

CERTIFICATE OF GMP COMPLIANCE

We certify herewith

that the company **Microsynth AG, Schützenstrasse 15, 9436 Balgach, Switzerland**, has been duly authorized to manufacture and distribute active pharmaceutical ingredients (APIs) and medicinal products

- quality control (chemical, physical, biochemical) of medicinal products as contract laboratory
- quality control (biological) of medicinal products as contract laboratory

The activities are restricted to the DNA sequencing with the Sanger method

that the company is keeping the required level for good practices in the manufacture of active pharmaceutical ingredients (APIs) and medicinal 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 15, 2018**;

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

Berne, July 20, 2018
No. 18-1647



Swissmedic, Swiss Agency for
Therapeutic Products



Dr. Georges Meseguer