

EC Certificate

Directive 93/42/EEC Annex II, excluding Section 4 Full Quality Assurance System Medical Devices

Registration No.: HD 60147078 0001

Report No.: 17054672 002

Manufacturer: BrosMed Medical Co., Ltd.

15th Building, SMEs Venture Park Songshan Lake, Hi-Tech Industrial

Development Zone

523808 Dongguan, Guangdong

P.R. China

Products: Medical Devices

(see attachment for products included)

Expiry Date: 2024-05-26

The Notified Body hereby declares that the requirements of Annex II, excluding section 4 of the directive 93/42/EEC have been met for the listed products. The above named manufacturer has established and applies a quality assurance system, which is subject to periodic surveillance, defined by Annex II, section 5 of the aforementioned directive. For placing on the market of class III devices covered by this certificate an EC design-examination certificate according to Annex II, section 4 is required.

Effective Date:

2020-04-24

Date:

2020-04-24

TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg
TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 93/42/EEC
concerning medical devices with the identification number 0197.



TÜV Rheinland LGA Products GmbH Tillystraße 2, 90431 Nürnberg Doc. 1/1, Rev. 0

Attachment to Certificate

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BrosMed Medical Co., Ltd.

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Products:

- -Introducer Sheath Sets
- -Connecting Tubings
- -Angiographic Syringes and Accessories
- -Introducer Needles
- -Non-vascular Guide Wires

Aspects of manufacture concerned with securing and maintaining sterile conditions:

- -Balloon Inflation Devices
- -Radial Artery Tourniquets
- -Y Connector Sets
- -Dose-control Syringes
- -Manifolds

Date: 2020-04-24

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