Certificate Number: PPE20161886

EU TYPE-EXAMINATION CERTIFICATE

This is to certify that INSPEC International B.V., Notified Body 2849, has evaluated the Personal Protective Equipment type(s) in respect of the product detailed on this certificate and deemed it(them) to be in compliance with Annex V (Module B) of the Personal Protective Equipment Regulation (EU) 2016/425 and the applicable Essential Health & Safety Requirements.

Manufacturer: **SPRO MEDICAL PRODUCTS (XIAMEN) CO., LTD**
West of 1-5th Floor, No. 139 Factory Bldg
TongAn Garden, TongAn Industry Area
Xiamen City
Fujian Province
China

Authorised Representative: **SPRO MEDICAL (EU) SP.Z O.O.**
Ul. Tadeusza Kos'ciuszki
14/4,84-200
Wejherowo
Poland

Compliance with the applicable Essential Health & Safety Requirements has been demonstrated as above, including examination in accordance with the harmonised standard below:

EN 149:2001 + A1:2009

Product description: Respiratory Protective Devices – Filtering half masks;

Models: GL001A and GL001

Date of initial certification: 11 March 2020
Date of current issue: 25 January 2022
Date of expiry: 11 March 2025

A. Diamond
Certification Manager

PRODUCTS
RvA C 657

Certificate Number: PPE20161886

Product details

Model identification:

Model	Class	Exhalation valve
GL001A	FFP2 NR	No
GL001	FFP2 NR	Yes

Key: NR Non-re-useable / limited to single shift use only

Technical file reference: TF20161886

Test reports: 1.18.10.73

Category: III

Category III product must also have an approval decision from a Notified Body to demonstrate conformity with Module C2 or D of Council Regulation (EU) 2016/425.

Classification: See above table

Accessories: None

Certificate amendment record

Date	Description
25/01/2022	Addition of Authorised Representative. Previous version was issued by NB0194.
11/03/2020	Initial issue

Conditions attached to the issue of this certificate:

1. The manufacturer / authorised representative shall undertake to fulfil the obligations arising out of the Personal Protective Equipment Regulation (EU) 2016/425, and with INSPEC's Regulations governing this Module.

2. The manufacturer / authorised representative shall inform INSPEC of any planned changes to the product or technical file which may affect the validity of this certificate, before any such change is made.

3. Marking and instructions have been assessed in the English language only. It is the manufacturer's / authorised representative's responsibility to obtain and supply language versions acceptable to the country where the product is to be sold.

4. For category III product, the manufacturer must obtain and maintain an approval decision to Module C2 or Module D prior to placing product on the Union market.

5. This certificate remains the property of INSPEC and may be withdrawn if any of the conditions attached to its issue are not complied with.

6. This certificate may be copied or reproduced by the certificate holder, complete and without omissions or additions, and in accordance with INSPEC's terms of business.