

FC Certificate

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

Registration No.: DD 60109667 0001

Report No.: 16805222 001

Manufacturer: Shenzhen Dian Fong Abrasives

Dental Burs

Company Limited West 4th Floor, Block 2 No. 5, TianHua Rd.

PingHu, LongGang District Shenzhen

518111 Guangdong China

(see attachment for additional site included)

Expiry Date: 2021-05-22

Products:

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 III band class III devices covered by this certificate an EC type-examination certificate are according to Annex III is required.

Effective Date: 2016-06-24

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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.