

**K090249 MODIFICATION TO: MICRO-INDUCTION 1000 SYSTEM**Apr 3, 2009  
60 days to decisionK090249 · Product code: **DPS** · Cardiovascular  
Source: <https://www.510kdatabase.net/k090249/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrocardiograph (DPS)
Date received	Feb 2, 2009
Decision date	Apr 3, 2009
Days to decision	60 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Harbinger Medical, Inc.</b>
Location	Eden Prairie, MN, US
Contact	HAROLD HOIUM
510(k) history	2 submissions · 2 cleared · 2002-2009

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k090249/> 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 23, 2026