TÜVRheinland®

Certificate

The Certification Body of
TÜV Rheinland LGA Products GmbH

hereby certifies that the organization

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

has established and applies a quality management system for medical devices for the following scope:

Manufacture and Distribution of
Dental Root-canal Instruments

Proof has been furnished that the requirements specified in

EN ISO 13485:2016

are fulfilled. The quality management system is subject to yearly surveillance.

Effective Date:	2018-09-13
Certificate Registration No.:	SX 60130604 0001
An audit was performed. Report No.:	17063017 002
This Certificate is valid until:	2021-07-23

Certification Body

DAkkS
Deutsche
Akkreditierungsstelle
D-ZM-14169-01-02

Digitally signed by Perminov Andrei
Date: 2022.02.24 18:01:23 EET
Reason: MoldSign Signature
Location: Moldova

Date 2018-09-13

X. Ren

TÜV Rheinland LGA Products GmbH
TÜVRheinland®
Zertifizierungsstelle

TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg
Tel.: +49 221 806-1371 Fax: +49 221 806-3935 e-mail:cert-validity@de.tuv.com http://www.tuv.com/safety

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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Date 2018-09-13

TÜV Rheinland LGA Products GmbH
TÜVRheinland
Zertifizierungsstelle
X. Ren

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