

# EC Declaration of Conformity

according to the Regulation (EU) 2017/745

**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

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SRN: CN-MF-000001682

**Trademarks:** COXO, YUSENDEMENT, YSDENT, CODENTAL

**EC Representative:**

**Lotus NL B.V. located**

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

E-mail: [peter@lotusnl.com](mailto:peter@lotusnl.com)

**We declare under our sole responsibility that**

**the medical  
device(s)**

Product Name: Air Abrasion System with spray

Model: CA-1

Basic UDI-DI: no, applying

**of class**

Class I , rule 1

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 ISO 13485:2016

EN ISO 14971:2019

EN ISO 20608:2018

ISO 15223-1:2021

EN 1041:2008+A1:2013

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Conformity assessment route:

Annex II+ III

Notified Body:

N/A

Signed this Day/ 25 of Month/ 10 of Year/ 2021 , Place (FoShan), PR China

**Represented by:**

Title: General Manager

Name: (Mr) Zheng Yongliang

Signature:

Official Seal:

