EC Certificate

Production Quality Assurance
MDD Annex V

Registration No.:	DD 2185275-1
Manufacturer:	Foshan Suntem Medical Instrument Co., Ltd. 1 Flat, No. 7, C District Sanshui Industry Park Foshan City 528137 Guangdong P.R. China
Products:	Dental Units, Ear, Nose & Throat Treatment Units
	Replaces Approval, Registration No.: DD 60115433 0001

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.

Report No.:	10917976-100
Effective date:	2021-01-21
Expiry date:	2024-05-26
Issue date:	2021-01-21

TÜV Rheinland LGA Products GmbH Zertifizierungsstelle

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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.

Digitally signed by Timus Andrei
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