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CERTIFICATE OF GMP COMPLIANCE

We certify herewith

that the company **Amino AG, Fabrikation pharmazeutischer und chemischer Produkte, Wiesenstrasse 21, 5412 Gebenstorf** with its site **Amino AG, Fabrikation pharmazeutischer und chemischer Produkte, Ämmenmattstrasse 2, 3123 Belp, Switzerland**, has been duly authorized to manufacture and distribute medicinal products and investigational medicinal products;

that the company is manufacturing the following dosage forms:

- liquid dosage forms including aseptically prepared forms restricted to aseptical products for topical use (i.e. eye drops, eye water Zeller) and terminally sterilised forms
- semi-solid dosage forms including aseptically prepared forms restricted to aseptical products for topical use (i.e. ointments for wounds and eyes) and terminally sterilised forms
- investigational medicinal products

that the finished medicinal products put on the market in Switzerland by the company are subject to appraisal and authorisation by our agency;

that the company is keeping the required level for good practices in the manufacture of medicinal products and investigational medicinal products 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) and the Directives of the European Commission;

that the manufacturing plant of the company is subject to official periodic inspections; the last regular inspection was conducted on **March 13-15, 2017**;

that the requirements regarding manufacture and quality control for medicinal products and investigational medicinal products for export are identical to those applicable to medicinal products and investigational medicinal products sold in Switzerland.

Berne, April 4, 2018
No. 18-0718

Swiss Agency for Therapeutic Products swissmedic

Swissmedic, Swiss Agency for Therapeutic Products

Dr. Alfred Ryf

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