

**K844966 CARDIOBEEPER PROFILE MONITOR**Feb 8, 1985  
46 days to decisionK844966 · Product code: **DXH** · CardiovascularSource: <https://www.510kdatabase.net/k844966/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transmitters And Receivers, Electrocardiograph, Telephone (DXH)
Date received	Dec 24, 1984
Decision date	Feb 8, 1985
Days to decision	46 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/k844966/>; 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