

**K092364 MODEL 220 PATIENT MAGNET**Nov 3, 2009  
90 days to decisionK092364 · Product code: **DTG** · Cardiovascular  
Source: <https://www.510kdatabase.net/k092364/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Magnet, Test, Pacemaker (DTG)
Date received	Aug 5, 2009
Decision date	Nov 3, 2009
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Cyberonics, Inc.</b>
Location	Houston, TX, US
Contact	KAREN STUEBER
510(k) history	1 submissions · 1 cleared · 2009-2009

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k092364/> 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 June 22, 2026