

K813101 CARDIOBEEPER MEMORY MONITORNov 27, 1981
24 days to decisionK813101 · Product code: **DXY** · CardiovascularSource: <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

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Device record: <https://www.510kdatabase.net/k813101/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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