EC Certificate

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

Registration No.: DD 60152718 0001

Report No.: 15079360 011

Manufacturer:

Chengdu Rich Science
Industry Co., Ltd.
No. 66, Tianqindong Street, West
Hi-tech Development Zone
Chengdu
611731 Sichuan
P.R. China

Products:

Single Use Blood Collection Needles;
Aspects of manufacture concerned with securing and maintaining sterile conditions:
Single Use Specimen Collection Swabs

Replaces Approval, Registration No.: DD 60145132 0001

Expiry Date: 2024-05-26

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: 2021-03-18

Date: 2021-03-18

Notified Body

Fuxiu Sheng

TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg

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