



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

This is to certify that:

SPRO Medical Products (Xiamen) Co., Ltd. No.139 Factory Bldg TongAn Garden **TongAn Industrial Area** Xiamen Fujian 361100 China

Holds Certificate Number:

CE 754045

In respect of:

#### For the manufacture of Respiratory Protective Devices as detailed 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: 2022-02-21 Latest Issue: 2022-02-21 Effective Date: 2022-02-21 Expiry Date: 2027-02-21

Page: 1 of 3



## ...making excellence a habit."

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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.

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

No. CE 754045

#### **Design location:**

SPRO Medical Products (Xiamen) Co., Ltd. No.139 Factory Bldg TongAn Garden TongAn Industrial Area Xiamen Fujian 361100 China

#### **Production Location:**

SPRO Medical Products (Xiamen) Co., Ltd. No.139 Factory Bldg TongAn Garden TongAn Industrial Area Xiamen Fujian 361100 China

#### **Product Specification:**

The products covered by the scope of this Certificate conform to the following standards:

Standard

#### **Product Type**

EN 149:2001 + A1:2009

Filtering half mask, respiratory protective device, to protect against particles.

- Last item -

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Page: 2 of 3

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No. CE 754045

#### **Certificate Amendment Record:**

Issue	date	Comments
133uc	uute	Commence

February 2022 First issue **BSI Review No.** 

2797:22:3490934

#### **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 a quality system certified by a recognized certification organisation.

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Page: 3 of 3

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