FDA 510(k) Premarket Notification

K821223 CR-9

CLEARED

Jun 17, 1982
51 days to decision

K821223 · 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