

K822246 NONINVASIVE TEMPORARY PACEMAKERAug 6, 1982
8 days to decisionK822246 · Product code: **DRO** · CardiovascularSource: <https://www.510kdatabase.net/k822246/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pacemaker, Cardiac, External Transcutaneous (non-invasive) (DRO)
Date received	Jul 29, 1982
Decision date	Aug 6, 1982
Days to decision	8 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/k822246/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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