

# EU Type Examination Certificate

This is to certify that:

ZHEJIANG DONGMENG MEDICAL EQUIPMENT  
CO., LTD.  
Dongmeng Industrial Area Wuniu  
Yongjia, Wenzhou  
Zhejiang  
325000  
China

Holds Certificate Number:

CE 729864

In respect of:

**Model EYTV139110051-02 Particulate Respirator.**  
**To technical specification Annex II (EHSR) of the PPE Regulation (EU) 2016/425.**  
**PPE for use by healthcare professionals as per Commission recommendation 2020/403.**

on the basis that BSI carried out the relevant Type Examination procedures under the requirements with the Regulation (EU) 2016/425 of the European Parliament and Council relating to Personal Protective Equipment Regulation (PPE) Annex V (Module B) and meets the relevant health and safety requirements specified in Annex II

For and on behalf of BSI, a Notified  
Body for the above Regulation  
(Notified Body Number 2797):

Previous Notified Body: BSI 0086

First Issued: 2020-06-30

Latest Issue: 2020-06-30

Drs. Dave Hagenaaars, Managing Director

Effective Date: 2020-06-30

Expiry Date: 2021-06-30

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No. CE 729864

## Product Specification

**Product Name:** Particulate Respirators.

**Product Type:** Filtering half masks for use by Healthcare professionals.

**Model:** **EYTV139110051-02**

**Technical Specification:** Technical specification to satisfy Annex II of the PPE Regulation (EU) 2016/425.

**Product Description:** The respirator is non-reusable, secured to the face of the user by a pair of elasticated straps, and has no exhalation valve. The respirator is approved to GB 2626-2006 as KN95 product.

The respirator listed on this certificate is for use by healthcare workers, first responders and other personnel involved in the efforts to contain the COVID-19 virus and avoid its further spread.

The product covered by this certificate is not approved for industrial applications and the certificate is only valid as long as EU Commission recommendation sheet 2020/403 remains applicable.

**Product Assessments:** GB 2626-2006 Standard KN95

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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  
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## Certificate Administration Details

Technical File Reference: Technical File -dongmeng-EYTV139110051-02

## Certificate Amendment Record:

Issue date	Comments	BSI Review No.
June 2020	First issue.	2797:20:3218883

## 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 product, failure to do so could invalidate the Certificate in respect of product manufactured following the introduction of such changes.

The validity of the Certificate for the products is also dependent on the maintenance of the EU Conformity to Type based on Internal Production Control plus supervised product checks at random intervals, Annex VII (Module C2) as referenced on BSI issued Certificate CE 729865.

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