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CERTIFICATE OF GMP COMPLIANCE

We certify herewith

that the company **PUREXIS SA, Via Violino 1, 6928 Manno,** Authorisation No. 511260-102606338 with its site **PUREXIS SA, Via Violino 1, 6928 Manno, Switzerland,** Site No. 1005923 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 **22.06.2021** (dd.mm.yyyy).

No.	*Operation*	*Scope**
3	**MANUFACTURE OF ACTIVE SUBSTANCES**	
3.5	**General finishing steps**	
3.5.2	Primary packaging	-
3.5.3	Secondary packaging	-

* 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, **18.08.2021** (dd.mm.yyyy)
No. GMP-CH-1002439

Swissmedic, Swiss Agency for Therapeutic Products

Swiss Agency for Therapeutic Products
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