

**K884106 SUPERVISORY TELEMETRY ARRHYTHMIA TERMINAL**Mar 6, 1989  
158 days to decisionK884106 · Product code: **DSI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k884106/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Sep 29, 1988
Decision date	Mar 6, 1989
Days to decision	158 days
Third-party review	No

**APPLICANT**

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Company	<b>Siemens Medical Solutions USA, Inc.</b>
Location	Hoffman Estates, IL, US
Contact	HARRY K SCHWILL
510(k) history	778 submissions · 778 cleared · 1980-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k884106/>; 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 15, 2026