture resistance of the gloves after exposure to the challenge chemical. NOTE: Where the test specimens gave an increased puncture force after chemical exposure, the result is reported as a negative degradation.

pass

pass

ISO 374-5:2016





Resistance to bacteria and fungi

This information does not reflect the actual duration of protection in the workplace and the differentiation between mixtures and pure chemicals. The chemical and penetration resistance has been assessed under laboratory conditions from samples taken from the palm only and relates only to the chemical tested. It can be different if the chemical is used in a mixture. It is recommended to check that the gloves are suitable for the intended use because the conditions at the workplace may differ from the type test depending on temperature, abrasion and degradation. When used, protective gloves may provide less resistance to the dangerous chemical due to changes in physical properties. Movements, snagging, rubbing, degradation caused by the chemical contact etc. may reduce the actual use time significantly. For corrosive chemicals, degradation can be the most important factor to consider in selection of chemical resistant gloves. Before usage, inspect the gloves for any defect or imperfections.



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NON STERILE EXAMINATION AND PROTECTIVE GLOVES | TECHNICAL DATA

	SIZE	REF	_	GLOVE DIMENSIONS (EN 455)		
		100/90* pcs.	Width of p	palm	Total length	
	XS	6067500	<u>≤</u>	80 mm		
	S	6067526	80	± 10 mm		
	M	6067549	95 ± 10 mm		≥ 240 mm	
					£ 240 IIIII	
	L	6067565	110 ± 10 mm ≥ 110 mm			
	XL*	6067590				
PHYSICAL PROPERTIES				Min. specification	Typical value	
	Wall thicknes	SS	Finger	0.08 mm	0.14 mm	
			Palm	0.08 mm	0.12 mm	
			Cuff		0.08 mm	
	Force at brea	k	During shelf life	6 N	8.1 N after ageing	
	Elongation at	break	Before ageing	650%	816%	
			After ageing	500%	916%	
	Tensile streng	ıth	Before ageing	18 MPa	28 MPa	
	J	,	After ageing	14 MPa	25 MPa	
GLOVE DESIGN	Colour	natural white				
	Shape Cuff Surface finish Inner glove surface		straight fingers, ambidextrous fitting rolled rim, regular cuff fingertip textured polymer coated, powder-free			
	Outer glove surface		chlorinated			
GLOVE MATERIAL	Natural rubber latex (NRL)		Protein content ≤ 50 µg/g lower claims are not considered to be reliable given the expected process variation in manufacture and inter-laboratory testing (EN 455-3:2020)			
	Latex allergy risk		containing natural rubber latex which may cause allergic reactions including anaphylactic reactions			
ACCELERATORS	Zn-dithiocarb	pamate				
	Free of thiurames and mercaptobenzothiazoles (MBT)					
LOGISTIC INFORMATION	Dispenser pac	ck	100 / 90 pcs.	240 x 1	22 x 65 mm (L x W x H)	
	Transportatio	n carton	10 dispenser pack	s 340 x 2	49 x 250 mm (L x W x H)	
	Shelf life		3 years			
	Storage cond	itions	store at room temperature, protect from dust, humidity, sun light and ozone		d ozone	
			Packaging is made from recycled material			



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NON STERILE EXAMINATION AND PROTECTIVE GLOVES | BARRIER PROPERTIES – CHEMICALS



Tested by SATRA, UK in accordance with

EN 16523-1: Determination of material resistance to permeation by chemicals.

CHEMICAL	CAS REGISTRY NO.	PERMEATION PERFORMANCE LEVEL	BREAKTHROUGH TIME	
Acetone	67-64-1	not recommended	immediate	
Acetonitrile	75-05-8	not recommended	immediate	
Chloroform	67-66-3	not recommended	immediate	
Dichloromethane	75-09-2	not recommended	immediate	
Diethyl amine	109-89-7	not recommended	immediate	
Diethyl ether	60-29-7	not recommended	immediate	
Dimethylsulfoxide DMSO	67-68-5	not recommended	immediate	
Ethanol 70 %	64-17-5	not recommended	immediate	
Ethidium bromide 1 %	1239-45-8	level 6	> 480 min	
Ethyl acetate	141-78-6	not recommended	immediate	
Formaldehyde 37 %	50-00-0	level 5	> 240 min	
Gasoline	8032-32-4	not recommended	immediate	
Heptane-n	142-82-5	not recommended	immediate	
Hexane-n	110-54-3	not recommended	immediate	
Hydrogen peroxide 30 %	7722-84-1	level 3	> 60 min	
Methanol p.a.	67-56-1	not recommended	immediate	
Nitric acid 10%	7697-37-2	level 1	> 10 min	
Nitric acid 65%	7697-37-2	level 1	> 10 min	
Sodium hydroxide 40 %	1310-73-2	level 6	> 480 min	
Sulphuric acid 47 %	7664-93-9	level 1	> 10 min	
Sulphuric acid 96%	7664-93-9	level 1	> 10 min	
Toluene	108-88-3	not recommended	immediate	
Trichlorethane	71-55-6	not recommended	immediate	
Xylene	95-47-6	not recommended	immediate	