

## EC DECLARATION OF CONFORMITY

Name and address of the manufacturer: Foshan COXO Medical Instrument Co., Ltd.  
No. 17, Guangming Ave., New Light Source Industrial Base,  
Nanhai National High-tech Zone, Foshan 528226, Guangdong  
P.R. China

EC Representative: Lotus NL B.V.  
Koningin Julianaplein 10, 1e Verd, 2595AA, The Hague,  
Netherlands.  
E-mail: peter@lotusnl.com

Trademarks: **COXO, YUSENMENT, YSDENT, CODENTAL**

We declare under our sole responsibility that

the medical device: **Product Name:** High-speed Air Turbine Handpieces  
**Model:** CX207, CX207-G, CX207-2, CX207-A, CX207-A-2,  
CX207-B, CX207-B-2, CX207-C, CX207-C-2,  
CX207-F, CX207-W, CX207-W-2

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 60151346 0001**

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

(FoShan ), PR China 2020-12-08

Place, date

Title: General Manager  
Name: (Mr) Zheng Yongliang  
Signature: 

Name and function

