



TÜVRheinland®

## EC Certificate

Directive 93/42/EEC Annex II, excluding Section 4  
Full Quality Assurance System  
Medical Devices

Registration No.: HD 60109212 0001

Report No.: 17050363 001

**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  
**Products:** Medical Devices

(see attachment for products included)

**Expiry Date:** 2020-12-08

The Notified Body hereby declares that the requirements of Annex II, excluding section 4 of the directive 93/42/EEC have been met for the listed products. The above named manufacturer has established and applies a quality assurance system, which is subject to periodic surveillance, defined by Annex II, section 5 of the aforementioned directive. For placing on the market of class III devices covered by this certificate an EC design-examination certificate according to Annex II, section 4 is required.

**Effective Date:** 2016-05-27

**Date:** 2016-05-27



**TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg**

TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 93/42/EEC concerning medical devices with the identification number 0197.



Doc. 1/1, Rev. 1

**TÜV Rheinland**  
**LGA Products GmbH**  
**Tillystraße 2, 90431 Nürnberg**

**Attachment to  
Certificate**

**Registration No.:** HD 60109212 0001  
**Report No.:** 17050363 002

**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

**Products:**

- Root Apex Locators
- Endo Motors
- Pulp Testers
- High-speed Air Turbine Handpieces
- Dental low speed handpieces including straight and geared angle handpieces and airmotors
- Dental Implant Systems
- Dental Electrical Motors

**Date:** 2017-05-12



S. Liu