

K822903 AUTOECHONov 1, 1982
34 days to decisionK822903 · Product code: **DXG** · CardiovascularSource: <https://www.510kdatabase.net/k822903/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Computer, Diagnostic, Pre-programmed, Single-function (DXG)
Date received	Sep 28, 1982
Decision date	Nov 1, 1982
Days to decision	34 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/k822903/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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