EC Certificate Full Quality Assurance System: Certificate US98/13201

The management system of

Electromedical Products International, Inc.

2201 Garrett Morris Parkway,
Mineral Wells, TX, 76067, United States

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC

on medical devices, Annex II (excluding Section 4)

For the following products

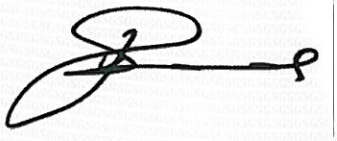
Cranial electrotherapy stimulation devices for the treatment of anxiety disorders of all levels of severity, mild depression accompanying anxiety and insomnia associated with anxiety and pain.

Where the above scope includes class III medical device(s), a valid EC Design Examination Certificate according to Annex II (Section 4) is a mandatory requirement for each device in addition to this certificate to place that device on the market.

This certificate is valid from 12 April 2016 until 12 September 2020
and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 12 September 2018
Issue 9. Certified since 11 May 1998

Certification is based on reports numbered WW/MW 07692

Authorised by



SGS United Kingdom Ltd, Notified Body 0120
202B Worle Parkway, Weston-super-Mare, BS22 6WA UK
t +44 (0)1934 522917 f +44 (0)1934 522137 www.sgs.com

SGS CE 02 0215