FDA 510(k) Premarket Notification

K012241 CARDIONET AMBULATORY ECG MONITOR WITH ARRHYTHMIA DETECTION, MODEL CN1001

CLEARED

Feb 1, 2002
199 days to decision

K012241 · Product code: **DSI** · Cardiovascular
Source: https://www.510kdatabase.net/k012241/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Jul 17, 2001
Decision date	Feb 1, 2002
Days to decision	199 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Cardionet, Inc.**
Location	San Diego, CA, US
Contact	DONALD V CANAL
510(k) history	6 submissions · 6 cleared · 2002-2010