



Conformity to Type based on Quality Assurance of the **Production Process**

This is to certify that:

Supreme Visors Limited Unit 2 Aston Fields Road Whitehouse Industrial Estate Runcorn WA7 3DL United Kingdom

Holds Certificate Number:

CE 754388

In respect of:

The manufacture of Respiratory Protective Equipment to the standards specified on the continuation sheet.

on the basis that BSI carried out the quality assurance assessment under the requirements with the Regulation (EU) 2016/425 of the European Parliament and Council relating to Personal Protective Equipment Regulation (PPE) Annex VIII (Module D)

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

Drs. Dave Hagenaars, Managing Director

First Issued: 2021-08-13 Latest Issue: 2021-08-13 Effective Date: 2021-08-13 Expiry Date: 2026-08-13

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...making excellence a habit."

This certificate has been issued by and remains the property of BSI Group The Netherlands B.V., John M. Keynesplein 9, 1066 EP Amsterdam, The Netherlands and should be returned immediately upon request. To check its validity telephone +31 20 3460780. An electronic certificate can be authenticated <u>online</u>.

BSI Group The Netherlands B.V., registered in the Netherlands under number 33264284, at John M. Keynesplein 9, 1066 EP Amsterdam, The Netherlands A member of BSI Group of Companies.

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No. CE 754388

Manufacturing location:

Supreme Visors Limited Unit 2 Aston Fields Road Whitehouse Industrial Estate Runcorn WA7 3DL United Kingdom

Product Specifications

The Respiratory Protective Equipment covered by the Module D scope of this Certificate conform to the following standards:

Standard

Product Type

EN 14594:2005 Respiratory protective devices. Continuous flow compressed air line breathing apparatus

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Certificate Amendment Record:

Issue date Comments

August 2021 First issue **BSI Review No.**

2797:21:3492918

Certificate validity

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

The validity of the Certificate is also dependent on the maintenance of an ISO 9001 quality system certified by a recognized certification organisation.

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