

EC Declaration of Conformity

according to the Regulation (EU) 2017/745

Manufacturer:

Foshan COXO Medical Instrument Co., Ltd.

Site 1: BLDG 4, District A Guangdong New Light Source Industrial Base,
South of Luocun Avenue Nanhai District Foshan 528226 Guangdong
China

Site 2: No. 17, Guangming Ave., New Light Source Industrial Base,
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Tel: +86 (757) 66692058 Fax: +86 (757) 81800058

Web: www.coxotec.com Email: coxotec@163.com

SRN: CN-MF-000001682

Trademarks: COXO, YUSENDET, YSDENT, CODENTAL

EC Representative:

Lotus NL B.V. located

Koningin Julianaplein 10, 1e Verd, 2595AA, The Hague, Netherlands.

E-mail: peter@lotusnl.com

We declare under our sole responsibility that

**the medical
device(s)**

Product Name: Teeth Whitening Accelerator

Model	C-Bright	C-Bright Ia
Basic UDI-DI	697426789060PQ	697426789061PS

of class

Class I, rule 13

according to annex VIII of Regulation (EU) 2017/745

(non-sterile, non-measurement, non-reusable surgical)

meets the provisions of the Regulation (EU) 2017/745

The products comply
with requirements of
relevant harmonized
standards:

EN 60601-1:2006+ A11:2011+A1:2013+A12:2014	EN 60601-1-2:2015
EN ISO 14971:2012	EN ISO 15223-1:2016
EN 1041:2008+A1:2013	

Conformity assessment route:

Annex II+ III

Notified Body:

N/A

Signed this Day/ 25 of Month/ 05 of Year/ 2021 , Place (FoShan), PR China

Represented by:

Title: General Manager

Name: (Mr) Zheng Yongfang

Signature:

Official Seal:

