



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

Registration No.: DD 60128426 0001

Report No.: 15096250 002

Manufacturer: Yancheng Huida Medical Instruments
Co., Ltd.
No. 2, Kuiyuan Road
Xuefu Town, Yandu District
Yancheng City
224035 Jiangsu
China

Products:

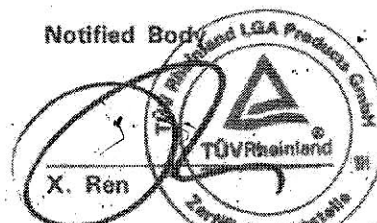
- Sterile Blood Lancets
- Disposable Surgical Blades (with and without handle)

Expiry Date: 2023-05-09

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: 2018-05-10

Date: 2018-05-10



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.

