CERTIFICATE OF REGISTRATION

N° 38111–0

GMED certifies that the quality management system developed by

Arthrosurface, Inc.
28 Forge Parkway
FRANKLIN, MA 02038 UNITED STATES

Facility identifier (REPs-generated) : F001525

for the activities

Conception et développement, fabrication et distribution d'implants orthopédiques et d'ancillaires pour le domaine de la chirurgie orthopédique.

Design and development, manufacture and distribution of orthopedic implants and ancillary instruments for the area of orthopedic surgery

performed on the location(s) of

Arthrosurface, Inc. 28 Forge Parkway, Franklin, MA 02038 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
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

Début de validité / *Effective date* **March 18th, 2021 (included)**
Valable jusqu'au / *Expiry date :* **March 17th, 2024 (included)**
Etabli le / *Issued on :* **March 18th, 2021**

MDSAP MEDICAL DEVICE SINGLE AUDIT PROGRAM

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

GMED SAS GMED GROUPE LNE GMED SAS

DocuSigned by:
Beatrice Lys
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On behalf of the President
Béatrice LYS
Technical Director

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