EC Certificate

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

Registration No.: DD 60140547 0001

Report No.: 17037638 009

Manufacturer:

BDC Dental Corporation Ltd.
Part 3, No. 1 Guanchong Section, Shilian Rd., Shiqi Town
Panyu District
Guangzhou
511450 Guangdong
China

Products:

High-speed Air Turbine Handpieces, Straight Handpieces, Angle Handpieces and Air Motors

Replaces Approval, Registration No.: DD 60128002 0001

Expiry Date: 2024-03-16

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: 2019-07-18

Date: 2019-07-18

Notified Body

Dipl.-Ing. I. Munkier

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.