EC Certificate

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

TÜVRheinland®

Registration No.: DD 60130603 0001

Report No.: 17063017 002

Manufacturer:
Huizhou Videya Technology Co. Ltd.
Factory of Songshan Villagers Group,
Baishi Village,
Qiuchang Street Office, Huiyang District
Huizhou
516221 Guangdong
China

Products: Dental Root-canal Instruments

Expiry Date: 2023-07-23

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: 2018-09-13

Date: 2018-09-13

Notified Body

X. Ren

TÜV Rheinland LGA Products GmbH

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: 2020.11.30 15:51:57 EET
Reason: MoldSign Signature
Location: Moldova