CERTIFICATE OF REGISTRATION

N° 35605–2

GMED certifies that the quality management system developed by

VIANT MEDICAL, LLC

4545 Kroemer Road

Fort Wayne, IN 46818 UNITED STATES

Facility identifier (REPs-generated) : F004047

for the activities

Conception, développement, fabrication et distribution d'implants d'orthopédie et de traumatologie, d'instruments et de boîtiers de stérilisation, de plateaux et de couvercles

Design, development, manufacturing, and distribution of orthopaedics and trauma implants, instruments and sterlization cases, trays and lids

performed on the location(s) of

Viant Medical, LLC 4545 Kroemer Road, Fort Wayne, IN 46818 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* **December 9th, 2021 (included)**

Valable jusqu'au / *Expiry date :* **December 8th, 2024 (included)**

Etabli le / *Issued on :* **November 24th, 2021**

DocuSigned by:
Beatrice Lys
EF33BDA9BAA04A3...

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 35605-1

GMED_MDSAP2016-F-V0-07-