FDA 510(k) Premarket Notification

K822388 LIFE TRACE 14

CLEARED

Aug 24, 1982
15 days to decision

K822388 · Product code: **DRT** · Cardiovascular
Source: https://www.510kdatabase.net/k822388/

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	Aug 24, 1982
Days to decision	15 days
Third-party review	No

APPLICANT

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