CERTIFICATE OF REGISTRATION

Electromedical Products International, Inc.

2201 Garrett Morris Parkway
Mineral Wells, Texas 76067 UNITED STATES

REPs Facility ID: F003926

UL Medical Regulatory Services of UL LLC®(UL) issues this certificate to the Firm named above, after auditing the Firm's quality management system and finding it in conformance per the defined scope with respect to:

ISO 13485:2016

with additional regulatory requirements listed on final page of this certificate.

Design and manufacture of TENS/CES equipment.

For Australia: Design and manufacture of TENS/CES equipment for the indication of pain.
For Canada: Design and manufacture of TENS/CES equipment for the indication of pain and insomnia in the areas of personal use.
For the United States: Design and manufacture of TENS/CES equipment for the indications of acute, chronic and post traumatic pain control and anxiety, insomnia and depression.

MDSAP
MEDICAL DEVICE SINGLE AUDIT PROGRAM

Authorized by

Deborah Jennings-Conner
Global Regulatory Director
UL Life and Health Sciences
UL LLC

Check Certificate Status: here

File Number	A28461	Cycle Start Date	September 13, 2021
Certificate Number	1785.210913	Effective Date	September 13, 2021
Initial Issue Date	September 13, 2018	Expiry Date	September 12, 2022

This quality system registration is included in UL's Directory of Registered Firms and applies to the provision of goods and/or services as specified in the scope of registration from the address(es) shown above. By issuance of this certificate the firm represents that it will maintain its registration in accordance with the applicable requirements. This certificate is not transferable and remains the property of UL Medical and Regulatory Services of UL LLC.
Certificates may be verified by visiting the Online Certifications Directory on UL.com.

UL REGISTERED FIRM

UL Medical and Regulatory Services UL, LLC is an MDSAP Recognized Auditing Organization

UL LLC
333 Pfingsten Road
Northbrook, IL 60062-2096 USA

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Additional Regulatory Requirements

Australia:
- Therapeutic Goods (Medical Devices) Regulations, 2002, Schedule 3 Part 1 (excluding Part 1.6) – Full Quality Assurance Procedure

Canada:
- Medical Devices Regulations – Part 1- SOR 98/282

United States:
- 21 CFR 820
- 21 CFR 803
- 21 CFR 806
- 21 CFR 807 – Subparts A to D
- 21 CFR 821 (where applicable)

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