Declaration of Conformity - MotionWatch

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

MANUFACTURER		AUTHORISED REPRESENTATIVE	
Name of Company	**Address**	**Name of Company**	**Address**
CamNtech Limited	Riband House Manor Farm Fenstanton Cambridgeshire PE28 9JD, UK	Qarad EC-REP BV	Pas 257 2440 Geel Belgium
SRN: TBA		**SRN:** BE-AR-000000040	

PRODUCT IDENTIFICATION		
Product Name	**Catalogue No.**	**Basic UDI-DI**
MotionWatch 8	04-111	++B329MTN012J
MotionWatch R (Wrist)	04-121	
MotionWatch R (Loop)	04-125	
Intended Purpose		**Representative Photos**
The MotionWatch is a compact, lightweight, body-worn activity monitoring device that may be used to document physical movement associated with applications in physiological monitoring. The device is intended to monitor limb or body movements during daily living and sleep. The MotionWatch can be used to assess activity in any instance where quantifiable analysis of physical motion is desired.		 MotionWatch 8  MotionWatch R

RISK CLASS FOR MEDICAL DEVICES
Class 1 (active, non-measuring, non-sterile) Annex VIII of regulation 2017/745; rules 1, 13 Conformity Assessment: Annex II and Annex III of regulation 2017/745

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

- Medical Device Regulation (MDR) 2017/745
- RoHS Directive 2011/65/EU
- WEEE Directive 2012/19/EU

COMPANY REPRESENTATIVE: Howard Smith

TITLE: Director **SIGNATURE:**

PLACE: United Kingdom **DATE:** 05/01/2021