

**K872249 SIRECUST MODEL 960 AND 961 PATIENT MONITORING**Jul 21, 1987  
39 days to decisionK872249 · Product code: **DRT** · Cardiovascular  
Source: <https://www.510kdatabase.net/k872249/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Jun 12, 1987
Decision date	Jul 21, 1987
Days to decision	39 days
Third-party review	No

**APPLICANT**

---

Company	<b>Medical Electronics Laboratories, Inc.</b>
Location	Danvers, MA, US
Contact	MARVIN ROSENBAUM
510(k) history	2 submissions · 2 cleared · 1987-1988

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k872249/>; 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 July 7, 2026