

K890330 MEDILOG 6000-FD REPORT GENERATORApr 20, 1989
87 days to decisionK890330 · 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

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