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CERTIFICATE OF GMP COMPLIANCE

We certify herewith

that the company **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, the manufacturing licence excluding sterile products and including 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 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 **June 25, 2015**;

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

Berne, September 1, 2015
No. 15-1983

Swiss Agency for Therapeutic Products
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Michel Keller

Schweizerisches Heilmittelinstitut
Institut suisse des produits thérapeutiques
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