VAST.REHAB
REHABILITATION WITH BIOFEEDBACK

CE

Declaration of Conformity

Manufacturers Name:	BRONTES PROCESSING SP. Z O.O.
Manufacturers Address:	Ul. Przewozowa 32 44-100 Gliwice Poland
Basic UDI-DI:	59044968178VREU
Name of the Device:	VAST.REHAB
Classification:	Class I (MDR, Annex VIII Rule 11)
Conformity assessment route:	Manufacturer uses the following procedures for the CE-labeling of their products according the Regulation MDR 2017/745: Class I: EC conformity declaration according to art. 52 para. 7 + annex II + annex III.

This declaration of conformity is issued under the sole responsibility of BRONTES PROCESSING SP. Z O.O.

We hereby declare that he medical device(s) specified above meet the provision of the Regulation (EU) MDR 2017/745 for medical devices. This declaration is supported by the Quality System approval to ISO 13485:2016.

Andrzej Czech, CEO

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Gliwice, 26 October 2021