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CERTIFICATE OF GDP COMPLIANCE

We certify herewith

that the company **ebi-pharm ag, Lindachstrasse 8c, 3038 Kirchlindach,** Authorisation No. 511304-102618263 with its site **ebi-pharm ag, Lindachstrasse 8c, 3038 Kirchlindach, Switzerland**, Site No. 1000453 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 Guidelines on Good Distribution Practice of Medicinal Products for Human Use (2013/C 343/01) of the European Commission and with the requirements of the European GMP Part II (Basic Requirements for Active Substances used as Starting Materials);

that the company is subject to official periodic inspections; the last regular inspection has been performed on **07.05.2019** (dd.mm.yyyy).

No.	*Operation*
S.2	**IMPORT OF MEDICINAL PRODUCTS (WITHOUT LABILE BLOOD PRODUCTS)**
S.2.2	**Import of ready-to-use medicinal products, including market release**
S.2.2.1	Medicinal products (without immunological and blood products)
S.2.2.2	Immunological medicinal products
S.2.6	**Outsourcing of manufacture of medicinal products as contract giver**
	The authorised activities do not include the storage of medicinal products
S.4	**WHOLESALE DISTRIBUTION OF MEDICINAL PRODUCTS (WITHOUT LABILE BLOOD PRODUCTS)**
S.4.1	**Wholesale distribution of non- ready-to-use medicinal products**
S.4.1.1	Medicinal products (without immunological and 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)
S.4.6	**Outsourcing of manufacture of medicinal products as contract giver**
	The authorised activities 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)
	The authorised activities do not include the storage of medicinal products

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Berne, **24.09.2019** (dd.mm.yyyy)
No. GDP-CH-1000472

Swissmedic, Swiss Agency for Therapeutic Products

Dr. Georges Meseguer

Swiss Agency for Therapeutic Products
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CERTIFICATE OF GDP COMPLIANCE

We certify herewith

that the company **ebi-pharm ag, Lindachstrasse 8c, 3038 Kirchlindach,** Authorisation No. 511304-102618263 with its site **ebi-pharm ag, Nüchternweg 2, 3038 Kirchlindach, Switzerland**, Site No. 1001420 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 Guidelines on Good Distribution Practice of Medicinal Products for Human Use (2013/C 343/01) of the European Commission and with the requirements of the European GMP Part II (Basic Requirements for Active Substances used as Starting Materials);

that the company is subject to official periodic inspections; the last regular inspection has been performed on **07.05.2019** (dd.mm.yyyy).

No.	*Operation*
S.2	**IMPORT OF MEDICINAL PRODUCTS (WITHOUT LABILE BLOOD PRODUCTS)**
S.2.2	**Import of ready-to-use medicinal products, including market release**
S.2.2.1	Medicinal products (without immunological and blood products)
S.2.2.2	Immunological medicinal products
S.2.6	**Outsourcing of manufacture of medicinal products as contract giver**
S.4	**WHOLESALE DISTRIBUTION OF MEDICINAL PRODUCTS (WITHOUT LABILE BLOOD PRODUCTS)**
S.4.1	**Wholesale distribution of non- ready-to-use medicinal products**
S.4.1.1	Medicinal products (without immunological and 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)
S.4.6	**Outsourcing of manufacture of medicinal products as contract giver**
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)

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Berne, **24.09.2019** (dd.mm.yyyy)
No. GDP-CH-1000473

Swissmedic, Swiss Agency for Therapeutic Products

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CERTIFICATE OF GDP COMPLIANCE

We certify herewith

that the company **ebi-pharm ag, Lindachstrasse 8c, 3038 Kirchlindach,** Authorisation No. 511304-102618263 with its site **ebi-pharm ag, Nüchternweg 1-3, 3038 Kirchlindach, Switzerland**, Site No. 1003419 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 Guidelines on Good Distribution Practice of Medicinal Products for Human Use (2013/C 343/01) of the European Commission and with the requirements of the European GMP Part II (Basic Requirements for Active Substances used as Starting Materials);

that the company is subject to official periodic inspections; the last regular inspection has been performed on **07.05.2019** (dd.mm.yyyy).

No.	*Operation*
S.4	**WHOLESALE DISTRIBUTION OF MEDICINAL PRODUCTS (WITHOUT LABILE BLOOD PRODUCTS)**
S.4.1	**Wholesale distribution of non- ready-to-use medicinal products**
S.4.1.1	Medicinal products (without immunological and 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)
S.4.6	**Outsourcing of manufacture of medicinal products as contract giver**

Berne, **24.09.2019** (dd.mm.yyyy)
No. GDP-CH-1000474

Swissmedic, Swiss Agency for Therapeutic Products

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