

K833162 AMBULATORY MONITORING DEVICEJan 10, 1984
116 days to decisionK833162 · Product code: **DSI** · Cardiovascular
Source: <https://www.510kdatabase.net/k833162/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Sep 16, 1983
Decision date	Jan 10, 1984
Days to decision	116 days
Third-party review	No

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k833162/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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