FDA 510(k) Premarket Notification

K852884 MEDILOG M1

CLEARED

Jul 26, 1985
17 days to decision

K852884 · 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