

**K933603 LEADWIRE**Jan 18, 1994  
175 days to decisionK933603 · Product code: **DSA** · CardiovascularSource: <https://www.510kdatabase.net/k933603/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cable, Transducer And Electrode, Patient, (including Connector) (DSA)
Date received	Jul 27, 1993
Decision date	Jan 18, 1994
Days to decision	175 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Sentry Medical Products, Inc.</b>
Location	Walker, MI, US
Contact	WILLIAM E BLAIR
510(k) history	13 submissions · 13 cleared · 1983-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k933603/> 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 1, 2026