Declaration of Conformity

PRODUCT IDENTIFICATION	
Product name	**Model/number**
ActiGraph	GT9X

MANUFACTURER		
Name of company	**Address**	**Representative**
ActiGraph, LLC	49 East Chase Street Pensacola, Florida 32502 United States of America	Adam Simon Regulatory Affairs Manager

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

CONFORMITY ASSESSMENT		
Device classification	**Route to compliance**	**Standards applied**
Class: 1 Self-Certified (active, non-measuring) Rule: 12 (Annex IX, III. Classification, Page 55)	-Annex VII of Medical Devices Directive (MDD) 93/42/EEC -Radio Equipment Directive (RED) 2014/53/EU	-EN 60601-1-2:2007, C:2010 -EN 61000-4-2:2009 -EN 61000-4-2:2001 -EN 61000-4-3:2006 A1:2008 & A2:2010 -EN 61000-4-8:2010 -EN 55011:2009, A1:2010 -EN 55016-2-3:2010 -EN 55022:2006 -EN 55032:2015 -EN 55016-2-3:2010 -EN 300 328-2 v1.9.1:2015 -EN 301 489-1 V2.1.1:2017 -EN 301 489-3 V1.4.1 -EN 301 489-17 V3.2.0:2017

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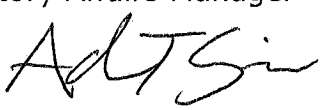
ActiGraph, LLC declares that the above mentioned products meet the provision of the Council Directive 93/42/EEC for Medical Devices.

COMPANY REPRESENTATIVE: Adam Simon

TITLE: Regulatory Affairs Manager

SIGNATURE: 

DATE: 2019-05-13