swissmedic

FREE SALES CERTIFICATE

Nr.: FSC-18-21573 | valid until: 26 August 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:

Surgical Instruments for Ophthalmology, according to the attached list (2 pages).

Therefore, the firm H. P. Braem AG, Industriestrasse 4, 9552 Bronschhofen, Switzerland,

in conformity with the medical devices law 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 26 August 2021

Bern, 27 August 2018

Swiss Agency for Therapeutic Products
Medical Devices Division

Swiss Agency for Therapeutic Products swissmedic

Claude-Philippe Petitpierre, Master of Law

Fee: CHF 300.00

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

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