

*GMED certifies that the quality management system developed by*

**Apyx Medical Corporation**

**5115 Ulmerton Rd.**

**Clearwater, FL 33760 UNITED STATES**

**D.U.N.S. identification number :10-122-3287**

*for the activities*

**Conception et fabrication de dispositifs électrochirurgicaux utilisés dans des procédures ouvertes et laparoscopiques destinées à être utilisées avec des générateurs électrochirurgicaux compatibles pour la délivrance d'énergie de radiofréquence et / ou de plasma d'hélium pour la**

*Design and Manufacture of Electrosurgical Devices used in Open and Laparoscopic Procedures intended to be used with compatible electrosurgical generators for the delivery of radiofrequency energy and/or helium plasma for cutting, coagulation and ablation of soft tissue*

*performed on the location(s) of*

**Apyx Medical Corporation 5115 Ulmerton Rd. Clearwater, FL 33760 USA**

**has been audited and found to conform to the requirements of the international standard ISO 13485 : 2016 and following regulatory requirements**

Australia	Therapeutic Goods (Medical Devices) Regulations, 2002, Schedule 3 Part 1 (excluding Part 1.6) - Full Quality Assurance Procedure
Brazil	RDC ANVISA n. 16/2013 RDC ANVISA n. 23/2012 RDC ANVISA n. 67/2009
Canada	Medical Devices Regulations - Part 1 - SOR 98/282
Japan	MHLW MO 169 PMD Act
United States	21 CFR 820 21 CFR 803 21 CFR 806 21 CFR 807 - -Subparts A to D

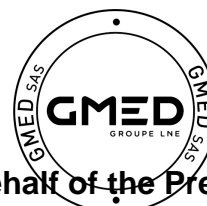
**Début de validité / Effective date March 19th, 2020 (included)**

**Valable jusqu'au / Expiry date :March 18th, 2023 (included)**

**Etabli le / Issued on : March 19th, 2020**



GMED is authorised under the Medical Devices Single Audit Program  
This certificate is issued according to the rules of GMED Certification  
The validity of this certificate can be verified on [www.gmed.fr](http://www.gmed.fr)



**On behalf of the President**  
**Béatrice LYS**  
**Technical Director**

DocuSigned by:  
*Béatrice Lys*  
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