

CERTIFICATE OF GMP COMPLIANCE

(for the Health Authorities of Taiwan)

We certify herewith

that the company
its site

廠名
廠址

is with

has been duly authorized to manufacture and distribute active pharmaceutical ingredients (APIs) as well investigational APIs, the manufacturing licence and including following types of active pharmaceutical ingredients:

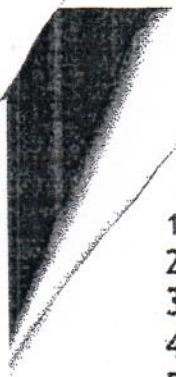
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that the company manufactures following API(s): see annex

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

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

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