

## EC DECLARATION OF CONFORMITY

Name and address of the manufacturer: Foshan SOCO Precision Instrument Co.,Ltd.  
2FL.Bldg 3,District A Guangdong New Light Source Industrial  
Base,Luocun Shishan Town,Nanhai District Foshan City  
528226 Guangdong China

EC Representative: Lotus NL B.V.  
Koningin Julianaplein 10,1e Verd,2595AA,The Hague,  
Netherlands.

We declare under our sole responsibility that

the medical device: **Product Name:** Dental Root Canal Instruments  
**Model:** SC, SC-PRO, SC plus, S-one plus, S-one pro,  
SX-F3, SC-GOLD

of class: Ila, rule 6  
according to annex IX of directive 93/42/EEC

meets the provisions of the directive 93/42/EEC as amended by Directive 2007/47/EC and its transpositions in national laws which apply to it. The declaration is valid in connection with the "final inspection report" of the device.

Conformity assessment procedure: **Directive 93/42/EEC Annex V**

Registration No.: **DD 2056447-1**

Notified Body: TÜV Rheinland LGA Products GmbH  
Tillystraße 2  
90431 Nürnberg  
Deutschland  
CE 0197

(FoShan ), PR China 2021-05-25

Place, date

Title: General Manager

Name: (Mr) Zeng Yong

Signature: 

Name and function

