

Technical product specification

Product name	Nitrile powder free glove	Version / Index no:
Spec code	NOF-040BK-N-3CZ	NOF-040BK-N-3CZ_Version H_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 Nitrile Butadiene Rubber (NBR) [not made with natural rubber latex]

Colour black Inside powder free **Outside** no treatment

Cuff / surface rolled cuff / finger textured

Shelf life 3 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 0.18 mm (double measured) / 0.09 mm (single measured) at finger thickness at palm 0.12 mm (double measured) / 0.06 mm (single measured) 0.10 mm (double measured) / 0.05 mm (single measured) at cuff

Glove width according to EN 455-2: median XS ≤ 80 mm, S 80 ± 10 mm, M 95 ± 10 mm, L 110 ± 10 mm, XL ≥

110 mm

Force at Break median ≥ 6 N (during shelf life according to EN 455-2) **Tensile Strength** min. 14 MPa after aging (according to ASTM D6319) Elongation at Break min. 400% after aging (according to ASTM D6319)

Residual powder / ≤ 2 mg (according to EN 455-3)

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 D6319, 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 or III - Regulation (EC) 1935/2004 on Food Contact Materials

Conformity to standards EN 420, EN ISO 374-1, EN 374-2, EN 16523-1, EN 374-4, EN ISO 374-5

(subject to labelling),

EN 455-1, EN 455-2, EN 455-3, EN 455-4, ISO 2859-1, ASTM D6319, ASTM

F1671



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Instructions and additional statements

Storage instruction Store in original packaging

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

This product contains accelerators (Dithiocarbamate types, Zincmercaptobenzothiazol) not to be used in a hypersensitivity of these

substances.

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

Medical Device Reporting team. E-Mail:

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

A./Wöss Director J. Glantschnig | Head of Regulatory Affairs and Contract Management

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Remark

Replaces all previous versions.

All standards references refer to the date of document issue.