# CERTIFICATE OF REGISTRATION

## N° 40112–0



*GMED certifies that the quality management system developed by*

**UNION PLASTIC**
**ZA Robert**
**43140 SAINT DIDIER EN VELAY FRANCE**

**Facility identifier (REPs-generated) : F007667**

*for the activities*

**Conception, fabrication et distribution d'adaptateurs de flacons pour transfert stérile et mélange de préparations médicamenteuses.**

*Design, manufacture and distribution on vial adapters for sterile transfer and mixing of medicinal preparations.*

*performed on the location(s) of*

**Site UP'PHARM, Z.A. Robert, 43140 SAINT DIDIER EN VELAY, FRA**
**Site UPE, Zone Industrielle Route de Souppes, 75570 CHATEAU-LANDON, FRA**

**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<br>21 CFR 803<br>21 CFR 806<br>21 CFR 807 - -Subparts A to D |

**Début de validité** / *Effective date* **January 10th, 2025 (included)**
**Valable jusqu'au** / *Expiry date :* **January 9th, 2028 (included)**
**Etabli le** / *Issued on :* **January 10th, 2025**

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

DocuSigned by:
Lionel DREUX
A1D80E08C60D47A...

GMED SAS GMED SAS

**Lionel DREUX**
**President**

GMED_MDSAP2016-F-V0-07-