

**K041340 GUIDANT MICROWAVE ABLATION SYSTEM**Jul 28, 2004  
69 days to decisionK041340 · Product code: **NEY** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k041340/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Ablation, Microwave And Accessories (NEY)
Date received	May 20, 2004
Decision date	Jul 28, 2004
Days to decision	69 days
Third-party review	Yes
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Guidant Corporation, Cardiac Surgery</b>
Location	S,Mta Clara, CA, US
Contact	NANCY GALLO
510(k) history	8 submissions · 8 cleared · 2002-2008

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k041340/> 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 25, 2026