

**K101242 GENESIS DM MODEL: 6053000A1**Jun 11, 2010  
39 days to decisionK101242 · Product code: **DRG** · Cardiovascular  
Source: <https://www.510kdatabase.net/k101242/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Transmitters And Receivers, Physiological Signal, Radiofrequency (DRG)
Date received	May 3, 2010
Decision date	Jun 11, 2010
Days to decision	39 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Honeywell Hommed, LLC</b>
Location	Houston, TX, US
Contact	EMILY HEI
510(k) history	7 submissions · 7 cleared · 2006-2016

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k101242/> 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 2, 2026