FDA 510(k) Premarket Notification

K840900 PORTA PAK 90

CLEARED

Jun 11, 1984
104 days to decision

K840900 · Product code: **LDD** · Cardiovascular
Source: 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