

## **Declaration of Conformity**

ID:BFB

Products: Biofeedback trainer MITSAR-BFB

Classification: class IIa (MDD, Annex IX)

Manufacturer: MITSAR Co. Ltd

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Confirmation is hereby given that the Biofeedback trainer MITSAR-BFBconforms to the essential requirements laid down in the Council Directive regarding the harmonization of statutory requirements of the member states on Medical Devices 93/42/EEC (MDD, Annex II) as amended by Directive 2007/47/EC concerning medical devices.

The Quality Management System meets the requirements of ISO 13485.



## The systemconforms to following standards:

MDD 93/42/EEC Medical Device Directive

EN ISO 14971:2012 Medical devices -

Application of risk management to medical devices

EN 60601-1:2006 Medical electrical equipment -- Part 1: General requirements

for basic safety and essential performance

EN 60601-1-2:2015 Medical electrical equipment. General requirements for basic

safety and essential performance. Collateral Standard. Electromagnetic disturbances. Requirements and tests

EN 60601-2-26:2003 Medical electrical equipment -- Part 2-26: Particular

requirements for the safety of electroencephalographs

Notified Body: Eurofins Expert Services Ltd

(previous name VTT Expert Services Ltd) Medical Devices, Notified Body No. 0537 Hermiankatu 6-8 H, FI-33720 Tampere, Finland P.O.BOX 345, FI-33101 Tampere, Finland

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Conformity Assessment

procedure for CE:

Council Directive 93/42/EEC (MDD, Annex II)

Certificate number: C-01-1121-734-20

Date of issue: 28.04.2020

Valid until: 28.05.2024

The declaration of conformity is issued under the sole responsibility of MITSAR Co.Ltd.

Signed for and on behalf of manufacturer:

Date: 11.05.2020

St. Petersburg

Alexander A. Grushvitskiy

CEO «MITSAR Co. Ltd» Tere