FDA 510(k) Premarket Notification

K925535 CARDIOBEEPER(R) CB-IIA

CLEARED

Jan 28, 1993
87 days to decision

K925535 · Product code: **DXH** · Cardiovascular
Source: https://www.510kdatabase.net/k925535/

SUBMISSION DETAILS

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

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