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

This is to certify that:

Dishang Medical Technology Co,. Ltd
Room 406-409, Block C
Innovation and Entrepreneurship Base
Huoju Rd.(No.213)
Torch Hi-tech Ind. Development Zone
Weihai City
Shandong
China

Holds Certificate Number: CE 727973

In respect of:

Medical protective clothing to Annex II of Regulation 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

For and on behalf of BSI, a Notified Body for the above Regulation (Notified Body Number 0086):

Chris Lewis - Certification Director, Product Certification

First Issued: 2020-05-08
Latest Issue: 2020-05-08

Effective Date: 2020-05-08
Expiry Date: 2021-05-08

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No. CE 727973

Product manufactured by:

Dishang Medical Technology Co Ltd
Room 406-409, Block C
Innovation and Entrepreneurship Base
Huoju Rd. (No.213)
Torch Hi-tech Industrial Development Zone
Weihai City
Shandong Province
China

Product details

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

Product type:	Protective Coverall for Protection against Infective Agents
Model and classifications:	Models covered in this recommendation: AY160 to 200 AW160 to 200 BY160 to 200 BW160 to 200
Technical Specification:	Technical specification to satisfy Annex II of the PPE Regulation (EU) 2016/425. BSI's PPE for Healthcare Professionals 2020/403 – Medical Clothing 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.
May 2020	First issue.	0086:20:3175488

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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