

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 2nd, 2025 (included)

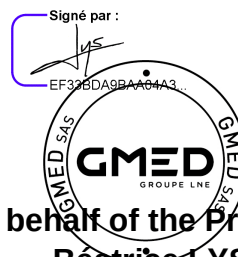
Valable jusqu'au / Expiry date : November 1st, 2028 (included)

Etabli le / Issued on : September 15th, 2025



GMED is recognised 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 39173-3



**On behalf of the President
Béatrice LYS
Technical Director**