

**K843503 AEGIS AMBULATORY MONITORING SYSTEM**Oct 26, 1984  
50 days to decisionK843503 · Product code: **DSI** · Cardiovascular  
Source: <https://www.510kdatabase.net/k843503/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Detector And Alarm, Arrhythmia (DSI)
Date received	Sep 6, 1984
Decision date	Oct 26, 1984
Days to decision	50 days
Third-party review	No

**APPLICANT**

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Company	<b>Aegis Medical Systems, Inc.</b>
Location	Marlton, NJ, US
Contact	JOHN R MORENO
510(k) history	1 submissions · 1 cleared · 1984-1984

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k843503/>; 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 8, 2026