



Declaration of Conformity

This European Declaration of Conformity is issued under the sole responsibility of the manufacturer.


MANUFACTURER

Name of Company	Address	SRN
ActiGraph L.L.C.	70 North Baylen Street, Suite 400, Pensacola, FL. 32502 United States of America	US-MF-000007954

AUTHORIZED REPRESENTATIVE

Name of Company	Address	SRN	Telephone/fax/email
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PRODUCT IDENTIFICATION

Product Name	Product Code / Catalog Number	Basic UDI-DI
ActiGraph	wGT3X-BT	0853048008008UP
Intended Purpose	Photo	
The ActiGraph activity monitors are small wrist-worn monitors designed for documenting physical movement associated with applications in physiological monitoring. The devices are intended to monitor the activity associated with movement during sleep. The ActiGraph activity monitors can be used to analyze circadian rhythms and assess activity in any instance where quantifiable analysis of physical motion is desirable.		



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RISK CLASS FOR MEDICAL DEVICES	
Device Classification	Common Specifications
Class: I, Self-Certified (active, non-measuring) Rules: 13 (Annex VIII)	-EN 60601-1 -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

Actigraph declares that the above-mentioned products meet the provision of the following EU legislation:

- Medical Device Regulation, MDR (EU) 2017/745
- Radio Equipment Directive (RED) 2014/53/EU
- Restriction of the use of certain hazardous substances (RoHS) Directive 2011/65/EU

COMPANY REPRESENTATIVE: Thomas Hartshorn

TITLE: Head of Quality Assurance & Compliance

SIGNATURE:

PLACE: Pensacola, FL. United States of America

DATE:

22 Mar 2024