

**K890461 CARDIOBEEPER IV**Feb 23, 1989  
24 days to decisionK890461 · Product code: **DXH** · CardiovascularSource: <https://www.510kdatabase.net/k890461/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transmitters And Receivers, Electrocardiograph, Telephone (DXH)
Date received	Jan 30, 1989
Decision date	Feb 23, 1989
Days to decision	24 days
Third-party review	No

**APPLICANT**

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Company	<b>Survival Technology, Inc.</b>
Location	Mchenry, IL, US
Contact	GARY W LEYLAND
510(k) history	10 submissions · 10 cleared · 1977-1993

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k890461/>, 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 July 6, 2026