FDA 510(k) Premarket Notification

K871971 MODIFIED PACE*AID MODEL 53

CLEARED

Jun 24, 1987
34 days to decision

K871971 · Product code: **DRO** · Cardiovascular
Source: https://www.510kdatabase.net/k871971/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pacemaker, Cardiac, External Transcutaneous (non-invasive) (DRO)
Date received	May 21, 1987
Decision date	Jun 24, 1987
Days to decision	34 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k871971/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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