

**K901251 MODEL 9101 RECORDER/ANALYZER SYSTEM**May 22, 1990  
67 days to decisionK901251 · Product code: **DSH** · CardiovascularSource: <https://www.510kdatabase.net/k901251/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Recorder, Magnetic Tape, Medical (DSH)
Date received	Mar 16, 1990
Decision date	May 22, 1990
Days to decision	67 days
Third-party review	No

**APPLICANT**

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Company	<b>Aequitron Medical, Inc.</b>
Location	Walker, MI, US
Contact	KENNETH B HERLAND
510(k) history	24 submissions · 22 cleared · 1984-1997

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k901251/>, 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 25, 2026