swissmedic

FREE SALES CERTIFICATE

Nr.: FSC-17-21638 | valid until: 18 July 2020

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:

- **EasyOne**
- **EasyOne Air**
- **Easy on-PC**
- **EasyOne Pro**
- **EasyOne Pro LAB**
  incl. Accessories and Consumables.

Therefore, the firm ndd Medizintechnik AG, Technoparkstrasse 1, 8005 Zürich, 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 18 July 2020

Bern, 19 July 2017

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

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