FDA 510(k) Premarket Notification

K935674 LIFEPAK 9 DEFIBRILLATOR/MONITOR

CLEARED

Dec 16, 1993
8 days to decision

K935674 · Product code: **LDD** · Cardiovascular
Source: https://www.510kdatabase.net/k935674/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Dec 8, 1993
Decision date	Dec 16, 1993
Days to decision	8 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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