ZERTIFIKAT ◆ CERTIFICATE ◆ 認証証書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

Product Service

EC Certificate

Production Quality Assurance System

Directive 93/42/EEC on Medical Devices (MDD), Annex V
(Devices in Class IIa, IIb or III)

No. G2 15 09 92454 001

Manufacturer: **Danmeter-Canada Enterprise Ltd**
2781 162 Street
Surrey BC V3Z 8E4
CANADA

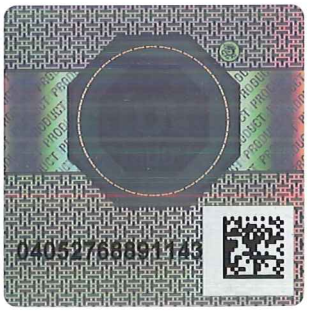
EC-Representative: **FH Service**
Falen 18B
5000 Odense C
DENMARK

Product Category(ies): **Non-invasive pain relief and muscle stimulation equipment**

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex V. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class IIb and III devices an additional Annex III certificate is mandatory. See also notes overleaf.

Report No.: 72107164

Valid from: 2016-04-06
Valid until: 2021-04-03



Date, 2016-04-06

Stefan Preiß

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

TÜV®

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Facility(ies):

Danmeter-Canada Enterprise Ltd
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