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