

FREE SALES CERTIFICATE

Nr.: FSC-18-21163

valid until: 27 June 2021

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) meets (meet) the legal requirements set out in the Swiss Medical Devices Ordinance and which incorporates the Medical Devices Directives of the European Union:

- **The sterile Surgical Sutures:**

Absorbable: AssuCryl® (PGA), AssuCryl® Rapid (PGA), AssuCryl® Lactin (PGLA), AssuCryl® MonoSlow (PDO), AssuCryl® MonoRapid (PGCL)

Non-absorbable: Astralen / Polyester, Astralen / Polyester Tape, Silk, Nylon / Polyamide, Supramid, Polypropylene, Steel, Pledgets (PTFE), AssuTopFiber®, Temporary Pacing-Wire

- **Sterile Microsurgical Knives**

Therefore, the firm ASSUT MEDICAL Sàrl, Avenue de Rochettaz 57, 1009 Pully-Lausanne, Switzerland (legal manufacturer); Assut Medical Sàrl, Av. de Lavaux 35, 1009 Pully-Lausanne, Switzerland (offices); Assut Medical Sàrl, Sur le Crêt 13, 2606 Corgémont, Switzerland (production site),

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 CE marked medical device(s) above-mentioned.

This certificate is valid until 27 June 2021

Bern, 28 June 2018

Swiss Agency for Therapeutic Products
Medical Devices Division



Claude-Philippe Petitpierre, Master of Law

Fee: CHF 300.00