

**K052240 CARDIONET AMBULATORY ECG MONITOR WITH
ARRHYTHMIA DETECTION, MODEL 1002**Oct 19, 2005
63 days to decisionK052240 · Product code: **QYX** · Cardiovascular
Source: <https://www.510kdatabase.net/k052240/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Outpatient Cardiac Telemetry (QYX)
Date received	Aug 17, 2005
Decision date	Oct 19, 2005
Days to decision	63 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Cardionet, Inc.
Location	San Diego, CA, US
Contact	JACK GAIKWAD
510(k) history	6 submissions · 6 cleared · 2002-2010

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k052240/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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