

CERTIFICATE OF GMP COMPLIANCE

We certify herewith

that the company **Swiss Stem Cell Foundation, Via Tesserete 48, 6900 Lugano, Switzerland** with its site **Swiss Stem Cell Foundation, Technoparkstrasse 1, 8005 Zürich, Switzerland**, has been duly authorized to manufacture and distribute transplant products, the manufacturing licence includes the following product categories:

- investigational cell therapy products for autologous use
- investigational tissue products for autologous use

that the finished transplant products put on the market in Switzerland by the company are subject to appraisal and authorisation by our agency;

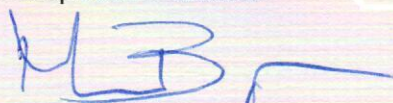
that the company is keeping the required level for good practices in the manufacture of pharmaceutical products according to the Swiss regulations in force. These regulations are in accordance with the requirements for good practices in the manufacture and quality control of the Pharmaceutical Inspection 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 **28 march 2012**;

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

Berne, December 16, 2014
No. 14-2432

Swissmedic, Swiss Agency for
Therapeutic Products



Dr. Martin Burgener