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CERTIFICATE OF GMP COMPLIANCE

We certify herewith

that the company **Oligopharm SA** with its site **Route de Montheron 8b, 1053 Cugy, Switzerland**, has been duly authorized to manufacture and distribute medicinal products, the manufacturing licence excluding sterile products and including following dosage forms:

- liquid dosage forms (restricted to solutions of oligoelements)
- solid dosage forms (restricted to the filling of hard capsules)

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 **May 10, 2011**;

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 1st, 2011
No. 11-1727

Swissmedic, Swiss Agency for Therapeutic Products

Michel Keller

Swiss Agency for Therapeutic Products
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