INSPEC

副COPY本

Certificate Number: PRE19161717

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:

Fido Masks Co., Ltd.
No. 31, Gong 1st Rd.,
Rinan Vil., Dajia Dist.,
Taichung City
43767
Taiwan

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

EN149:2001 + A1:2009

Product description: Respiratory Protective Devices – Filtering Half Masks;

Model: F539V

Date of initial certification: 24 October 2019
Date of current issue: 26 April 2024
Period of validity: 24 October 2024 - 24 October 2029

Certificate Signatory

PRODUCTS
RvA C 657

Product details

Model identification: F539V

Model description: Filtering half mask with valve fitted.

Technical file reference: TD-F539V (TF19161717)

Test reports: 1.15.08.01, 1.21.10.19

Category: III (three)

Classification: FFP3 NR

Options: None

Accessories: None

Key:

NR	Single shift use

Certificate Number: PPE19161717

Certificate amendment record

Date	Description
24/10/2019	Initial issue.
31/07/2023	Addition of alternative filter media and test report 1.21.10.19.
26/04/2024	Renewed following Simplified Review. Amendment to Company Address.

Conditions attached to the issue of this certificate:

1. This certificate alone, if Category II (two) PPE, forms INSPECs permission to the manufacturer to use the 'CE' conformity mark for compliant products to be placed on the European Union internal market. In this case, the manufacturer may affix the conformity mark to each PPE or on a document supplied with the PPE, and draw up a written EU declaration of conformity for each PPE model referencing this certificate as per Article 17.

2. 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.

3. The manufacturer / authorised representative shall inform INSPEC without delay of any planned changes to the product, technical file or manufacturer information which may affect the validity of this certificate, before any such change is made.

4. 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.

5. 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 European Union internal market.

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 This certificate remains the property of INSPEC and may be withdrawn if any of the conditions attached to its issue are not complied with.

7. 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.

8. The manufacturer shall not use its product certification in any manner as to bring INSPEC into disrepute, nor make any statements regarding the product certification that INSPEC considers misleading or unauthorised.

9. Upon suspension, withdrawal or termination of this certificate, the manufacturer must discontinue all advertising matter that references the product certification and take action to cease production of products.