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CERTIFICATE OF GMP COMPLIANCE

We certify herewith

that the company **Band-Genossenschaft, Riedbachstrasse 9, 3027 Bern,** Authorisation No. 512475-102656783 with its site **Band-Genossenschaft Abteilung Food & Pharma, Murtenstrasse 350, 3027 Bern, Switzerland,** Site No. 1005806 has been duly authorised to perform the manufacturing activities according to the table below;

that the company is keeping the required level for Good Manufacturing Practices for Medicinal Products (GMP) according to the Swiss regulations in force. These regulations are in accordance with the requirements for good practices in the manufacture and quality control of the Pharmaceutical Inspection Convention/Co-operation Scheme (PIC/S) as well as with The Good Manufacturing Practice requirements referred to in the Agreement of Mutual Recognition between the European Union/Canada and Switzerland;

that the manufacturing plant of the company is subject to official periodic inspections; the last regular inspection was conducted on **23.02.2022** (dd.mm.yyyy).

No.	*Operation*	*Scope**
1	**MANUFACTURE OF MEDICINAL PRODUCTS (WITHOUT LABILE BLOOD PRODUCTS)**	
1.5	**Packaging**	
1.5.2	Secondary packaging	H/V

* Scope of authorisation:
H/V Human and veterinary medicinal products, without investigational products
V Veterinary medicinal products only, without investigational products
I Human investigational medicinal products
– Not specified

Berne, **04.08.2022** (dd.mm.yyyy)
No. GMP-CH-1003411

Swissmedic, Swiss Agency for Therapeutic Products

Luxshana Santhirasegarar

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CERTIFICATE OF GDP COMPLIANCE

We certify herewith

that the company **Band-Genossenschaft, Riedbachstrasse 9, 3027 Bern,** Authorisation No. 512475-102656783 with its site **Band-Genossenschaft Abteilung Food & Pharma, Murtenstrasse 350, 3027 Bern, Switzerland**, Site No. 1005806 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 **23.02.2022** (dd.mm.yyyy).

No.	*Operation*	*Scope**
S.4	**WHOLESALE DISTRIBUTION OF MEDICINAL PRODUCTS (WITHOUT LABILE BLOOD PRODUCTS)**	
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)	-

* Scope of authorisation:
H/V Human and veterinary medicinal products, without investigational products
V Veterinary medicinal products only, without investigational products
I Human investigational medicinal products
– Not specified

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Berne, **04.08.2022** (dd.mm.yyyy)
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Swissmedic, Swiss Agency for Therapeutic Products

Luxshana Santhirasegarar

Schweizerisches Heilmittelinstitut
Institut suisse des produits thérapeutiques
Istituto svizzero per gli agenti terapeutici
Swiss Agency for Therapeutic Products

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