

**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

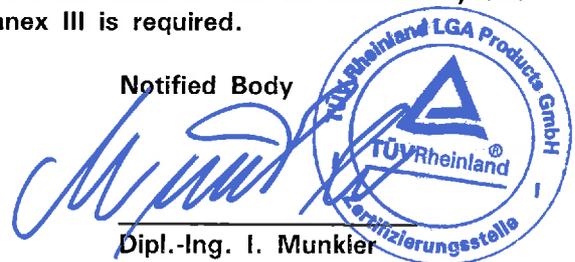
**Expiry Date:** 2024-03-16

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. Munkler

**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.