

K812145 CARDIORATER CR-3Sep 25, 1981
59 days to decisionK812145 · Product code: **DRT** · CardiovascularSource: <https://www.510kdatabase.net/k812145/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Jul 28, 1981
Decision date	Sep 25, 1981
Days to decision	59 days
Third-party review	No

APPLICANT

Company	Life Science Instrumentation, Inc.
Location	Mchenry, IL, US
510(k) history	20 submissions · 20 cleared · 1981-1985

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k812145/> 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 8, 2026