

K822390 LIFE TRACE 24Sep 21, 1982
43 days to decisionK822390 · Product code: **DRT** · CardiovascularSource: <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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k822390/> 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