

K821223 CR-9Jun 17, 1982
51 days to decisionK821223 · Product code: **DRT** · Cardiovascular
Source: <https://www.510kdatabase.net/k821223/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Apr 27, 1982
Decision date	Jun 17, 1982
Days to decision	51 days
Third-party review	No

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k821223/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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