FDA 510(k) Premarket Notification

K890330 MEDILOG 6000-FD REPORT GENERATOR

CLEARED

Apr 20, 1989
87 days to decision

K890330 · Product code: **DSI** · Cardiovascular
Source: https://www.510kdatabase.net/k890330/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Jan 23, 1989
Decision date	Apr 20, 1989
Days to decision	87 days
Third-party review	No

APPLICANT

Company	**Oxford Medilog, Inc.**
Location	Mchenry, IL, US
Contact	MICHAEL J SMITH
510(k) history	48 submissions · 48 cleared · 1978-1994