

**K871251 CIRCAMED FULL DISCLOSURE SYSTEM**May 22, 1987  
53 days to decisionK871251 · Product code: **DSI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k871251/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Mar 30, 1987
Decision date	May 22, 1987
Days to decision	53 days
Third-party review	No

**APPLICANT**

---

Company	<b>Circadian, Inc.</b>
Location	Mchenry, IL, US
Contact	MATT S MAGOUN
510(k) history	13 submissions · 13 cleared · 1980-1992

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k871251/>, Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 20, 2026