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ZLG-BS-244.10.08

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 091024 0003 Rev. 03

Manufacturer:

Aurum Epoch Technology Co., Ltd
Unit 894, Building 1 (South), Building 27, Xincheng Dongli, Tongzhou District, 101100 Beijing
PEOPLES'S REPUBLIC OF CHINA

Product Category(ies): **Medical Nd:YAG Laser Therapy Systems, Medical Intense Pulsed Light Therapy Systems, Medical Fractional CO2 Laser Therapy Systems, Medical Diode Laser Therapy Systems.**

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.: BJ1998307

Valid from: 2026-01-10
Valid until: 2030-05-26

Date, 2026-01-10

Christoph Dicks
Head of Certification/Notified Body

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

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No. G1 091024 0003 Rev. 03

Facility(ies):

Unit 894, Building 1 (South), Building 27, Xincheng Dongli, Tongzhou District, 101100 Beijing
PEOPLES'S REPUBLIC OF CHINA

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