

**K901592 MODEL MD-5 DOPPLER**Oct 22, 1990  
200 days to decisionK901592 · Product code: **DXK** · CardiovascularSource: <https://www.510kdatabase.net/k901592/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Echocardiograph (DXK)
Date received	Apr 5, 1990
Decision date	Oct 22, 1990
Days to decision	200 days
Third-party review	No

**APPLICANT**

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Company	<b>D. E. Hokanson, Inc.</b>
Location	Mchenry, IL, US
Contact	EUGENE HOKANSON
510(k) history	19 submissions · 19 cleared · 1979-2004

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k901592/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 18, 2026