

# **Technical product specification**

Product name	Vinyl powder free glove	Version / Index no:
Spec code	VCS-048CL-N-3CZ	VCS-048CL-N-3CZ_Version F_February
Date of issue	03.02.2020	2020_EN

#### **General information**

Type single use examination and disposable protective glove, non sterile

Labelling information printed on dispenser box
Shape ambidextrous - straight fingers

Material Polyvinylchlorid (PVC) [not made with natural rubber latex]

Colour transparent

Inside polymer coated / powder free

Outside no treatment
Cuff / surface rolled cuff / smooth

Shelf life 5 years

**Available sizes** XS (5-6) S (6-7) M (7-8) L (8-9) XL (9-10)

Dimensions, physical properties and biocompatibility

Glove length median ≥ 240 mm (according to EN 455-2)

Minimum wall<br/>thicknessat finger0.10 mm (double measured) / 0.05 mm (single measured)0.14 mm (double measured) / 0.07 mm (single measured)

at cuff 0.08 mm (double measured) / 0.04 mm (single measured)

Glove width according to EN 455-2: median XS  $\leq$  80 mm, S 80  $\pm$  10 mm, M 95  $\pm$  10 mm, L 110  $\pm$  10 mm, XL  $\geq$ 

110 mm

Force at Breakmedian ≥ 3.6 N (during shelf life according to EN 455-2)Tensile Strengthmin. 9 MPa after aging (according to ASTM D5250)

Elongation at Break min. 300% after aging (according to ASTM D5250)

Residual powder / ≤ 2 mg (according to EN 455-2)

**Powder content** 

### Performance requirements and inspection levels

Freedom from holes (Barrier) AQL ≤ 1.5

(as per EN 455-1, sampling in accordance with ISO 2859-1, G-1)

Dimensions and physical properties AQL 4.0 (as per ASTM D5250, sampling in accordance with ISO 2859-1, S-2)

Standards, guidelines & quality certificates

Quality certification ISO 9001, ISO 13485, ISO 14001

Conformity to regulations Upon request:

Medical Device Regulation (EU) 2017/745: Class I
 PPE Regulation (EU) 2016/425: Category I

- Regulation (EC) 1935/2004 on Food Contact Materials and Regulation (EU)

10/2011 on plastic materials for food contact

Conformity to standards EN 420 (subject to labelling), EN 455-1, EN 455-2, EN 455-3, EN 455-4, ISO

2859-1, ASTM F1671



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### Instructions and additional statements

Storage instruction

Store in original packaging in a dry and dark place at 10 °C to 30 °C. Refer to guidelines of storage of rubber products as described in ISO 2230:2002. Ensure that storage area is kept cool, dry and dust free, avoid ventilation and storage close to photocopy equipment. Copper ions discolour the glove. Protect gloves against ultraviolet light sources, such as sunlight and oxidizing agents. Storage above 30 °C will lead to accelerated aging and should be avoided.

Cautionary statement and ingredient information

No allergenic ingredients known.

For further information, a list of substances contained in the glove is available

upon request.

Reporting system

Medical device vigilance and reporting system

According to the official reporting criteria of the Medical Device Regulation, incidents caused by examination gloves must be reported immediately to our

L. Rieger

Head of Product Management

Medical Device Reporting team. E-Mail:

J. Glantschnig | Head of Regulatory Affairs and Contract Management

sempermed.complaints@semperitgroup.com or Tel.: +43 2630 310 0

Remark

Replaces all previous versions.

All standards references refer to the date of document issue.

Sempermed