**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 PROPRIOMED ®

**Type:** BIOSWING® PROPRIOMED ® 100

**Basic-UDI-DI:** 4260742520051

**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 Technical Manager



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Eduard Haider Dipl.-Ing. Alexander Haider