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CERTIFICATE OF GMP COMPLIANCE

We certify herewith

that the company **Lipomed AG, Fabrikmattenweg 4, 4144 Arlesheim, Switzerland**, has been duly authorized to manufacture and distribute medicinal products, active pharmaceutical ingredients (APIs) and investigational medicinal products, the manufacturing licence excluding sterile API and including following types of active pharmaceutical ingredients:

- highly active or sensitising APIs (Cladibrine and Deferiprone)

including following packaging activities:
- Secondary packaging of medicinal products

that the company is keeping the required level for good practices in the manufacture of 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 **June 13-15, 2016;**

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

Berne, May 15, 2017
No. 17-1018

Swissmedic, Swiss Agency for Therapeutic Products

Dr. Georges Meseguer

Swiss Agency for Therapeutic Products
swissmedic

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