



Product Service

EC Certificate

Full Quality Assurance System

Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
(Devices in Class IIa, IIb or III)

No. G1 090932 0010 Rev. 02

Manufacturer:

Qingdao DMD Medical Technology Co., LTD

No.873 Heyuan Road, Hetao street

Hongdao Economic Zone

266113 Qingdao

PEOPLE'S REPUBLIC OF CHINA

Product Category(ies): Ligating Clips(Ti), Ligating clips(Polymer), Suction-irrigation set, Endoscopic trocar, Aortic punch, Disposable clip applier, Orthopedic suction tubes, Disposable endoscopic surgical instruments, Disposable Veress Needle.

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See also notes overleaf.

Report No.:

BJ1994407

Valid from:

2019-10-18

Valid until:

2024-05-26

Date, 2019-10-18

1. Permit

Stefan Preiß

Head of Certification/Notified Body



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Facility(ies):

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