

**K903710 WASHINGTON CARDIAC MONITOR**Nov 9, 1990  
87 days to decisionK903710 · Product code: **DXK** · CardiovascularSource: <https://www.510kdatabase.net/k903710/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Echocardiograph (DXK)
Date received	Aug 14, 1990
Decision date	Nov 9, 1990
Days to decision	87 days
Third-party review	No

**APPLICANT**

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Company	<b>Washington Biolab, Inc.</b>
Location	Ames, IA, US
Contact	KI, PH.D.
510(k) history	1 submissions · 1 cleared · 1990-1990

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k903710/>, 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 July 6, 2026