

# EU- Declaration of conformity for medical devices

(According to Regulation (EU) 2017/745 for medical devices, Annex VIII of May 2017)

**Manufacturer:** HAIDER BIOSWING GmbH  
**SRN:** DE-MF-000008765

We hereby declare under our sole responsibility that the sensorimotor therapy device POSTUROMED®

**Type:** BIOSWING® POSTUROMED® compact

**Basic-UDI-DI:** 4260742520020

**Serial number:** 2021: 2521000000 to 2521999999  
2022: 2522000000 to 2522999999  
(further ongoing)

complies with the relevant provisions of the following directive:

## Regulation (EU) 2017/745 for medical devices

Classification according to Regulation (EU) 2017/745 for medical devices, Annex VIII:

**Class I non-active medical device,**

according to classification rule 1



95704 Pullenreuth, Germany, May 26th, 2021

Management

A handwritten signature in blue ink, appearing to read 'E. Haider', written over a dotted line.

Eduard Haider

Technical Manager

A handwritten signature in blue ink, appearing to read 'A. Haider', written over a dotted line.

Dipl.-Ing. Alexander Haider

BIOSWING Therapiesysteme

