

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: IIa, 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

