

EU Declaration of Conformity

Manufacturer	NDD Medizintechnik AG
	Technoparkstrasse 1
	CH-8005 Zürich, Switzerland
Product Family	Spirometers
Basic UDI-DI	764014219SpirometerBS
Single Registration Number	CH-MF-000015550
Authorized	NDD Medizintechnik GmbH
Representative	Endersbacher Strasse 49
	DE-71334 Waiblingen
	Germany
	SNR: DE-AR-000032322
Product Trade Name &	Easy on-PC
Catalogue Number	REF: 2700-3
CND code	Z121501 – Spirometry Instruments
Classification	Class IIa according to (EU) 2017/745, Annex VIII, Rule 10
Common Specifications	See List of Applied Standards

We hereby declare our sole responsibility for the EU Declaration of Conformity.

The devices covered by this declaration are in conformity with the European Medical Device Regulation (EU) 2017/745 as well as other relevant Union legislations that make provisions for the issuing of a declaration of conformity.

ndd Medizintechnik AG follows the Conformity Assessment procedure based on a Quality Management pursuant to Regulation (EU) 2017/745, Annex IX, which involves the intervention of the Notified Body:

TÜV SÜD Product Service GmbH, Notified Body 0123

Ridlerstrasse 65, 80339 Munich, Germany

This Declaration of Conformity is applicable to all products listed and released after the Date of Issuance of this Declaration of Conformity, and is valid until the expiry date of the CE Certificate G10 005204 0004.

Andreas Senn

Director of Quality Services

Michael Bencak

CEO

Zurich, 30. May 2023