

K130401 ALERTWATCH: ORFeb 7, 2014
353 days to decisionK130401 · Product code: **MHX** · Cardiovascular
Source: <https://www.510kdatabase.net/k130401/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Physiological, Patient(with Arrhythmia Detection Or Alarms) (MHX)
Date received	Feb 19, 2013
Decision date	Feb 7, 2014
Days to decision	353 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Alertwatch, LLC
Location	Clarksville, MD, US
Contact	DONNA-BEA TILLMAN
510(k) history	2 submissions · 2 cleared · 2014-2016

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