Certificate CN23/00006794

The quality management system of

BEIJING UNT Technology Co., Ltd.

M2-1 Area, Xinggu Development Zone, Pinggu District, 101200, Beijing, P.R. China
Facility number: F007060

has been assessed and certified as meeting the requirements of

MDSAP (ISO 13485:2016)

Australia: Therapeutic Goods (Medical Devices) Regulations 2002, Schedule 3, Part 1 - Full Quality Assurance System
Brazil: RDC ANVISA n. 665/2022 - Good Manufacturing Practices; RDC ANVISA n. 551/2021; RDC ANVISA n. 67/2009 - Vigilance
Canada: Medical Device Regulations SOR/98-282, Part 1
Japan: MHLW Ministerial Ordinance No.169 (2004) as amended by MHLW Ordinance No. 60 (2021); Japan PMD Act (as applicable)
USA: 21 CFR Part 803 - Medical Device Reporting; 21 CFR Part 806 - Reports of Corrections and Removals; 21 CFR Part 807 (Subparts A to D) - Establishment Registration and Device Listing, 21 CFR Part 820 - Quality System Regulation

For the following activities
Design, manufacture and servicing of Diode Laser Therapy System used in areas of general and plastic surgery and in dermatology

This certificate is valid from Effective date 2024-02-22 until Expiry date 2026-12-18 and remains valid subject to satisfactory surveillance audits.

Issue 2. Certified since 2023-12-19

L Henderson

Authorised by
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SGS UK LTD is recognised under the Medical Devices Single Audit Program. The validity of this certificate can be verified at www.SGS.com.

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