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 **SRN:** CN-MF-000001682
EC Representative:	Lotus NL B.V. Koningin Julianaplein 10, 1e Verd, 2595AA, The Hague, Netherlands. E-mail: peter@lotusnl.com **SRN:**NL-AR-000000121

Trademarks: **COXO, YSDENT**

We declare under our sole responsibility that

the medical device:	**Product Name:** Dental ultrasonic surgical Device **Model:** C-EXPLORER **Basic UDI-DI:** 697426789352C **GMDN:** 47010 **MDA Code:** 0311 Active non-implantable dental devices **Intended Use:** This product is intended only for use in the field of dentistry, for surgery to expose and dissect oral tissue structures or endodontic treatments (e.g. periodontal， gap, gingiva, bone, jaw, extractions, implantations)
of class:	IIa, according to Rule 9, AnnexVIII, Regulation (EU) 2017/745

meets the provisions of Medical Device Regulation (EU) 2017/745. The declaration is valid in connection with the "final inspection report" of the device.

Conformity assessment procedure:	Annex IX (Full QMS), Regulation (EU) 2017/745
Registration No.:	**HZ 2182788-1**
Notified Body:	**TÜV Rheinland LGA Products GmbH** **Tillystraße 2** **90431 Nürnberg** **Deutschland** **CE 0197**

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

(FoShan), PR China 2024-10-12

Place, date

Name and function