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CERTIFICATE OF GMP COMPLIANCE

We certify herewith

that the company **AromaSan Sàrl, Route de Montheron 8 B, 1053 Cugy, Switzerland**, has been duly authorized to manufacture and distribute active pharmaceutical ingredients (APIs), the manufacturing licence excluding sterile API restricted to repackaging of essential oils as active pharmaceutical ingredients (APIs);

that the company is keeping the required level for good practices in the manufacture of 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 **July 8, 2011**;

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

Berne, March 12, 2012
No. 12-536

Swissmedic, Swiss Agency for Therapeutic Products

Dr. Alfred Ryf

Swiss Agency for Therapeutic Products
swissmedic

Schweizerisches Heilmittelinstitut
Institut suisse des produits thérapeutiques
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Swiss Agency for Therapeutic Products

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