

EC DECLARATION OF CONFORMITY

Name and address of the manufacturer: **BLDG 4, District A, Guangdong New Light Source Industrial Base, South of Luocun Avenue, Nanhai District, Foshan, 528226 Guangdong, China**

We declare under our sole responsibility that

the medical device: **Product Name: High-speed air turbine handpieces
Model : CX207**

of class: **Ila , rule 9**

according to annex IX of directive 93/42/EEC

meets the provisions of the directive 93/42/EEC 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.: **HD601092120001**

Notified Body: **TÜV Rheinland LGA Products GmbH
Tillystraße 2
90431 Nürnberg
Deutschland
CE 0197**

(FoShan), PR China 2016-05-27

Place, date

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

Name and function

