FDA 510(k) Premarket Notification

K822903 AUTOECHO

CLEARED

Nov 1, 1982
34 days to decision

K822903 · Product code: **DXG** · Cardiovascular
Source: 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