Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
ZLG-BS-244.10.08

TÜV SÜD Product Service

EC Certificate

Production Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex V
(Devices in Class IIa, IIb or III)

No. G2 104331 0002 Rev. 00

Manufacturer:

FOSHAN AKOS MEDICAL INSTRUMENT CO., LTD
Room 301, 3F, Unit A, No. 4 Zone B
HAO SCIENCE PARK, Guicheng Street
Nanhai district
528200 Foshan, Guangdong
PEOPLE'S REPUBLIC OF CHINA

Product Category(ies):

Dental high speed air turbine handpiece,
Dental Low speed air turbine handpiece

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex V. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class IIb and III devices an additional Annex III certificate is mandatory. See also notes overleaf.

Report No.: GZ1939601

Valid from: 2020-03-09
Valid until: 2024-05-26

Date, 2020-03-09

Christoph Dicks
Head of Certification/Notified Body

TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany