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CERTIFICATE OF GMP COMPLIANCE

We certify herewith

that the company **Eurofins BioPharma Product Testing Switzerland AG, Parkstrasse 10, 5012 Schönenwerd,** Authorisation No. 511435-102671921 with its site **Eurofins BioPharma Product Testing Switzerland AG, Parkstrasse 10, 5012 Schönenwerd, Switzerland,** Site No. 1102792 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 **13.01.2022** (dd.mm.yyyy).

No.	*Operation*	*Scope**
1	**MANUFACTURE OF MEDICINAL PRODUCTS (WITHOUT LABILE BLOOD PRODUCTS)**	
1.6	**Quality control testing**	
1.6.2	Microbiological: non-sterility	H/V, I
1.6.3	Chemical/Physical	H/V, I
1.6.4	Biological	H/V, I
3	**MANUFACTURE OF ACTIVE SUBSTANCES**	
3.6	**Quality control testing**	
3.6.1	Physical / Chemical testing	-
3.6.2	Microbiological: testing (excluding sterility testing)	-
3.6.4	Biological Testing	-

* 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, **28.03.2022** (dd.mm.yyyy)
No. GMP-CH-1003080

Swissmedic, Swiss Agency for Therapeutic Products

[redacted]

Marianne Baumann

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