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CERTIFICATE OF GMP COMPLIANCE

We certify herewith

that the company **Osmopharm SA, via alle Fornaci, 6930 Bedano, Switzerland**, has been duly authorized to manufacture and distribute medicinal products and investigational medicinal products, the manufacturing licence excluding sterile products and including following dosage forms:

- solid dosage forms

including following packaging activities:

- packaging of investigational medicinal products

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 **April 5-6, 2016**;

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

Berne, July 6, 2016
No. 16-1270

Swissmedic, Swiss Agency for Therapeutic Products

Dr. Alfred Ryf

Swiss Agency for Therapeutic Products
swissmedic

Schweizerisches Heilmittelinstitut
Institut suisse des produits thérapeutiques
Istituto svizzero per gli agenti terapeutici
Swiss Agency for Therapeutic Products

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