

**K153719 LifeStream Management System, LifeStream Management Suite, LifeStream Patient Manager, LifeStream Manager, LifeStream Web Manager, LifeStream**Jun 3, 2016  
158 days to decisionK153719 · Product code: **DRG** · Cardiovascular  
Source: <https://www.510kdatabase.net/k153719/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transmitters And Receivers, Physiological Signal, Radiofrequency (DRG)
Date received	Dec 28, 2015
Decision date	Jun 3, 2016
Days to decision	158 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Honeywell Hommed, LLC</b>
Location	Houston, TX, US
Contact	Greg Lillegard
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/k153719/> 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