

AliveCor, Inc.

KardiaMobile System

**DECLARATION OF CONFORMITY
MEDICAL DEVICES**

We, AliveCor, Inc, hereby declare that the products listed below are in conformity with all relevant provisions of Council Directive 93/42/EEC, as amended September 21, 2007 (M5), concerning Medical Devices.

This Declaration of Conformity is made under Annex II, excluding Section 4, of this directive according to EC Conformity Certificate No. G1171197418005, issued on January 15, 2018 and delivered by TÜV located at TÜV SÜD Product Service GmbH, Ridlerstraße 65, 80339 München, Germany (Notified Body No. 0123).

In addition, we ensure and declare that the distributed CE marked products, as mentioned and falling within Class IIa, Rule 10, meet the provisions of the EC-Directive which apply to them, including an Authorized Representative. The Authorized Representative is Obelis SA, located at BD General Wahis 53 1030, Brussels Belgium.

This declaration is based on the application of the Quality System approved for the manufacture and final inspection of the products concerned, in accordance with Annex II, excluding Section 4, of Directive 93/42/EEC. This declaration is supported by the Quality System certification based on the harmonized standards EN ISO 13485:2016, certificate number Q50974180008 Rev.00, issued on September 4, 2019 and delivered by TÜV located at TÜV SÜD Product Service GmbH, Ridlerstraße 65, 80339 München, Germany (Notified Body No. 0123).

This declaration covers the KardiaMobile System devices and concerns the following products (GMDN:11413):

KardiaMobile	AC-009
KardiaMobile 6L	AC-019
Kardia App for iOS devices	002001
Kardia App for Android devices	002002
KardiaBand	AC-011
Kardia watch App	015001

This declaration is valid from January 15, 2018 to the time of the expiration of the above reference EC Conformity Certificate for all products described here above, bearing the CE marking and manufactured at the following site(s):

AliveCor, Inc.
444 Castro Street, Suite 600
Mountain View, CA, 94041
USA



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November 2, 2020

Date