

CE

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Verify the validity with the QR code



EU TYPE EXAMINATION CERTIFICATE

Certificate No: 2163-PPE-901

Respiratory protective devices, filtering half masks to protect against particles manufactured by
Fuzhou Chunlan Medical Equipment Co., Ltd.
 SF08-55, Fusheng Building, No. 3 Fuxing Avenue, Gulshan Town, Jixian District, Fuzhou City,
 Fujian Province, China
 are tested and evaluated according to:

**EN 149:2001 + A1:2009 Respiratory Protective Devices -
 Filtering Half Masks to Protect Against Particles -
 Requirements, Testing, Marking**

Based on the type examination conducted with the evaluation of test reports, technical file
 according to Personal Protective Equipment Regulation (EU) 2016/425 Annex 5, it is approved
 that the product meets the requirements of the regulation.

Product Definition

Model: CL-P1
 Filtering half mask
 Classification: FFP2 NR

Here by the manufacturer is allowed to use notified body number (2163) and can fix CE mark, as
 shown below, on the Category III product models given above, with:

- Issuing an appropriate EU Declaration of Conformity according to Personal Protective
 Equipment Regulation (EU) 2016/425 Annex 9.
- Ongoing successful performance in fulfilment of the requirements set out in Personal
 Protective Equipment Regulation (EU) 2016/425 and harmonised standards, entered
 by assessments based on Annex 7 (Module C2) or Annex 8 (Module D) of the
 regulation no later than 1 year from the beginning of serial production.

This certificate is initially issued on 30/06/2020 and will be valid for 5 years, if there is no
 change in the relevant harmonised standard affecting the essential health and safety
 requirements.



Serhat KACMAZ
 UNIVERSAL CERTIFICATION
 Director

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CERTIFICATE OF CONFORMANCE

Certificate No: 2163-PPE-901/01

Respiratory protective devices, filtering half masks to protect against particles manufactured by
Fuzhou Chunlan Medical Equipment Co., Ltd.
 SF08-55, Fusheng Building, No. 3 Fuxing Avenue, Gulshan Town, Jixian District, Fuzhou City,
 Fujian Province, China
 Continue to fulfil the requirements of

**EN 149:2001 + A1:2009 Respiratory Protective Devices -
 Filtering Half Masks to Protect Against Particles -
 Requirements, Testing, Marking**

Based on the evaluation of test reports and internal quality control audit reports according to EN
 149+A1:2009 and Personal Protective Equipment Regulation (EU) 2016/425 Annex VII
 (Module C2). This certificate implies that the manufactured products shown below are in
 conformance with the approved EU Type Examination model and meets the requirements of the
 regulation.

Product Definition

Model	Class	EU Type Examination Certificate		
		Serial No	Date	Issuing NB No
CL-P1	FFP2 NR	2163-PPE-901	30.06.2020	2163

Here by the manufacturer is allowed to use notified body number (2163) and can fix CE mark, as
 shown below, on the Category III product models given above, with:

- Issuing an appropriate EU Declaration of Conformity according to Personal Protective
 Equipment Regulation (EU) 2016/425 Annex 9.
- Taking all measures necessary so that the manufacturing process and its monitoring
 ensure the homogeneity of production and conformity of the manufactured PPE with the
 type described in the EU type examination certificate.

This certificate is issued on 30/06/2020 and will be valid for one year, until 29/06/2021 if the
 manufacturer makes no major change in the product designs and manufacturing processes
 affecting the product performance on the essential health and safety requirements.

Serhat KACMAZ
 UNIVERSAL CERTIFICATION
 Director