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:	Wellkang Ltd. The Black Church, St. Mary's Place, Dublin 7, D07 P4AX, Ireland
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 II, excluding Section 4**
Registration No.:	**HD 60109212 0001**
Notified Body:	**TÜV Rheinland LGA Products GmbH** **Tillystraße 2** **90431 Nürnberg** **Deutschland** **CE 0197**

(FoShan), PR China 2020-2-19

Place, date

Title: General Manager

Name: (Mr) Zheng Yongliang

Signature:

Name and function