

K884645 PATIENT MONITOR RIGEL MODEL MULTICARE 429Feb 2, 1989
87 days to decisionK884645 · Product code: **DRT** · Cardiovascular
Source: <https://www.510kdatabase.net/k884645/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Nov 7, 1988
Decision date	Feb 2, 1989
Days to decision	87 days
Third-party review	No

APPLICANT

Company	Rigel Medical Electronics, Inc.
Location	Mill Valley, CA, US
Contact	MICHAEL E FEWER
510(k) history	6 submissions · 6 cleared · 1984-1989

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k884645/>; 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 July 7, 2026