FDA 510(k) Premarket Notification

K844966 CARDIOBEEPER PROFILE MONITOR

CLEARED

Feb 8, 1985
46 days to decision

K844966 · Product code: **DXH** · Cardiovascular
Source: https://www.510kdatabase.net/k844966/

SUBMISSION DETAILS

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

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