

EC DECLARATION OF CONFORMITY

Name and address of the manufacturer: Foshan COXO Medical Instrument Co., Ltd.
BLDG 4, District A Guangdong New Light Source Industrial
Base, South of Luocun Avenue Nanhai District Foshan 528226
Guangdong 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

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

Place, date

Name and function