



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

Registration No.: DD 60154538 0001

Report No.: 15096114 006

Manufacturer: Shanghai Bojin Medical Instrument
Co., Ltd.
A Zone of F2, C Zone of F1
Building 6
No. 125, Longpan Road, Jiading District
201801 Shanghai
P.R. China

Products:

- Medical Saw Blades
- Medical Drill Bits
- Arthroscopic Surgery Blades
- Medical Electrical Saw Drills

Replaces Approval, Registration No.: DD 60117721 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: 2021-04-14

Date: 2021-04-14

Notified Body

Jason Pan



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.