VITACORE

EU DECLARATION OF CONFORMITY

This declaration of conformity is issued under the sole responsibility of the manufacturer:

Vitacore Industries Inc.
(151-8518 Glenlyon Parkway, Burnaby, BC V5J 0B6, CANADA)

Object of the declaration:
CAN99 Particulate/Surgical Respirator

The object of the declaration described is in conformity with the relevant Union harmonisation legislation:
PPE Regulation 2016/425 & EN149:2001+A1:2009 FFP3 NR.

References to the relevant harmonised standards used, including the date of the standard, or references to the other technical specifications, including the date of the specification, in relation to which conformity is declared:
PPE Regulation 2016/425 & EN149:2001+A1:2009 FFP3 NR .

Where applicable, the notified body **BSI Group CE 2797**, Say Building, John M. Keynesplein 9, 1066 EP Amsterdam, Netherlands performed the EU type examination (Module B) and issued the EU type-examination certificate, **CE 741958.**

Where applicable, the PPE is subject to the conformity assessment procedure -conformity to type based on internal production control plus supervised product checks at random intervals (Module C2), Certificate **CE 741959** under surveillance of the notified body BSI Group NB number: **CE 2797.**

Authorized by:

Yang Fei,

Director of Research and Development
Vitacore Industries Inc.
8518 Glenlyon Parkway, V5J 0B6, CANADA, on 7/4/2021

EU Type Examination Certificate

This is to certify that:

Vitacore Industries Inc
#151-8518 Glenlyon Parkway
Burnaby
Canada
British Columbia
V5G 0B6
Canada

Holds Certificate Number: CE 741958

In respect of:

Vitacore Industries Inc range of filtering half mask to EN149:2001+A1:2009 Respiratory protective devices. Filtering half mask to protect against particles model(s) CAN99

on the basis that BSI carried out the relevant Type Examination procedures under the requirements with the Regulation (EU) 2016/425 of the European Parliament and Council relating to Personal Protective Equipment Regulation (PPE) Annex V (Module B) and meets the relevant health and safety requirements specified in Annex II

For and on behalf of BSI, a Notified Body for the above Regulation (Notified Body Number 2797):

Drs. Dave Hagenaars, Managing Director

First Issued: 2021-04-22
Latest Issue: 2021-04-22

Effective Date: 2021-04-22
Expiry Date: 2026-04-22

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EU Type Examination Certificate

No. CE 741958

Product specification

The respiratory protective filtering half mask covered by the scope of this Module B Certificate and the Harmonized European Standards to which the products are approved are to the following specifications:

Product Name	Particulate Respirator
Product Type:	Disposable filtering half mask respirator with no exhalation valve.
Models:	CAN99
Classification:	FFP3 NR
Technical Specification:	EN 149:2001+A1:2009 Respiratory protective devices. Filtering half mask to protect against particles.
Product Description:	The respirator is a non-reusable half mask for protection against particles, secured to the face of the user by a pair of elasticated head straps, and has no exhalation valve. The respirator is FFP3 NR class

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EU Type Examination Certificate

No. CE 741958

Technical File Reference: TF001

Certificate Amendment Record and BSI internal Review relating to this Certificate

Issue date	Comments	BSI Review No.
April 2021	First issue.	2797:21:3347364

Note: The Certificate holder is responsible for keeping the Notified Body advised of changes to any aspect of the overall process used in the manufacture of the product.

Monitoring of manufactured PPE

The validity of the Certificate for the products is also dependent on the maintenance of the EU Conformity to Type based on Internal Production Control plus supervised product checks at random intervals, Annex VII (Module C2) as referenced on BSI issued Certificate CE 741959

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Conformity to Type based on Internal Production Control plus supervised product checks at random intervals

This is to certify that:

Vitacore Industries Inc
#151-8518 Glenlyon Parkway
Burnaby
Canada
British Columbia
V5G 0B6
Canada

Holds Certificate Number: CE 741959

In respect of:

The respiratory protective device covered by the scope of this Module C2 Certificate and the Technical Specification to which the product is manufactured are as follows:

on the basis that BSI carried out the supervised production checks at random intervals under the requirements with the Regulation (EU) 2016/425 of the European Parliament and Council relating to Personal Protective Equipment Regulation (PPE) Annex VII (Module C2)

For and on behalf of BSI, a Notified Body for the above Regulation (Notified Body Number 2797):

Drs. Dave Hagenaars, Managing Director

First Issued: 2021-04-22
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PRODUCTS
RvA C 646

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Conformity to Type based on Internal Production Control plus supervised product checks at random intervals

No. CE 741959

Manufacturing Site:

Vitacore Industries Inc.
8518 Glenlyon Parkway
V5J 0B6
Canada

Product details

The respiratory protective device covered by the scope of this Module C2 Certificate conform to the following standards:

Standard	Product type
EN 149:2001+A1:2009	Respiratory protective devices – Filtering half mask to protect against particles.

Certificate Amendment Record and BSI internal Review relating to this Certificate

First Issued: 2021-04-22 Effective Date: 2021-04-22

Latest Issue: 2021-04-22 Expiry Date: 2026-04-22

Conformity to Type based on Internal Production Control plus supervised product checks at random intervals

No. CE 741959

Issue date	Comments	BSI Review No.
April 2021	First issue.	2797:21:3347359

Certificate validity

The Certificate holder is responsible for ensuring that the Notified Body is advised of changes to any aspects of the overall quality system utilized in the manufacture of the products, failure to do so could invalidate the Certificate in respect of product manufactured after the introduction of such changes.

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