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CERTIFICATE OF GMP COMPLIANCE

We certify herewith

that the company **Cannapharm AG, 3400 Burgdorf,** Authorisation No. 511581-102615809 with its site **Cannapharm AG, Burgergasse 50, 3400 Burgdorf, Switzerland,** Site No. 1002173 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 **15.08.2019** (dd.mm.yyyy).

No.	*Operation*	*Scope**
1	**MANUFACTURE OF MEDICINAL PRODUCTS (WITHOUT LABILE BLOOD PRODUCTS)**	
1.5	**Packaging**	
1.5.2	Secondary packing	I
S.1.8	**Blinding of medicinal products for clinical trials**	
The authorised activites solely include the issuing of contracts for the manufacture of magistral formula medicinal 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

Berne, **24.09.2019** (dd.mm.yyyy)
No. GMP-CH-1000464

Swissmedic, Swiss Agency for Therapeutic Products

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