EC Certificate

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

Registration No.: DD 60132095 0001

Report No.: 50170533 002

Manufacturer:
Xi'an Haiye Medical Equipment
Co., Ltd.
Feng Jing Industrial Park
Xi'an city
710300 Shaanxi
P.R. China

Products:
Video Laryngoscopes;
Aspects of manufacture concerned with securing and maintaining sterile conditions:
Disposable Video Laryngoscopes Blades

Expiry Date: 2023-11-09

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: 2019-08-29

Date: 2019-08-29

Notified Body

Wenxiang Zhang

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.