

## 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.  
+31644168999

We declare under our sole responsibility that

the medical device: **Product Name:** High-speed Air Turbine Handpieces  
**Model:** SCHG01, SCHG03, SCHG18, SCHF22, SCHF19

of class: Ila, rule 9  
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.

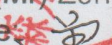
Conformity assessment procedure: **Directive 93/42/EEC Annex V**

Registration No.: **DD 60125716 0001**

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

(FoShan ), PR China 2021-7-12

Place, date

Title: General Manager  
Name: (Mr) Zeng Yong  
Signature: 

Name and function

