

**K925535 CARDIOBEEPER(R) CB-IIA**Jan 28, 1993  
87 days to decisionK925535 · Product code: **DXH** · CardiovascularSource: <https://www.510kdatabase.net/k925535/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transmitters And Receivers, Electrocardiograph, Telephone (DXH)
Date received	Nov 2, 1992
Decision date	Jan 28, 1993
Days to decision	87 days
Third-party review	No
Summary / Statement	Summary

**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/k925535/> 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