

## CERTIFICATE OF GMP COMPLIANCE

We certify herewith

that the company **DISCH AG, Bahnhofstrasse 21, 5504 Othmarsingen, Switzerland**, has been duly authorized to manufacture and distribute medicinal products and investigational medicinal products, the manufacturing licence excluding sterile products and including following dosage forms:

- solid dosage forms

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 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 **August 19-21, 2014**;

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

Berne, December 8, 2014  
**No. 14-2307**

Swissmedic, Swiss Agency for  
Therapeutic Products

Dr. Alfred Ryf

A handwritten signature in blue ink, appearing to read "Dr. Alfred Ryf", is written over the printed name.