

EC Certificate Full Quality Assurance System: Certificate ES19/86817

The management system of

## CAPENERGY MEDICAL, S.L.

C/Mare de Deu de Montserrat, 41 bis - Pasaje 1º Derecha Polígono Industrial Font Santa 08970 Sant Joan Despí, Barcelona, Spain

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

Diathermy equipments for treatment of hard tissue lesions, soft tissue lesions, tendons and ligaments, Chronic Prostatitis, Dysmenorrhea and Chronic Pelvic Inflammatory Disease Pain: C 50, C 100, C200, C 300, C 400, C 500 and C25

Equipos de diatermia para el tratamiento de lesiones de tejidos duros, tejidos blandos, tendones y ligamentos, prostatitis crónica, dismenorrea y dolor pélvico crónico: C 50, C100, C 200, C 300, C 400, C 500 and C25.

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 30 September 2020 until 24 May 2024 and remains valid subject to satisfactory surveillance audits.

Issue 2. Certified since 08 May 2014 and first certified by SGS Belgium NV since 26 November 2019

Certification is based on reports numbered ES/BCN 188505

Authorised by

SGS Belgium NV, Notified Body 1639

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LPMD5007 - Certificate CE1639 Annex II-4\_EN rev. 02

Page 1 of 1





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