

## PRODUCTION CONFORMITY TO TYPE CERTIFICATE

This is to certify that INSPEC International B.V., Notified Body 2849, has assessed the quality system of the manufacturer and deemed it to be in compliance with Annex VIII (Module D) of the Personal Protective Equipment Regulation (EU) 2016/425.

Manufacturer: **Shanghai Dasheng Health Products Manufacture Co. Ltd**  
No. 228 Shihui Road  
Songjiang  
Shanghai  
201613  
China

The scope of the certification is for:

**The manufacture of respiratory protective devices.**

as per the Product Type(s) and Site(s) listed on page 2.

This certificate, while valid, serves as authorisation to the manufacturer from INSPEC International B.V. to affix our notified body identification number, 2849, to each individual item of PPE that is in conformity with the type described in the type-examination certificates.

Date of initial certification: 26 August 2019  
Date of current issue: 26 March 2021  
Date of expiry: 31 March 2024



Certification Manager

## Product Types

The PPE produced under this quality system was designed taking into consideration the following standard(s) / technical specification(s):

Standard	Product type
EN149	Filtering half masks to protect against particles

The Type-examination certificate(s) number(s) for the PPE subject to our assessment of the quality system are detailed within internal INSPEC International B.V. documentation.

## Sites

The quality system incorporates the following premise(s):

Site	Name / Physical Address
Primary office & Production location:	As per certificate holder address.

**Certificate amendment record**

Date	Description
26/08/2019	Initial Issue
26/03/2021	Re-issued following re-assessment. Previous version was issued by NB0194.

**Conditions attached to the issue of this certificate**

1. The manufacturer / authorised representative shall undertake to fulfil the obligations arising out of the quality system, and of the Personal Protective Equipment Regulation (EU) 2016/425, Annex VIII, and with INSPEC's Regulations governing this Module.
2. The manufacturer / authorised representative shall inform INSPEC of any planned changes to the quality system, wherein INSPEC will proceed with evaluation of the proposals as per Annex VIII, 3.5.
3. For the certificate to remain valid audits and visits must be conducted, as per Annex VIII, 4.
4. The manufacturer may affix INSPEC's notified body identification number, 2849, to each PPE, and draw up a written EU declaration of conformity for each PPE model referencing this certificate and the Type-examination certificate(s) as per Annex VIII, 5.
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.