

Conformity to Type based on Internal Production Control plus supervised product checks at random intervals

This is to certify that:

Zagor Is Gv. Ekip. San. Tic. Ltd. Sti
Fevzi Cakmak Mahallesi
Mimar Sinan Caddesi
No:40 Dembeller Plaza
Istanbul, Pendik
34899
Turkey

Holds Certificate Number:

CE 732024

In respect of:

For the manufacture of particulate respirators to technical specification to satisfy Annex II of the PPE Regulation (EU) 2016/425.

on the basis that BSI carried out the supervised production checks at random intervals under the requirements with the Regulation (EU) 2016/425 of the European Parliament and Council relating to Personal Protective Equipment Regulation (PPE) Annex VII (Module C2)

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

Drs. Dave Hagenaaers, Managing Director

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Expiry Date: 2021-11-12

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Product manufactured by:

ZAGOR Is Guvenligi Ekipmanlari San. Tic. Ltd. Sti.
Fevzi Cakmak Mahallesi, Mimar Sinan Caddesi, No:40 Dembeller Plaza
Istanbul
Pendik
34899
Turkey

Product details

The respiratory protective device covered by the scope of this Module C2 Certificate and the Technical Specification to which the product is manufactured are as follows:

Product type:	Particulate filtering half masks for use by Healthcare professionals.
Model and classifications:	ZGR 5030 FFP3 V NR and ZGR 5035 FFP3 NR
Technical Specification:	Technical specification to satisfy Annex II of the PPE Regulation (EU) 2016/425. BSI's PPE for Healthcare Professionals 2020/403 – RPE Technical Specification.

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A member of BSI Group of Companies.

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Certificate Administration Details:

Certificate Amendment Record and BSI internal Review relating to this Certificate

Issue date	Comments	BSI Review No.
November 2020	First issue	2797:20:3253603

Certificate validity

The Certificate holder is responsible for ensuring that the Notified Body is advised of changes to any aspects of the overall quality system utilized in the manufacture of the products, failure to do so could invalidate the Certificate in respect of product manufactured after the introduction of such changes.

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