FDA 510(k) Premarket Notification

K905510 LIFEPAK 9 DEFIBRILLATOR & CARDIAC MONITOR/MODIFIED

CLEARED

Dec 20, 1990
13 days to decision

K905510 · Product code: **LDD** · Cardiovascular
Source: https://www.510kdatabase.net/k905510/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Dec 7, 1990
Decision date	Dec 20, 1990
Days to decision	13 days
Third-party review	No

APPLICANT

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