

**K802033 OXFORD INTENSIVE CARE PATIENT MON. SYS**Sep 26, 1980  
35 days to decisionK802033 · Product code: **DSI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k802033/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Aug 22, 1980
Decision date	Sep 26, 1980
Days to decision	35 days
Third-party review	No

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k802033/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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