

EC Certificate

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

Registration No.: DD 2056447-1

Manufacturer: Foshan SOCO Precision Instrument

Co., Ltd.

2FI. Bldg 3, District A, Guangdong New Light Source Industrial Base Luocun, Shishan Town, Nanhai District

Foshan City

528226 Guangdong

P.R. China

Products: High-speed Air Turbine Handpieces, Dental Low Speed Handpieces

including Straight and Geared Angle Handpieces and Air Motors, Dental

Root Canal Instruments

Replaces Approval, Registration No.: DD 60144326 001

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.

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TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 93/42/EEC concerning medical devices with the identification number 0197.