

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

Registration No.: DD 60134639 0001

Report No.: 15096311 002

Manufacturer: Huaian Helen Medical Instrument

Co., Ltd.

Group 3, Zhengtai Village Matou Town, Huaiyin District

Huaian

223345 Jiangsu

China

Products: - Sterile Blood Lancets

- Disposable Surgical Blades & Scalpels with Plastic Handle

Expiry Date: 2023-09-28

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: 2018-12-21

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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.