

EC Certificate



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

Registration No.: DD 2110099-1

Manufacturer: Nantong Jianan Medical Products
Co., Ltd.
2nd Floor, 7th Building,
Electronic Industry Park,
No.9, Xindong Road, Nantong Economic and
Technological Development Zone,
226010 Jiangsu
P.R. China

Products: Lap Sponges;
Aspects of manufacture concerned with securing and maintaining sterile
conditions: Sterile Gauze Sponges



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.

Report No.: 15063749 011
Effective date: 2021-03-29
Expiry date: 2023-11-03
Issue date: 2021-03-29



Fuxiu Sheng
TÜV Rheinland LGA Products GmbH
Tillystraße 2 · 90431 Nürnberg · Germany

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

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The scope of certification includes the following manufacturing sites:

No.	Location	Product groups manufactured
/01	Nantong Jianan Medical Products Co., Ltd. No.55, Fuxing Road, Nantong Economic And Technological Development Zone, 226009 Jiangsu P.R. China	Lap Sponges; Aspects of manufacture concerned with securing and maintaining sterile conditions: Sterile Gauze Sponges

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