

**K960849 MAGNO ALARM**Mar 26, 1997  
400 days to decisionK960849 · Product code: **DTG** · Cardiovascular  
Source: <https://www.510kdatabase.net/k960849/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Magnet, Test, Pacemaker (DTG)
Date received	Feb 20, 1996
Decision date	Mar 26, 1997
Days to decision	400 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Cardiac Assist Devices, Inc.</b>
Location	Cleveland, OH, US
Contact	RASSOLL RASHIDI
510(k) history	3 submissions · 3 cleared · 1997-2002

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k960849/> 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 3, 2026