

Conformity to Type based on Quality Assurance of the Production Process

This is to certify that:

MFA Is Guvenligi Medikal
Tekstil Imalat Tic. San. ltd. Sti
Elvanpazarcik Beldesi
Hayat Mah. Baruthane
Cad. No:21/1 Merkez
Zonguldak
67100
Turkey

Holds Certificate Number:

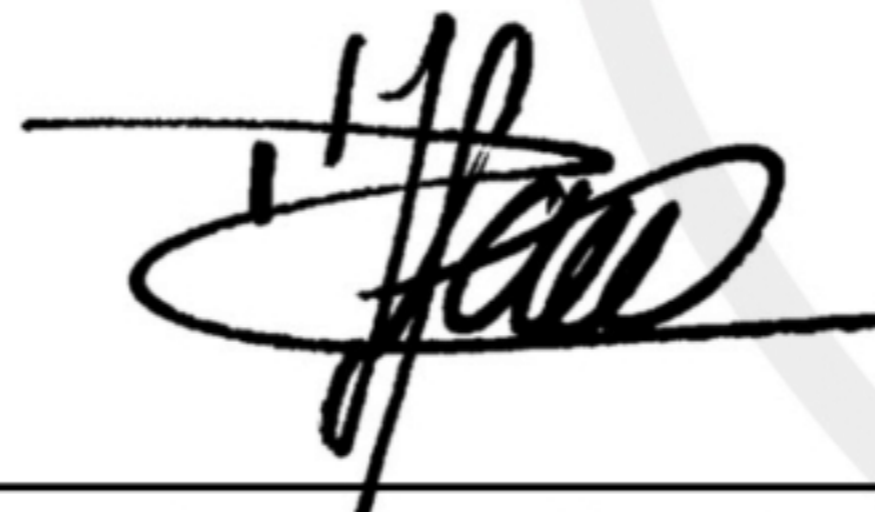
CE 688435

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: 2018-05-11

Latest Issue: 2020-10-28

Effective Date: 2020-10-28

Expiry Date: 2023-05-11

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No. CE 688435

PPE Module D Certificate holding location:

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Elvanpazarcik Beldesi Hayat Mah. Baruthane
Cad. No:21/1 Merkez
Zonguldak
67100
Turkey

Manufacturing Site:

MFA Is Guvenligi Medikal
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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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Certificate Amendment Record

Issue date	Comments	BSI Project Number
October 2020	Change of certification and manufacturing location	2797:20:3289267
September 2019	Addition of standards EN 136:1998+AC: 1999, BS EN 140:1999, BS EN 143:2000 & BS EN 14387:2004+A1:2008	0086:19:9783670
April 2018	First Issue	0086:18:8889024

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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