FDA 510(k) Premarket Notification

CLEARED

K052240 CARDIONET AMBULATORY ECG MONITOR WITH ARRHYTHMIA DETECTION, MODEL 1002

Oct 19, 2005
63 days to decision

K052240 · 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