

CERTIFICATE OF GMP COMPLIANCE

We certify herewith

that the company **SPAGYRIK Produktions AG, Bachweg 3, 3400 Burgdorf, Switzerland**, has been duly authorized to manufacture and distribute medicinal products and active pharmaceutical ingredients (restricted to homeopathics, spagyrics and phytotherapeutics), the manufacturing licence excluding sterile products and including following dosage forms:

- liquid dosage forms
- semi-solid dosage forms (restricted to manufacturing of ointment bases)

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 pharmaceutical products and active pharmaceutical ingredients 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 7-8, 2016**;

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

Berne, July 6, 2016
No. 16-1294

Swissmedic, Swiss Agency for
Therapeutic Products



Dr. Alfred Ryf

