

CERTIFICATE OF GMP COMPLIANCE

We certify herewith

that the company Passafaro Analytic Lab with its site of Im Kloster 9a, 8240 Thayngen, Switzerland, has been duly authorized to manufacture medicinal products, the manufacturing licence including following activities:

- Quality control (chemical, physical and biochemical) of medicinal products

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 **January 27, 2004**;

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.

Bern, July 19, 2004 No. **04-781**

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Swissmedic, Swiss Agency for Therapeutic Products

Michel Keller