EC Declaration of Conformity

Cynosure, LLC
5 Carlisle Road
Westford, MA 01886

certified according to
Council Directive 93/42/EEC of June 14 1993 concerning medical devices as amended by 2007/47/EC, Annex II Section 3.2 as certified by CERTIFICATE # **CE 41319433-05.**

Council Directive 2011/65/EU of the European Parliament and of the Council of 8 June 2011 on the restriction of the use of certain hazardous substances in electrical and electronic equipment as certified by Cynosure, LLC.

Cynosure, LLC hereby DECLARES that the

Model designation: **Picosure and PicoSure Pro Laser System**

Model Number: **105-7012-000, 100-7012-010, 105-7051-000, 100-7051-010**

comply with the Swedish Medical Product Agency regulation LVFS 2003:11- transposing European Medical Devices Directive 93-42-EEC.

Authorized EU Representative:

Cynosure BV
Veemarkt 143, 1019 CC
Amsterdam
The Netherlands
T: +31206246159

CÔNG TY TNHH THIẾT BỊ Y TẾ INAMED

Digitally signed by CÔNG TY TNHH THIẾT BỊ Y TẾ INAMED
DN: E=phung.inamed@gmail.com, OID.0.9.2342.19200300.100.1.1=MST:0312927632, CN=CÔNG TY TNHH THIẾT BỊ Y TẾ INAMED, O=CÔNG TY TNHH THIẾT BỊ Y TẾ INAMED, L="Tầng 8, Phòng 8.1, Tòa Nhà E.Town, Số 364 Cộng Hòa, Phường 13, Quận Tân Bình, Thành Phố Hồ Chí Minh, Việt Nam", S=TP Hồ Chí Minh, C=VN
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Notified Body and Identification Number: Intertek Semko AB (0413)
PO Box 1103
164 22 Kista
Sweden

Device Class: IIb per rule 9
Valid From: November 30, 2021

Kevin O'Connell
Sr. Director, Regulatory Affairs