

## **EU Declaration of Conformity**

Manufacturer NDD Medizintechnik AG

Technoparkstrasse 1

CH-8005 Zürich, Switzerland

**Product Family** Barriette

**Basic UDI-DI** 764014219BarrietteBV

Single Registration Number n/a

> NDD Medizintechnik GmbH **Endersbacher Strasse 49**

DE-71334 Waiblingen, Germany **Authorized Representative** 

SRN: DE-AR-000032322

**Product Trade Name & Catalogue Number** 

3050-0 (basic unit), 3050-1, 3050-2 (packaging

configurations)

**DLCO Barriette:** 

**FRC Barriette:** 

3150-0 (basic unit), 3150-1, 3150-2 (packaging

configurations)

**CND** code Z12150185 Spirometry Instruments - Consumables

Classification Class I according to (EU) 2017/745, Annex VIII, Rule 1

**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 procedure related to the EU Declaration of Conformity set out in Annex IV of Regulation (EU) 2017/745.

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 a revised Declaration of Conformity is issued.

Michael Bencak

CEO

Andreas Senn

Director Quality, Regulatory Affairs & Clinical Affairs

17. Aug. 2023

DoC Barriette MDR V02, V02 Form: ID:7000029, Rev:00