

EU Declaration of Conformity

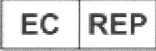
We hereby declare under our sole responsibility that the EvalTech system meets the relevant provisions of the following European Union Directives:

- Council **Directive 93/42/EEC** of 14 June 1993 concerning medical devices as amended by Directive 2007/47/EC (**MDD**)
- **Directive 2006/42/EC** of the European Parliament and of the Council of 17 May 2006 on machinery as amended by Regulation (EU) 2019/1243
- **Directive 2011/65/EU** of the European Parliament and of the Council of 8 June 2011 on the restriction of the use of certain hazardous substances in electrical and electronic equipment (**RoHS**)
- **Directive 2014/53/EU** of the European Parliament and of the Council of 16 April 2014 on radio equipment (Radio Equipment Directive)

The EvalTech has undergone a conformity assessment procedure required by the MDD and is manufactured in harmony with the Technical Documentation compiled as defined in the relevant Directives and retained by BTE.

Product information in regard to the **MDD** and **RoHS** Directives:

Manufacturer 	BTE Technologies 7455-L New Ridge Road Hanover, MD 21076, USA www.btetechnologies.com	Telephone: 410.850.0333 Email: Service@btetechnologies.com
Product Identification	Device Trade Name: EvalTech Device Name: ER Model: EvalTech	
UDI-DI	10850390007328	
EMDN (CND) code	Z120616 - PHYSICAL THERAPY AND REHABILITATION SYSTEMS	
Intended Purpose	The EvalTech is intended to be used for musculoskeletal testing. Applications include occupational and physical therapy and industrial rehabilitation.	
Device Classification (MDD)	Class I	
Classification Rule (MDD)	Rule 12	

Route to Compliance (MDD)	Annex VII of the Medical Devices Directive
Device Classification (MDR)	Class IIa
Classification Rule (MDR)	Rule 11
CE Marking Provision	<p>Under Medical Device Regulation (EU) 2017/745 (MDR), the device will be up-classified to class IIa due to changed software classification rules. Based on the MDR Article 120 §3, the EvalTech can be placed on the EU market as a class I device until May 26, 2024 provided that the device</p> <ul style="list-style-type: none"> • will continue to comply with the MDD, • there will be no significant changes in the design and intended purpose, and • the device will comply with the MDR requirements for post market surveillance, vigilance, and registration of economic operators and of devices
Authorized Representative 	<p>Emergo Europe Prinsessegracht 20 2514 AP, The Hague The Netherlands</p> <p>Telephone: +31.70.345.8570 Emails: EmergoEurope@ul.com EmergoVigilance@ul.com</p>

The device is CE marked since 2008.

Signed for on behalf of BTE Technologies



Ewa Kaczanowska
PRRC/Regulatory Manager
BTE Technologies

Hanover, MD

May 20, 2021



The Technology of Human Performance

Addendum to the original Declaration of Conformity

Per 31 January 2023, the address of our EU Authorized Representative as listed on the original DoC has changed.

OLD ADDRESS AUTHORIZED REPRESENTATIVE		
Name of company	Address	Telephone/email
Emergo Europe	Prinsessegracht 20 2514 AP The Hague The Netherlands	+31.70.345.8570 - phone EmergoEurope@ul.com

NEW ADDRESS AUTHORIZED REPRESENTATIVE		
Name of company	Address	Telephone/email
Emergo Europe	Westervoortsedijk 60 6827 AT Arnhem The Netherlands	+31.70.345.8570 - phone EmergoEurope@ul.com

COMPANY REPRESENTATIVE:

Ewa Kaczanowska
Regulatory Manager/PRRC

June 2, 2023