CERTIFICATE OF REGISTRATION

N° 39173–3

GMED certifies that the quality management system developed by

Clinisys, Inc.

3300 E. Sunrise Drive

Tucson, AZ 85718 UNITED STATES

Facility identifier (REPs-generated) : F006509

for the activities

Conception, développement, fabrication, distribution, installation, service et support de systèmes logiciels d'information clinique et de produits logiciels associés

Design, Development, Manufacture, Distribution, Installation, Servicing and Support of Clinical Information Systems and Associated Software Products.

performed on the location(s) of

3300 E. Sunrise Drive Tucson, AZ 85718 USA

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

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* **November 15th, 2024 (included)**

Valable jusqu'au / *Expiry date :* **November 1st, 2025 (included)**

Etabli le / *Issued on :* **November 15th, 2024**

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

Modifie le certificat 39173-2

GMED SAS GMED GROUPE LNE GMED SAS

DocuSigned by: Béatrice LYS EF33BDA9BAA04A3...

On behalf of the President

Béatrice LYS

Technical Director

GMED_MDSAP2016-F-V0-07-

MDSAP_2024 (DOC-834) Ver. 1

Approved By:

Leentje Swinnen - Author

December 12, 2024 5:42 AM MST ce2d0290-a1b0-4612-8637-4b6b2d7224fc

Version History:

Author	Effective Date	Ver.	Status
Leentje Swinnen	December 12, 2024 5:42 AM MST	1	Published
Tami Hix-Bostic	May 17, 2023 8:27 AM MST	0	Superseded