

K934067 POWRPAK REPLACEMENT BATTERIES FOR PHISIO-CONTROL LIFEPAK 5/10

Oct 7, 1993
48 days to decision

K934067 · Product code: DRK · Cardiovascular
Source: <https://www.510kdatabase.net/k934067/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, High Energy, (including Paddles) (DRK)
Date received	Aug 20, 1993
Decision date	Oct 7, 1993
Days to decision	48 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	National Custom Ent., Inc.
Location	Burnsville, MN, US
Contact	MIKE BENYO
510(k) history	12 submissions · 11 cleared · 1993-2001

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k934067/>. Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 2, 2026