

K833751 PORTA PULSE III #D320Jan 11, 1984
77 days to decisionK833751 · Product code: **LDD** · CardiovascularSource: <https://www.510kdatabase.net/k833751/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Oct 26, 1983
Decision date	Jan 11, 1984
Days to decision	77 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k833751/> 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 May 26, 2026