
Inlimited
PROMOTION

CRANBERRY EVOLVE 300 NITRILE POWDER FREE GLOVES

The Evolve 300 Nitrile Powder Free Examination Gloves are Cranberry's latest and softest gloves yet, combining comfort and tensile strength without sacrificing tactile sensitivity. Cranberry's exclusive EvoSoft™ formulation gives Evolve a unique silk-like attribute that is both soft and strong with textured fingertips for precise gripping under all operating conditions.

CRANBERRY EVOLVE 300 NITRILE POWDER FREE GLOVES

FORMFITTING
EvoSoft™ formulation delivers a silk-like attribute, soft and strong. Fingertip textured in a Royal Blue color.

SILK THIN
Our stretchiest gloves yet, high tensile strength and excellent comfort, while not sacrificing tactile sensitivity.

300 GLOVES PER BOX
Ultra 300 Saver Pack reduces storage space and packaging waste.

CRANBERRY EVOLVE 300 NITRILE POWDER FREE GLOVES

- Exclusive EvoSoft™ formulation delivers silk-like attribute, soft yet strong.
- Superior tensile strength for maximum stretch.
- Strong Cuff for Tear Resistance.
- Ultra Space Saver Pack (300 gloves per box).
  - Sizes XS, S, M, L, XL
  - 300 gloves/box, 10 boxes/case; *XL: 250 gloves/box.

CRANBERRY EVOLVE 300 NITRILE POWDER FREE GLOVES

300 evolve®
Nitrile Powder Free
Examination Gloves

Features	Benefits
Exclusive EvoSoft™ formulation	Provides unprecedented soft and strong comfort during extended wear
Textured Fingertips	Delivers good grip
300 Space Saver Pack	Maximize storage space and effciency

Physical Data

Dimension	Extra-Small	Small	Medium	Large	Extra-Large
Length (mm) min.	230	230	230	230	230
Palm Width (mm)	70±10	80±10	95±10	110±10	120±10
Thickness (mm) min.					
~ Finger	0.05	0.05	0.05	0.05	0.05
~ Palm	0.05	0.05	0.05	0.05	0.05
~ Cuff	0.04	0.04	0.04	0.04	0.04
Tensile Strength (MPa), min					
~ Unaged	16	16	16	16	16
~ Aged	16	16	16	16	16
Elongation (%), min					
~ Unaged	500	500	500	500	500
~ Aged	400	400	400	400	400

2020

CERTIFICATE OF REGISTRATION

This certifies that:

is registered with the U.S. Food and Drug Administration for FY 2020 pursuant to Title 21, 807 et seq. o
United States Code of Federal Regulations:

Establishment Registration:

DUNS No.:	67-169-5763
Device Classification Name:	NITRILE EXAMINATION GLOVE
Product Code:	LYY
Regulation Number:	880.6250
Official Correspondent and U.S. Agent:	Registrar Corp

Registrar Corp will confirm that such registration remains effective upon request and presentation o certificate until the end of the year stated above, unless said registration is terminated after issuance o certificate. Registrar Corp makes no other representations or warranties, nor does this certificate mak representations or warranties to any person or entity other than the named certificate holder, for whose benefit it is issued. This certificate does not denote endorsement or approval of the certificate-holder's d or establishment by the U.S. Food and Drug Administration. Registrar Corp assumes no liability to any pa or entity in connection with the foregoing.

Pursuant to 21 CFR 807.39, "Registration of a device establishment or assignment of a registration nu does not in any way denote approval of the establishment or its products. Any representation that creat impression of official approval because of registration or possession of a registration number is misle and constitutes misbranding."

The U.S. Food and Drug Administration does not issue a certificate of registration, nor does the U.S. and Drug Administration recognize a certificate of registration. Registrar Corp is not affiliated with the Food and Drug Administration.

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David Lennarz
Executive Director
Registrar Corp
Dated: January 20, 20

Certificate of Registration

In accordance with European Communities Council Directive 93/42/EEC as amended by 2007/47/EC concerning Medical Devices as transposed into European national law by the member states.

We hereby declare that:

- **An examination has been made of this organisation's Declaration of Conformity(s) and where appropriate Notified Body certification(s) exist.**
- **The EU Authorised Representative contract has been fulfilled.**
- **Device registrations for the medical devices mentioned within this certificate have duly been completed with an EU Competent Authority.**

Therefore, these devices have met the requirements of the council directive and the CE mark may be applied to the products listed below.

Certificate No: CE/THA/1999/09/03	Issue Date: 01st April 2020	Expiry Date: * 31st March 2021

**Please note, due to the implementation date of the new medical device regulation (EU 2017/745) this certificate is subject to a review of the client's technical documentation before the 26th May 2020, whereupon a new Certificate of Registration is issued once compliance to the medical device regulation has been achieved.*

Legal Manufacturer	EU Authorised Representative (EC REP)
Thailand	Advena Limited, Tower Business Centre, 2nd Flr, Tower Street, Swatar, BKR 4013 Malta.

Product Details, Names or Trade Names	MCCAA Device Registration Reference(s)
Examination Gloves	DVC-MT-20-02-000061

Competent Authority
Malta Competition and Consumer Affairs Authority (MCCAA) Mizzi House, National Road, Blata l-Bajda, HMR 9010 Malta. Tel: +356 2395 2000 Email: info@mccaa.org.mt

This certificate is issued by:	Authorised Signature:
Advena Limited Tower Business Centre, 2nd Flr, Tower Street, Swatar, BKR 4013. Malta. Tel: +44 1926 800153 Email: info@advenamedical.com Registered in Malta No. C 76865	Anthony Kirby – Managing Director (Malta)

This certificate is subject to the organisation maintaining their documentation in compliance with the directive stated in this certificate.

CERTIFICATE OF COMPLIANCE

✓ This is to certify that the manufacturing/distribution system of :

[redacted]

✓ Has been found to conform to the Manufacturing , marketing and quality management system standard:

ISO 9001: 2015

✓ This certificate is valid for the following scope:

MARKETING OF SURGICAL, EXAMINATION AND INDUSTRIAL
LATEX GLOVES, NITRILE GLOVES
VINYL GLOVES
PE GLOVES
IN BOTH POWDERED AND NON POWDERED FORMS

This certificate was issued electronically and remains the property of BSI and is bound by the

[redacted]

Singapore is a subsidiary of British Standards Institution.

For and on behalf of BSI:

Managing Director, Philip Kee

Latest Issue: 27JAN/2020

Originally registered: 27/JAN/2020 Expiry Date: 27/JAN/2021

[redacted]

EC CERTIFICATE

PRODUCTION QUALITY ASSURANCE

DIRECTIV 93/42/EEC ON MEDICAL DEVICE (MDD),ANNEX V

(DEVICES IN CLASS IN STERILE CONDITIONS, STERILIZED SYSTEMS OR PROCEDURE PACKS)

[redacted]

- MANUFACTURER/APPLICANT

[redacted]

- EC REPRRESENTIVE:

[redacted]

- PRODUCTS/CATEGORY

DISPOSABLE MEDICAL LATEX EXAMINATION GLOVES,
NITRILE EXAMINATION GLOVES
VINYL EXAMINATION GLOVES
LATEX SURGICAL GLOVES

- ATTESTATION

THE CERTIFICATION BODY OF BSI SINGAPORE DECLARES THAT THE AFOREMENTIONED APPLICANT/MANUFACTURER HAS IMPLEMENTED A QUALITY ASSURANCE SYSTEM FOR MANUFACTURE IN ACCORDANCE WITH MDD ANNEX V. THIS QUALITY ASSURANCE SYSTEM COVERS THOSE ASPECTS OF MANUFACTURE CONCERNED WITH SECURING AND MAINTAINING STERILE CONDITIONS OF THE RESPECTIVE DEVICES/DEVICE CATERGORIES AND CONFORMS TO THE REQUIREMENT OF THIS DIRECTIVE. IT IS SUBJECT TO PERIODICAL SUIVELLANCE. SEE ALSO NOTES OVERLEAF

Singapore Headquarters: 4[redacted]

subsidiary of British Standards Institution.

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Managing Director, Philip Kee

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