# EC Certificate

TÜVRheinland®

**Production Quality Assurance**
**Directive 93/42/EEC on Medical Devices, Annex V**

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| Registration No.: | DD 2056447-1 |
| Manufacturer: | Foshan SOCO Precision Instrument Co., Ltd.<br>2FI. Bldg 3, District A, Guangdong New Light Source Industrial Base<br>Luocun, Shishan Town, Nanhai District<br>Foshan City<br>528226 Guangdong<br>P.R. China |
| Products: | High-speed Air Turbine Handpieces, Dental Low Speed Handpieces including Straight and Geared Angle Handpieces and Air Motors, Dental Root Canal Instruments<br><br>Replaces Approval, Registration No.: DD 60144326 001 |

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Date: 2023.08.23 15:50:31 EEST
Reason: MoldSign Signature
Location: Moldova

The Notified Body hereby declares that the requirements of Annex V 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 V, section 4 of the aforementioned directive. For placing on the market of class IIb and class III devices covered by this certificate an EC type-examination certificate according to Annex III is required.

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| Report No.: | 10918784-100 |
| Effective date: | 2021-05-25 |
| Expiry date: | 2024-05-26 |
| Issue date: | 2021-05-25 |

TÜV Rheinland LGA Products GmbH Zertifizierungsstelle TÜVRheinland®

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