

CERTIFICATE OF GMP COMPLIANCE

We certify herewith

that the company **Laboratoire homéopathique D. Schmidt-Nagel SA, Rue du Pré-Bouvier 27, CH-1242 Satigny, Switzerland** with its site of **Rue du Pré-Bouvier 27, CH-1217 Meyrin, Switzerland**, has been duly authorized to manufacture and distribute medicinal products, the manufacturing licence excluding sterile products and being restricted to small quantities of homeopathics, phytopharmaceuticals and trace (oligo) elements in following dosage forms:

- liquid dosage forms
- semi-solid forms
- solid dosage forms

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 **November 23, 2006**;

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, December 15, 2006
No. **06-1509**



Swissmedic, Swiss Agency for
Therapeutic Products



Dr. Georges Meseguer