

**K072385 HEARTEVENT, MODEL AT**Jan 11, 2008  
140 days to decisionK072385 · Product code: **DXH** · Cardiovascular  
Source: <https://www.510kdatabase.net/k072385/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transmitters And Receivers, Electrocardiograph, Telephone (DXH)
Date received	Aug 24, 2007
Decision date	Jan 11, 2008
Days to decision	140 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Diagnostic Devices Pty, Ltd.</b>
Location	Eastgardens, AU
Contact	HARRY PLATT
510(k) history	1 submissions · 1 cleared · 2008-2008

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k072385/> 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