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Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
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ZLG-BS-244.10.08

TÜV SÜD 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 002325 0002 Rev. 01

**Manufacturer:**

**Beijing Globalipl Development Co., Ltd.**
F-8 Qunyinghui Building
Jinyuan Rd.32
Daxing Economic Development Zone
102628 Beijing
PEOPLE'S REPUBLIC OF CHINA

**Product Category(ies):** **Intense Pulsed Light for medical use + Diode Laser Equipment for medical use, CO2 Fractional Laser Equipment for medical use.**

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.:** BJ18752021

**Valid from:** 2023-10-28
**Valid until:** 2027-12-31

**Date,** 2023-10-28

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

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**No. G1 002325 0002 Rev. 01**

**Facility(ies):**

Beijing Globalipl Development Co., Ltd.
F-8 Qunyinghui Building, Jinyuan Rd.32, Daxing Economic Development Zone, 102628 Beijing, PEOPLE'S REPUBLIC OF CHINA

Beijing Globalipl Development Co., Ltd.
No. 1678, Chongdeyi Avenue, Decheng District, 253071 Dezhou City, Shandong Province, PEOPLE'S REPUBLIC OF CHINA


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