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KONFORMITÄTS-ERKLÄRUNG / DECLARATION OF CONFORMITY

To whom it may concern

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Münsingen, 21.03.2023

EU DECLARATION OF CONFORMITY

We, Sensopro AG (CHRN-MF-20000082; SRN CH-MF-000010122), declare under our sole responsibility that the medical devices with Basic UDI-DI 7649991448SensoproLunaGV:

Device name	**UDI-DI**
Sensopro Luna Physio	7649991448021 7649991448038 7649991448045
Sensopro Tube Medical (8 pcs.) Aquamarine / weak	7649991448090
Sensopro Tube Medical (8 pcs.) Azure blue / medium	7649991448106
Sensopro Tube Medical (8 pcs.) Coral-red / strong	7649991448113
Sensopro Tube Aquamarin/Aquamarine	7649991448168
Sensopro Tube Azurblau/Azure blue	7649991448175
Sensopro Tube Korall-Rot/coral-red	7649991448182
Sensopro Seat Physio	7649991448120
Sensopro Video Kit Montagebalken Physio	7649991448137

Intended Purpose: Treatment of deficits in postural balance in order to prevent falls.

Risk Class: Class I

Rule (according to Annex VIII): Rule 1

meet all applicable provisions of the Regulation (EU) 2017/745 (EU MDR).

Authorized representative: MT Promedt Consulting GmbH, Altenhofstrasse 80, 66386 St. Ingbert, Germany, SRN DE-AR-000000085

Applied common specifications: n.a.

Conformity assessment procedure: EU MDR Art. 52(7)

Florian Kuchen

Florian Kuchen, PRRC