EU- Declaration of conformity for medical devices

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

www.bioswing.de

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

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

Type:	BIOSWING® PROPRIOMED ® 2
Basic-UDI-DI:	4260742520044
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

CE

95704 Pullenreuth, Germany, May 26th, 2021

Management

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Eduard Haider

Technical Manager

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Dipl.-Ing. Alexander Haider

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