

**K012241 CARDIONET AMBULATORY ECG MONITOR WITH  
ARRHYTHMIA DETECTION, MODEL CN1001**

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k012241/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 21, 2026