

**DECLARATION OF CONFORMITY**

**Name of Manufacturer** Aidian Oy

**Address** Koivu-Mankkaan tie 6 B, P.O. Box 83, FI-02101 Espoo, Finland

**Devices Concerned**

<b>QuikRead go® CRP System</b>	
<b>QuikRead go® CRP</b>	<b>Cat.No. 135171, 135174, 135175, 133891, 135172, 154057, 135173</b>
<b>QuikRead CRP Control</b>	<b>Cat.No. 68296</b>
<b>QuikRead go CRP Control</b>	<b>Cat. No. 153764</b>
<b>QuikRead go CRP Control High</b>	<b>Cat. No. 137071, 153763</b>
<b>QuikRead Capillaries</b>	<b>Cat.No. 67962</b>
<b>QuikRead Plungers</b>	<b>Cat.No. 67966</b>

**Declaration** We hereby declare that the above-mentioned devices comply with the In Vitro Diagnostic Medical Devices Directive 98/79/EC

Classification: General / Other IVD

**Date of Issue** February 4, 2020

**Aidian Oy**



Juho Himberg  
President



p.p. Niina Vuorikallas  
Vice President, Quality Management