

**K903940 TYPE 9210 S/ECG**Dec 4, 1991  
464 days to decisionK903940 · Product code: **DSI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k903940/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Aug 27, 1990
Decision date	Dec 4, 1991
Days to decision	464 days
Third-party review	No

**APPLICANT**

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Company	<b>S &amp; W Medico Teknik</b>
Location	Walker, MI, US
Contact	METTE JORGENSEN
510(k) history	46 submissions · 46 cleared · 1984-1995

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k903940/>, 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 26, 2026