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CERTIFICATE OF GMP COMPLIANCE

We certify herewith

that the company **Frike Pharma AG, Auenstrasse 11, 8617 Mönchaltorf, 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

including following packaging activities:

- Primary packaging of medicinal products
- Secondary packaging of medicinal products

all activities are limited to medicinal products which are sold in Switzerland as non-prescription medicine;

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 **February 24-26, 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 17, 2015
No. 15-2117

Swiss Agency for Therapeutic Products
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Swissmedic, Swiss Agency for Therapeutic Products

Dr. Alfred Ryf

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

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