

**K031987 POWERHEART AED G3, MODEL 9300**Jul 30, 2003  
33 days to decisionK031987 · Product code: **MKJ** · Cardiovascular  
Source: <https://www.510kdatabase.net/k031987/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Jun 27, 2003
Decision date	Jul 30, 2003
Days to decision	33 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Cardiac Science</b>
Location	Edina, MN, US
Contact	KENNETH OLSON
510(k) history	1 submissions · 1 cleared · 2003-2003

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k031987/> 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