

## **EC** Certificate

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

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

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.

**Notified Body**