

## EU Declaration of Conformity

<b>Manufacturer</b>	<b>NDD Medizintechnik AG</b> Technoparkstrasse 1 8005 Zürich, Switzerland
<b>Product Family</b>	Breathing Tubes
<b>Basic UDI-DI</b>	764014219BreathingTubes6L
<b>Single Registration Number</b>	CH-MF-000015550
<b>Authorized Representative</b>	<b>NDD Medizintechnik GmbH</b> Endersbacher Strasse 49 DE-71334 Waiblingen Germany  SNR: DE-AR-00032322
<b>Product Trade Name &amp; Catalogue Number</b>	<b>Spirette M (8mm):</b> 2050-78, 2050-7 <b>Filter Adapter:</b> 2030-01, 2030-02, 2030-11, 2030-21 <b>Spirette FA Pro/LAB:</b> 2050-70, 2050-71, 2050-72, 2050-73
<b>CND code</b>	Z12150185 Spirometry Instruments - Consumables
<b>Classification</b>	Class I according to (EU) 2017/745, Annex VIII, Rule 1
<b>Common Specifications</b>	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.

n dd 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.



Andreas Senn  
Director of Quality Services

Zurich, 30. May 2023



Michael Bencak  
CEO