



CERTIFICATE OF GDP COMPLIANCE

We certify herewith

that the company **Spirig HealthCare AG, Industriestrasse 30, 4622 Egerkingen**, Authorisation No. 512295-102758127 with its site **Spirig HealthCare AG, Industriestrasse 28/30, 4622 Egerkingen, Switzerland**, Site No. 1106335 has been duly authorised to distribute medicinal products resp. API / intermediates according to the table below;

that the company is keeping the required level for Good Distribution Practices for Medicinal Products (GDP) according to the Swiss regulations in force. These regulations are in accordance with the requirements of the following documents:

- Guidelines of the European Commission on Good Distribution Practice of Medicinal Products for Human Use (2013/C 343/01)
- Commission Implementing Regulation (EU) 2021/1248 on Good Distribution Practice for Veterinary Medicinal Products
- Guidelines of the European Commission on Good Distribution Practice of active substances for medicinal products for human use (2015/C 95/01)
- Commission Implementing Regulation (EU) 2021/1280 on Good Distribution Practice for active substances for veterinary medicinal products.

that the company is subject to official periodic inspections; the last regular inspection has been performed on **10.02.2026** (dd.mm.yyyy);

that this certificate reflects the status of the premises at the time of the inspection noted above and should not be relied upon to reflect the compliance status if more than five years have elapsed since the date of that inspection.

The authenticity of this certificate may be verified in SwissGMDP. If it does not appear, please contact Swissmedic.

No.	Operation	Scope*
S.2	IMPORT OF MEDICINAL PRODUCTS (WITHOUT LABILE BLOOD PRODUCTS)	
S.2.1	Import of non- ready-to-use medicinal products	
S.2.1.1	Medicinal products (intermediates, without immunological and blood products)	H/V
S.2.2	Import of ready-to-use medicinal products, including market release	
S.2.2.1	Medicinal products (without immunological and blood products)	H/V
S.2.2.2	Immunological medicinal products	H/V
S.2.3	Import of ready-to-use medicinal products, excluding market release	
S.2.3.1	Medicinal products (without immunological and blood products)	H/V
S.2.3.2	Immunological medicinal products	H/V
S.2.3.4	The import of ready-to-use medicinal products, excluding market release, is restricted to:	

No.	Operation	Scope*
S.2.3.4.2	the import on behalf of the marketing authorisation holder	H/V
S.2.6	Outsourcing of manufacture of medicinal products as contract giver	H/V
The authorised importation operations do not include the storage of medicinal products		
S.4	WHOLESALE DISTRIBUTION OF MEDICINAL PRODUCTS (WITHOUT LABILE BLOOD PRODUCTS)	
S.4.2	Wholesale distribution of ready-to-use medicinal products, including market release	
S.4.2.1	Medicinal products (without immunological and blood products)	H/V
S.4.2.2	Immunological products	H/V
S.4.3	Wholesale distribution of ready-to-use medicinal products, excluding market release	
S.4.3.1	Medicinal products (without immunological and blood products)	H/V
S.4.3.2	Immunological products	H/V
S.4.3.3	Blood products	H/V
S.4.6	Outsourcing of manufacture of medicinal products as contract giver	H/V
The authorised wholesale operations do not include the storage of medicinal products		
S.5	EXPORT OF MEDICINAL PRODUCTS (WITHOUT LABILE BLOOD PRODUCTS)	
S.5.2	Export of ready-to-use medicinal products	
S.5.2.1	Medicinal products (without immunological and blood products)	H/V
S.5.2.2	Immunological products	H/V
The authorised exportation operations do not include the storage of medicinal products		

* Scope of authorisation:

H/V	Prefix S: Human and veterinary medicinal products/ Prefix ST: Human TpP/GT/GVO, without investigational products
V	Veterinary medicinal products only, without investigational products
I	Prefix S: Human investigational medicinal products/ Prefix: ST: Human Investigational TpP/GT/GVO
-	Not specified

Bern, **21.04.2026** (dd.mm.yyyy)

No. GDP-CH-1008351

Swissmedic, Swiss Agency for Therapeutic Products.