swissmedic

CERTIFICATE OF GMP COMPLIANCE

We certify herewith

that the **Laboratoire homéopathique D. Schmidt-Nagel SA, rue du Pré-Bouvier 27, 1242 Satigny** with its site **Laboratoire homéopathique D. Schmidt-Nagel SA, rue du Pré-Bouvier 27, 1217 Meyrin, Switzerland**, has been duly authorized to manufacture and distribute medicinal products;

that the company is manufacturing the following dosage forms:

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

the activities are limited to homeopathic medicinal products and trace elements

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 medicinal 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 **June 14, 2017**;

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

Berne, November 7, 2017
No. 17-2140

Swissmedic, Swiss Agency for Therapeutic Products

Dr. Alfred Rüf

Swiss Agency for Therapeutic Products
swissmedic

Swissmedic | Hallerstrasse 7 | Postfach | CH-3000 Bern 9 | www.swissmedic.ch | Tel. +41 58 462 02 11 | Fax +41 58 462 02 12