FDA 510(k) Premarket Notification

K833751 PORTA PULSE III #D320

CLEARED

Jan 11, 1984
77 days to decision

K833751 · Product code: **LDD** · Cardiovascular
Source: 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