

K840900 PORTA PAK 90Jun 11, 1984
104 days to decisionK840900 · Product code: **LDD** · CardiovascularSource: <https://www.510kdatabase.net/k840900/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Feb 28, 1984
Decision date	Jun 11, 1984
Days to decision	104 days
Third-party review	No

APPLICANT

Company	Medical Research Laboratories, Inc.
Location	Mchenry, IL, US
510(k) history	19 submissions · 15 cleared · 1981-2002

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Device record: <https://www.510kdatabase.net/k840900/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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