

**K871572 RHYTHMSCAN**Feb 22, 1988  
306 days to decisionK871572 · Product code: **DSI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k871572/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Apr 22, 1987
Decision date	Feb 22, 1988
Days to decision	306 days
Third-party review	No

**APPLICANT**

---

Company	<b>Brentwood Instruments, Inc.</b>
Location	Mchenry, IL, US
Contact	JAMES OBERMAYER
510(k) history	21 submissions · 20 cleared · 1981-1990

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k871572/>, 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 26, 2026