FDA 510(k) Premarket Notification

K902478 LIFEPAK 300 AUTOMATIC ADVISORY DEFIBRILLATOR

CLEARED

Oct 31, 1990
149 days to decision

K902478 · Product code: **LDD** · Cardiovascular
Source: https://www.510kdatabase.net/k902478/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Jun 4, 1990
Decision date	Oct 31, 1990
Days to decision	149 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