



## 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.

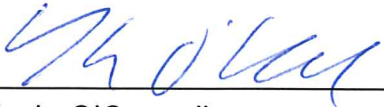
Cynosure, LLC hereby DECLARES that the  
Model designation: **Picosure and PicoSure Pro Laser System**  
Model Number: **105-7012-000, 100-7012-010, 105-7051-000, 100-7051-010**

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

Authorized EU Representative: Cynosure BV  
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Amsterdam  
The Netherlands  
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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