# 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

Notified Body

X. Ren

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.

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