

DECLARATION OF CONFORMITY



We, **TELEMED UAB**
Dariaus ir Gireno str. 42, Vilnius, LT-02189 Lithuania

Declare under our sole responsibility that:

Equipment	Ultrasound scanners	Probes	
	<i>MicrUs EXT-1H</i>	<i>LV8-4L65S-3</i>	<i>MC4-2R20S-3</i>
		<i>L12-5L40S-3</i>	<i>MC8-4R20S-3</i>
		<i>C5-2R60S-3</i>	<i>MC10-5R10S-3</i>
		<i>L15-6L25S-3</i>	<i>LD12-5L40S-3</i>
		<i>MCV9-5R10S-3</i>	<i>LVD8-4L65S-3</i>
Software	<i>Echo Wave II</i>		
Drivers	<i>TELEMED Drivers Package</i>		

Classification: **Class IIa** (in compliance with Annex II, Art.11 Medical Device Directive)
are in conformity with:

Essential Requirements of Council Directive 93/42/EEC (**Medical Device Directive**)

- IEC 60601-1: 2005, Part 1: General requirements for basic safety and essential performance.
- IEC 60601-1-2: 2007, Part 1: General requirements for basic safety and essential performance, 2.Collateral standard: Electromagnetic compatibility - Requirements and tests
- IEC 60601-2-37:2007 Particular requirements for the basic safety and essential performance of ultrasonic medical diagnostic and monitoring equipment
- ISO-10993-1:2009, Biological Evaluation of Medical Devices, Part 1: Evaluation and Testing within a risk management process.
- ISO 10993-5:2009 Biological evaluation of medical devices -- Part 5: Tests for in vitro cytotoxicity
- ISO-10993-10:2010, Biological Evaluation of Medical Devices, Part 10: Tests for irritation and skin sensitization
- IEC 62304: 2006 Medical device software -- Software life cycle processes
- ISO 14971:2012 Medical devices -- Application of risk management to medical devices

The compliance with the Council Directive 93/42/EEC is under the monitoring of the Notified Body:
MEDCERT GmbH Pilatuspool 2 20355 Hamburg, code: 0482

Date of issue: 12.03.2018

Dmitry Novikov, president