EC Certificate

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

Registration No.: DD 60114908 0001

Report No.: 15096001 001

Manufacturer:
Hangzhou Qiandao Medical
Equipment Co., Ltd.
No.8, Xiatang Road, Chengdong
Village, Yangxi Street
311607 Jiande City, Zhejiang
China

Products: Disposable Liposuction Cannula

Replaces Approval, Registration No.: DD 60040180 0001

Expiry Date: 2021-08-25

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.

Notified Body

Effective Date: 2016-12-02

Date: 2016-12-02

X. Ren

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.