

K840353 PACEMAKER PULSE GENERATOR W/ELECTRODEApr 23, 1984
88 days to decisionK840353 · Product code: **DRO** · CardiovascularSource: <https://www.510kdatabase.net/k840353/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pacemaker, Cardiac, External Transcutaneous (non-invasive) (DRO)
Date received	Jan 26, 1984
Decision date	Apr 23, 1984
Days to decision	88 days
Third-party review	No

APPLICANT

Company	Ross Research, Inc.
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1982-1984

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Device record: <https://www.510kdatabase.net/k840353/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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