

Conformity to Type based on Quality Assurance of the Production Process

This is to certify that:

MFA Is Guvenligi Medikal A.S.
Elvanpazarcik Beldesi Hayat Mah. Baruthane Cad. No:21/1
Merkez
Zonguldak
67990
Turkey

Holds Certificate Number:

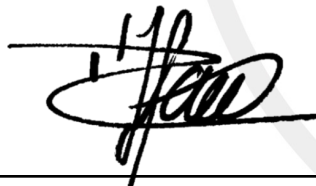
CE 758381

In respect of:

For the manufacture of RPE to the standards listed on the continuation sheet.

on the basis that BSI carried out the quality assurance assessment under the requirements with the Regulation (EU) 2016/425 of the European Parliament and Council relating to Personal Protective Equipment Regulation (PPE) Annex VIII (Module D)

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



Drs. Dave Hagenaaers, Managing Director

First Issued: 2022-01-11

Latest Issue: 2022-01-11

Effective Date: 2022-01-11

Expiry Date: 2027-01-11

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No. CE 758381

PPE Module D Certificate holding location:

MFA Is Guvenligi Medikal A.S.
Elvanpazarcik Beldesi Hayat Mah. Baruthane
Cad. No:21/1 Merkez
Zonguldak
67100
Turkey

Manufacturing Site:

MFA Is Guvenligi Medikal A.S.
Elvanpazarcik Beldesi Hayat Mah. Baruthane
Cad. No:21/1 Merkez
Zonguldak
67100
Turkey

Product Specification

The products are covered by the scope of this Module D certificate, conform to the following standards:

Standard

EN 136:1998/AC:2003
EN 140:1998/AC:1999
EN 143:2000/A1:2006
EN 149:2001+A1:2009
BS EN 14387:2004+A1:2008

Product Type

Respiratory protective devices – Full face masks.
Respiratory protective devices – Half masks and quarter masks.
Respiratory protective devices – Particle filters.
Respiratory protective devices – Filtering half masks to protect against particles.
Respiratory protective devices – Gas filter(s) and combined filter(s).

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A member of BSI Group of Companies.

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No. CE 758381

Certificate Amendment Record

| Issue date | Comments | BSI Project Number |
|--------------|--------------|--------------------|
| January 2022 | First issue. | 2797:21:3541518 |

Note: The Certificate Holder is responsible for keeping the Notified Body advised of changes to any aspect of the overall process in the manufacture of the product.



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