

K852884 MEDILOG M1Jul 26, 1985
17 days to decisionK852884 · Product code: **DSI** · Cardiovascular
Source: <https://www.510kdatabase.net/k852884/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Jul 9, 1985
Decision date	Jul 26, 1985
Days to decision	17 days
Third-party review	No

APPLICANT

Company	Oxford Medilog, Inc.
Location	Mchenry, IL, US
Contact	JOHN K LAURIE
510(k) history	48 submissions · 48 cleared · 1978-1994

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k852884/>; 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 May 21, 2026