Declaration of Conformity

Legal Identity:

Palomar Medical Technologies, Inc
15 Network Dr.
Burlington
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USA
Tel: + 1 781-993-2300
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Authorized Representative:

CRS Medical GmbH
Ringstrasse 2
35614 Asslar
Germany
49-6441-38331-29
Palomar_eu@crs-medical.com

Equipment Identification

Type of Equipment: Intense Pulsed Light & Laser System

Model Number (s):

1500-5000 Starlux® 500 Pulsed Light and Laser System
5520-G036 LuxG Handpiece
5520-Y026 LuxY Handpiece
5520-RS26 LuxRS Handpiece
5520-V026 LuxV Handpiece
5520-R026 LuxR Handpiece
5520-YS06 LuxYS Handpiece
5520-1064P Lux1064+ Handpiece
1520-1064 Lux1064 Handpiece
5520-IR36 LuxDeepIR Handpiece
1520-1541 Lux1540 Handpiece
5520-2940 Lux2940 Handpiece
5520-1440 Lux1440 Handpiece
5520-PR06 PowR Handpiece
5520-G046 MaxG Handpiece

We, Palomar Medical Technologies, Inc., hereby declare that the devices mentioned above comply with the Swedish Medical Product Agency regulation LVFS 2003:11- transposing European Medical Devices Directive 93-42-EEC

Date of Validity: 2/18/2010
Classification: IIb According to Rule 9
Conformity Assessment Procedure: Annex II

Name of Authorized Signatory: Steven D Armstrong
Position Held in Company: Senior Vice President of Operations

Signature