

**K914576 KELLER CARDIAC MONITOR MEDIL KMS870**Dec 30, 1991  
76 days to decisionK914576 · Product code: **DRT** · Cardiovascular  
Source: <https://www.510kdatabase.net/k914576/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiometer & Rate Alarm) (DRT)
Date received	Oct 15, 1991
Decision date	Dec 30, 1991
Days to decision	76 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

---

Company	<b>Keller Medical Specialties Products, Inc.</b>
Location	Antioch, IL, US
Contact	JAY KELLER
510(k) history	6 submissions · 6 cleared · 1990-1999

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k914576/> 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 25, 2026