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

This is to certify that:

Guangxi Anran Medical equipment Co., Ltd
Room 3-1,No.3,Workshop
No.1,yinxiang Road
Liunan District ,Liuzhou City
Guangxi
545000
China

Holds Certificate Number: CE 730098

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 Hagenaars, Managing Director

Previous Notified Body: BSI 0086

First Issued: 2020-07-09

Effective Date: 2020-07-09

Latest Issue: 2020-07-09

Expiry Date: 2021-07-09

PRODUCTS RvA C 646

...making excellence a habit.™

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No. CE 730098

Product manufactured by:

Guangxi Anran Medical equipment Co., Ltd.
Room 3-1, No.3, Workshop
No.1, Yinxiang Road
Liunan District
Liuzhou City
Guangxi,
545000
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:	ANRAN9501A 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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No. CE 730098

Certificate Administration Details:

Certificate Amendment Record and BSI internal Review relating to this Certificate

Issue date	Comments	BSI Review No.
July 2020	First issue.	2797:20:3219728

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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