EC Certificate

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

Registration No.: DD 60128002 0001

Report No.: 17037638 006

Manufacturer:

BDC Dental Corporation Ltd.
17th, Xiangshan Road, Science City
Guangzhou
510663 Guangdong
China

Products:

Dental Air Turbine Handpieces

Replaces Approval, Registration No.: DD 60084076 0001

Expiry Date: 2019-03-16

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: 2018-05-10

Date: 2018-05-10

Notified Body

S. Liu

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.