CERTIFICATE OF REGISTRATION

N° 36681–1

GMED certifies that the quality management system developed by

ANIKA THERAPEUTICS, Inc.

32 Wiggins Avenue, Bedford,

MA 01730 UNITED STATES

Facility identifier (REPs-generated) : F004920

for the activities

Conception et fabrication d'acide hyaluronique stérile pour application chirurgicale, ophtalmique, orthopédique et de produit viscoélastique pour injection intradermique.

Design and manufacture of sterile hyaluronic acid for surgical, orthopaedic ophthalmic application, and viscoelastic product for intradermal injection.

performed on the location(s) of

ANIKA THERAPEUTICS, Inc. - 32 Wiggins Avenue, Bedford, MA 01730 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
United States	21 CFR 820 21 CFR 803 21 CFR 806 21 CFR 807 - -Subparts A to D 21 CFR 821 (where applicable)

Début de validité / *Effective date* **March 29th, 2022 (included)**

Valable jusqu'au / *Expiry date :* **March 28th, 2025 (included)**

Etabli le / *Issued on :* **March 21st, 2022**

DocuSigned by:
Beatrice Lys
EF33BDA9BAA...
GMED SAS GMED SAS

On behalf of the President

Béatrice LYS

Technical Director

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

Renouvelle le certificat 36681-0

GMED_MDSAP2016-F-V0-07-