FDA 510(k) Premarket Notification

K813101 CARDIOBEEPER MEMORY MONITOR

CLEARED

Nov 27, 1981
24 days to decision

K813101 · Product code: **DXY** · Cardiovascular
Source: https://www.510kdatabase.net/k813101/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implantable Pacemaker Pulse-generator (DXY)
Date received	Nov 3, 1981
Decision date	Nov 27, 1981
Days to decision	24 days
Third-party review	No

APPLICANT

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