

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, Langsha Luocun Shishan Town,
Nanhai District, Foshan City, Guangdong Province,
China.

Trademarks: **COXO, YUSENMENT, YSDENT, CODENTAL**

We declare under our sole responsibility that

the medical device: **Product Name: Root Apex Locators**
Model : C-Root I, C-Root I(III), C-Root I(V), C-Root I(VI), C-Root i+

of class: **Ila , rule 10**

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 2017-05-12

Place, date

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

Name and function

