

K821673 HEART*AID, ALS, MODEL 95Aug 6, 1982
60 days to decisionK821673 · Product code: **DRO** · CardiovascularSource: <https://www.510kdatabase.net/k821673/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pacemaker, Cardiac, External Transcutaneous (non-invasive) (DRO)
Date received	Jun 7, 1982
Decision date	Aug 6, 1982
Days to decision	60 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/k821673/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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