

**K052767 AFIBALERT ATRIAL FIBRILLATION DETECTOR**Jun 23, 2006  
266 days to decisionK052767 · Product code: **DXH** · CardiovascularSource: <https://www.510kdatabase.net/k052767/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transmitters And Receivers, Electrocardiograph, Telephone (DXH)
Date received	Sep 30, 2005
Decision date	Jun 23, 2006
Days to decision	266 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Lechnologies Research, Inc.</b>
Location	Sussex, WI, US
Contact	TIM LOHMAN
510(k) history	1 submissions · 1 cleared · 2006-2006

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k052767/> 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 30, 2026