

K790287 ANALYZER, MEDILOG IIFeb 28, 1979
20 days to decisionK790287 · Product code: **DSI** · CardiovascularSource: <https://www.510kdatabase.net/k790287/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Feb 8, 1979
Decision date	Feb 28, 1979
Days to decision	20 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k790287/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 21, 2026