TÜVRheinland®

EC Certificate

Directive 93/42/EEC Annex V
Production Quality Assurance
Medical Devices

Registration No.: DD 60144229 0001

Report No.: 15076053 008

Manufacturer:
Zhejiang Bangli Medical
Products Co., Ltd.
No. 118 Yuegui South Road
City West New District
321300 Yongkang City, Zhejiang Province
China

Products:
Aspects of manufacture concerned with securing and maintaining sterile conditions of Wound Plasters, Wound Dressings, First Aid Kits

Replaces Approval, Registration No.: DD 60097501 0001

Expiry Date: 2024-05-27

The Notified Body hereby declares that the requirements of Annex V 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 V, section 4 of the aforementioned directive. For placing on the market of class IIb and class III devices covered by this certificate an EC type-examination certificate according to Annex III is required.

Effective Date: 2019-11-14

Date: 2019-11-14

Notified Body

TÜV Rheinland LGA Products GmbH Zertifizierungsstelle III TÜVRheinland

Fuxiu Sheng

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.

Digitally signed by Guțan Ghenadie
Date: 2019.12.13 12:36:23 EET
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