
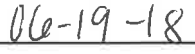


# Declaration of Conformity

<b>Manufacturer:</b> WR Medical Electronics Co.	<b>Address:</b> 1700 Gervais Ave Maplewood, MN 55109 USA
<b>Product Group:</b> Quantitative Sweat Testing Device – Neurological	
<b>Product Family:</b> Autonomic and Sensory Testing Systems	
<b>Device Name:</b> Q-Sweat™	
<b>Product Part Number(s):</b> 5188	
<b>Device Classification Per MDD:</b> Class I – Measuring, per Rule 1	
<b>Year of Manufacture:</b> 2018	
<b>RoHS2 Declaration:</b> The Q-Sweat™ conforms to the Directive 2011/65/EU of the European Parliament and of the Council of 8 June 2011, Restriction of Hazardous Materials (RoHS). Conformance is based on declarations received from our suppliers that the products and raw materials they supply comply with 2011/65/EU and do not contain substances as outlined in Annex II of the directive.	
<b>RoHS2 Declaration Based On:</b> Directive 2011/65/EC	
<b>European Representative:</b> Medical Device Safety Service GmbH, Schiffgraben 41, 30175 Hannover, Germany	
<b>Notified Body:</b> Intertek Semko AB (0413)	
<b>Declaration:</b> WR Medical Electronics Co. hereby declares that the medical device specified above, to which this declaration relates, is in conformance with the essential requirements of Council Directive 93/42/EEC Medical Device Directive under Annex II (EC Declaration of Conformity; Full Quality Assurance System), and with Swedish National Legislation under LVFS 2003:11.	
<b>Declaration Based On:</b> Annex II of the Device Directive 93/42/EEC for Medical Devices	
<b>Certificate No.:</b> 41314493	<b>Issued by:</b> Intertek Semko AB
<b>Declaration of Conformance Issued By:</b> Ms. Amanda Johnsen, Operations Manager; WR Medical Electronics Co. 1700 Gervais Ave, Maplewood, MN, 55109, USA	
<b>Prepared By:</b> Quality Steering Team	
 _____ (Ms. Amanda Johnsen)	 _____ (Date)
Rev 1.8	