FDA 510(k) Premarket Notification

K822390 LIFE TRACE 24

CLEARED

Sep 21, 1982
43 days to decision

K822390 · Product code: **DRT** · Cardiovascular
Source: https://www.510kdatabase.net/k822390/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Aug 9, 1982
Decision date	Sep 21, 1982
Days to decision	43 days
Third-party review	No

APPLICANT

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