swissmedic

FREE SALES CERTIFICATE

Nr.: G-FSC-12-20670 valid until: 2 July 2015

The SWISS AGENCY FOR THERAPEUTIC PRODUCTS certifies herewith, that medical devices are regulated in Switzerland under the Federal Law on Medicinal Products and Medical Devices (Law on Therapeutic Products) of 15 December 2000 in force since 1 January 2002 and the Medical Devices Ordinance of 17 October 2001 in force since 1 January 2002.

The following medical device(s) system meets (meet) the legal requirements set out in the Swiss Medical Devices Ordinance and which incorporates the Medical Devices Directives of the European Community:

Medical Devices for Hemodialysis:

- **Fistula Needles**
- **Filters for Hemodialysis**
- **Blood Lines for Hemodialysis**

Therefore, the firm DIALIFE SA, Via al Fiume 3, 6807 Taverne, Switzerland,

in conformity with the laws of Switzerland is authorized to develop, manufacture and sell on the Swiss market and to export into any country the medical device(s) system above-mentioned.

This certificate is valid until 2 July 2015

Bern, 3 July 2012

Swiss Agency for Therapeutic Products
Medical Devices Division

Swiss Agency for Therapeutic Products

Claude-Philippe Pettipierre, Master of Law Fee: CHF 300.00

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Schweizerisches Heilmittelinstitut
Institut suisse des produits thérapeutiques
Istituto svizzero per gli agenti terapeutici
Swiss Agency for Therapeutic Products

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