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MOLDOVA EUROPEANĂ

EC DECLARATION OF CONFORMITY

The following EC declaration of conformity exemplifies the required content according to Directive 93/42/EEC.

EC DECLARATION OF CONFORMITY

Name and address of the manufacturer:	Tealth Foshan Medical Equipment Co.,Ltd. No.4, Qiling Road, Lutang Industrial Zone, Luocun, Shishan Town, Nanhai District, Foshan, Guangdong 528226, China
Name and address of Authorized Representative:	Shanghai International Holding Corp. GmbH (Europe) Eiffestrasse 80, 20537 Hamburg，Germany

We declare under our sole responsibility that

the medical device: Dental High speed air turbine handpiece

Model of high-speed air turbine handpieces: 2300PQL-M4, 2300PL-M4, 2300P-B2, 2300PL-B2, 2300P-45, 2300P-45B, 2300PL-45, 2300PQL-45, 2301PL-M4, 2301PL-B2, 2302PL-B2, 2302PQL-M4, 2302PQL-B2, 2303PL-M4, 2303PL-B2, 2304P-M4, 2304P-B2, 2305P-M4, 2305P-B2.

of class: IIa 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.: **CN19/41107**

Notified Body:

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Notified Body number: 1639

Li Xiuying Li Xiuying

Foshan, Guangdong — General Manager:

Place, date / 2021.9.16 — Name and function