

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

This is to certify that:

Unimama B.V.
Sloterweg 315
Badhoevedorp
1171 VC
The Netherlands

Holds Certificate Number:

CE 729821

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):

Previous Notified Body: BSI 0086

First Issued: 2020-07-08

Latest Issue: 2020-07-08

Drs. Dave Hagenaaars, Managing Director

Effective Date: 2020-07-08

Expiry Date: 2021-07-08

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No. CE 729821

Product manufactured by:

Guangzhou Carrot Mall Network Technologies Co., Ltd
Room 1406, No.4 Zhudian Road (Nansha Street),
Nansha District,
Guangzhou,
Guangdong Province,
China

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:	IRYS-01 FFP2 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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Certificate Administration Details:

Certificate Amendment Record and BSI internal Review relating to this Certificate

Issue date	Comments	BSI Review No.
July 2020	First issue.	2797:20:3218335

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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