

**K965101 CARDIOBEEPER CB-12L**Mar 17, 1997  
87 days to decisionK965101 · Product code: **DXH** · Cardiovascular  
Source: <https://www.510kdatabase.net/k965101/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transmitters And Receivers, Electrocardiograph, Telephone (DXH)
Date received	Dec 20, 1996
Decision date	Mar 17, 1997
Days to decision	87 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Meridian Medical Technologies, Inc.</b>
Location	Columbia, MD, US
Contact	SHEILA HEMEON-HEYER
510(k) history	3 submissions · 3 cleared · 1997-2003

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