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, YUSENDENT, 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:	IIa, 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