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

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 056988 0013 Rev. 00

Manufacturer:

Foshan Joinchamp Medical Device Co., Ltd.
No.1, Keyang Road, Nanzhuang Town
528000 Foshan City, Guangdong
PEOPLE'S REPUBLIC OF CHINA

Product Category(ies):

Pressure Steam Sterilizer, Integral Dental Unit

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.: SH19261EXT01

Valid from: 2019-09-24
Valid until: 2024-05-26

Date, 2019-09-24

Stefan Preiß
Head of Certification/Notified Body

ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

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

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

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Facility(ies):

Foshan Joinchamp Medical Device Co., Ltd.
No.1, Keyang Road, Nanzhuang Town, 528000 Foshan City,
Guangdong, PEOPLE'S REPUBLIC OF CHINA

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