

K820507 PACE* AID MODEL 50Aug 6, 1982
162 days to decisionK820507 · Product code: **DRO** · CardiovascularSource: <https://www.510kdatabase.net/k820507/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pacemaker, Cardiac, External Transcutaneous (non-invasive) (DRO)
Date received	Feb 25, 1982
Decision date	Aug 6, 1982
Days to decision	162 days
Third-party review	No

APPLICANT

Company	Cardiac Resucitator Corp.
Location	Mchenry, IL, US
510(k) history	17 submissions · 16 cleared · 1981-1988

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Device record: <https://www.510kdatabase.net/k820507/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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