

**K881273 HORIZON 2000 ARRHYTHMIA MONITORING OPTION**Jul 12, 1988  
109 days to decisionK881273 · Product code: **DSI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k881273/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Mar 25, 1988
Decision date	Jul 12, 1988
Days to decision	109 days
Third-party review	No

**APPLICANT**

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Company	<b>Mennen Medical, Inc.</b>
Location	Clarence, NY, US
Contact	THOMAS CONNELLY
510(k) history	34 submissions · 34 cleared · 1985-2004

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k881273/>; 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 21, 2026