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Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
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ZLG-BS-244.10.08

TÜV SÜD Product Service

EC Certificate

Full Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
(Devices in Class IIa, IIb or III)

No. G1 003076 0002 Rev. 02

Manufacturer: **Shinva Medical Instrument Co., Ltd.**
Xinhua Medical Scientific Zone
Zibo New & Hi-Tech Industrial Development Zone
255086 Zibo
PEOPLE'S REPUBLIC OF CHINA

Product Category(ies): **Autoclave, Rapid Automatic Washer-Disinfector and Dental Zirconia Ceramic.**

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See also notes overleaf.

Report No.: BJ19501033

Valid from: 2019-12-09
Valid until: 2023-03-18

Date, 2019-12-09

Christoph Dicks
Head of Certification/Notified Body

Digitally signed by Perminov Andrei
Date: 2022.06.22 12:40:30 EEST
Reason: MoldSign Signature
Location: Moldova

TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany

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Facility(ies):

Shinva Medical Instrument Co., Ltd.
No. 99 Beixin Road, Zibo New & Hi-Tech Zone, 255086 Zibo, PEOPLE'S REPUBLIC OF CHINA

Shinva Medical Instrument Co., Ltd.
Xinhua Medical Scientific Zone, Zibo New & Hi-Tech Industrial Development Zone, 255086 Zibo, PEOPLE'S REPUBLIC OF CHINA

Shinva Medical Instrument Co., Ltd.
No. 2009 Xinhua Ave, Zhoucun, 255300 Zibo, PEOPLE'S REPUBLIC OF CHINA

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