

CERTIFICATE OF GMP COMPLIANCE

We certify herewith

that the company **Eurofins Scientific AG, Parkstrasse 10, 5012 Schönenwerd, Switzerland**, has been duly authorized to manufacture medicinal products, the manufacturing licence including following activities:

- Quality control (chemical, physical and biochemical) of medicinal products as contract laboratory
- Quality control (microbiological) of medicinal products as contract laboratory excluding tests of sterility

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 inspection was conducted on **August 16-17, 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, December 5, 2016
No. 16-2328

Swissmedic, Swiss Agency for
Therapeutic Products

A blue ink signature of Dr. Georges Meseguer is written over the text of the Swiss Agency for Therapeutic Products.

Dr. Georges Meseguer