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CERTIFICATE OF GMP COMPLIANCE

We certify herewith

that the company **Cannapharm AG, 3400 Burgdorf** with its site **Cannapharm AG, c/o Markus Lüdi, Burgergasse 50, 3400 Burgdorf, Switzerland**, has been duly authorized to manufacture and distribute medicinal products, the manufacturing licence including following packaging activities:

- Secondary packaging of medicinal products including randomisation of medicinal products for clinical trials

that the company is keeping the required level for good practices in the manufacture of pharmaceutical products and 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 **January 20, 2015**;

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

Berne, October 26, 2016
No. 16-2012

Swissmedic, Swiss Agency for Therapeutic Products

Michel Keller

Swiss Agency for Therapeutic Products
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