EU Type Examination Certificate

This is to certify that:

Prenger Healthcare B.V.
PO BOX 74
2280 AB Rijswijk
The Netherlands

Holds Certificate Number: CE 751967

In respect of:

Protective Half Masks
To EN 149:2001 + A1:2009
See Continuation Sheets for Details

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: 2021-07-23
Latest Issue: 2021-07-23

Effective Date: 2021-07-23
Expiry Date: 2026-07-23

PRODUCTS
RvA C 646

...making excellence a habit.™

EU Type Examination Certificate

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Product Specification

Product Type:	Filtering half masks to protect against particles.
Model(s):	2003 2004 2005 (mask with exhalation valve)
Product description:	The particulate respirators are designed to protect against solid and non-volatile liquid particles. The masks are held on the face by a pair of elasticated straps. The models are single shift devices denoted by the classification symbol NR. The Dolomite Clogging option is denoted by the classification symbol D.
Technical specifications:	EN 149:2001+A1:2009 – Respiratory Protective Devices. Filtering half masks to protect against particles.

	Model	**Classification**
EN 149 Classification	2003	**FFP2 NR**
	2004	**FFP2 NR D**
	2005	**FFP3 NR D**

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Certificate Administration Details

Product initially approved on BSI certificates: CE 732331 and CE 739472

Certificate Amendment Record:

Issue date	Comments	BSI Review Number
July 2021	First issue.	2797:21:3476698

Certificate validity

The Certificate holder is responsible for ensuring that the Notified Body is advised of changes to any aspect of the overall processes utilised in the manufacture of the product, failure to do so could invalidate the Certificate in respect of product manufactured following the introduction of such changes.

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 D) as referenced on BSI issued Certificate CE 739473.

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