

*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

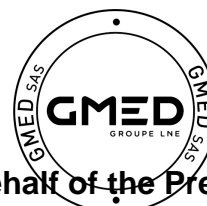
**Début de validité / Effective date March 18th, 2021 (included)**

**Valable jusqu'au / Expiry date : March 17th, 2024 (included)**

**Etabli le / Issued on : March 18th, 2021**



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)



DocuSigned by:

*Beatrice Lys*

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**On behalf of the President**

**Béatrice LYS**

**Technical Director**