



PRODUCT SPECIFICATION

***Patterson[®] Latex Powder-free Examination Gloves
With Vitamin E and Aloe***

PRODUCT

Latex examination glove
Polymer coating with vitamin E and aloe
Medical grade
Non-sterile
Powder-free
Textured surface

COUNTRY OF ORIGIN

Thailand

INTENDED USE

This is a disposable device intended for medical purposes that is worn on the examiner's hand to prevent contamination between patient and examiner

MATERIAL

Natural rubber latex

CAUTION: This product contains natural rubber latex which may cause allergic reactions

OUTER SURFACE

Halogenation/siliconization and extensive washing in water
Inside coated with synthetic material with vitamin E and aloe
No donning powder used

COMPONENTS

Natural rubber latex
Sulfur
Zinc Oxide
Organic accelerators (dithiocarbamates, benzothiazolate)

SHAPE

Straight fingers
Thumb and fingers in one plane
Ambidextrous

CUFF

Beaded (rolled rim)

COLOR

Natural (white)

SIZES

Extra small (XS), small (S), medium (M), large (L), extra large (XL)

Sempermed USA, Inc., 13900 49th Street North, Clearwater, Florida 33762
Tel: 800.366.9545 Fax: 800.763.5491

MARKING

Packaging marked to designated size (gloves not marked)

PACKAGING AND LABELING

Reorder Number 088-4635,088-4643,088-4650,088-4668,088-4676
100 pieces per box (90 pieces XL), 1000 pieces per case (900 pieces XL)

CONTROL NUMBER (LOT NO.)

Each packing unit (dispenser box) and outer carton bears a control number

EXAMPLE: 092009 1234 0098

Key: 092009 Production month and year
1234 Internal running order number
0098 Carton number

QUALITY CHARACTERISTICS*

All listed standards are used in their latest edition. Current data on physical properties is available upon request.

DESCRIPTION	SPECIFICATION	ASSURANCE ACTION
<u>Dimensions</u>		ASTM D 3578
<i>Overall length</i>	220 mm min (XS, S) 230 mm min (M, L, XL)	
<i>Width</i>	70 mm +/- 10 mm (XS) 80 mm +/- 10 mm (S) 95 mm +/- 10 mm (M) 111 mm +/- 10 mm (L) 115 mm +/- 10 mm (XL)	
<i>Thickness (single wall)</i>	<i>Finger:</i> 0.08 mm/3.2 mils min. <i>Palm:</i> 0.08 mm/3.2 mils min.	
<u>Biocompatibility</u>		ASTM D 5712
<i>Proteins</i>	50 micrograms or less of total water-extractable protein per gram	
<i>Inside pH</i>	7.0 +/- 1	Test method A1
<u>Physical properties</u>		ASTM D 412
<i>Tensile strength (before aging)</i> <i>(after aging)</i>	18 MPa min. 14 MPa min.	
<i>Elongation (before aging)</i> <i>(after aging)</i>	650% min. 500% min.	

*minimum acceptable requirements of ASTM and FDA where applicable. Gloves offered by Sempermed USA, Inc. meet or exceed these physical requirements. Independent laboratory test results are available upon request.

PERFORMANCE REQUIREMENTS FOR QUALITY CHARACTERISTICS

For reference purpose in accordance with ISO 2859 "Sampling Procedures for Inspection by Attributes"

INTERNAL ATTRIBUTIVE RELEASE INSPECTION

Sampling for examination in accordance with ISO 2859

Unit for *inspection*: one (1) glove

If several defects are found on one glove, only the most serious defect (i.e. lowest category) is evaluated. The acceptance criteria is based on the number of defectives observed in a sample

FINAL GLOVE RELEASE

Assurance action

ASTM D 3578: "Standard Specification for Rubber Examination Gloves"

ASTM D 5151: "Standard Test Method for Detection of Holes in Medical Gloves"

Sampling inspection and final release information

Major defects: highest concern non-conformities which prevent correct use of the product. AQL 1.5 (inspection level GI for leaks)

Minor defects: non-conformities of a lesser degree of concern, which do not prevent correct use of the product. AQL 4.0 (inspection level GI for visual defects aggregated)

PACKAGING, MARKING, GOOD DELIVERY INSPECTION

Assurance Action

Set-up and patrol inspection at packaging

Supervision of vehicle or vessel loading

Sempermed USA, Inc. is a certified participant of C-TPAT (U.S. Customs-Trade Partnership Against Terrorism)

GOOD MANUFACTURING PRACTICE

The gloves are manufactured in compliance with ISO 9001, ISO 13485, and US FDA 21 CFR part 820

MICROBIOLOGICAL CLEANLINESS CONTROL

The bioburdens of the finished gloves are monitored and recorded. Unusual contaminants are identified.

CAUTION: Non-sterile examination gloves are used in a variety of circumstances, including procedures where the surface of the glove contacts wounds, body cavities, or other possible routes of contamination. If conditions warrant, the user may wish to minimize the risk of infection. In this case we recommend the decontamination of the gloves prior to use by disinfectants or other effective methods.

STORAGE

According to ISO 2230 for Vulcanized Rubber

Store in a dry, ventilated area

Avoid direct sunlight, fluorescent lighting, heat, and moisture

Do not store above 100° F (38° C) as this will lead to accelerated aging

Long-term storage can result in pleats and stickiness

Copper ions discolor the glove

END OF DOCUMENT

