FDA 510(k) Premarket Notification

K890079 LIFEPAK 10 DEFIBRILLATOR/MONITOR

CLEARED

Feb 23, 1989
50 days to decision

K890079 · Product code: **LDD** · Cardiovascular
Source: https://www.510kdatabase.net/k890079/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Jan 4, 1989
Decision date	Feb 23, 1989
Days to decision	50 days
Third-party review	No

APPLICANT

Company	**Physio-Control Corp.**
Location	Mchenry, IL, US
Contact	SHERRI L POCOCK
510(k) history	80 submissions · 78 cleared · 1976-1999