EC 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
Company Limited
West 4th Floor, Block 2
No. 5, TianHua Rd.
PingHu, LongGang District
Shenzhen
518111 Guangdong
China

Products: Dental Burs

(see attachment for additional site included)

Expiry Date: 2021-05-22

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: 2016-06-24

Date: 2016-06-24

Notified Body

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.

Digitally signed by Jighili Tatiana
Date: 2023.05.24 16:28:04 EEST
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