FDA 510(k) Premarket Notification

K881153 LIFEPAK 9 DEFIBRILLATOR AND CARDIAC MONITOR

CLEARED

Jul 27, 1988
132 days to decision

K881153 · Product code: **LDD** · Cardiovascular
Source: https://www.510kdatabase.net/k881153/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Mar 17, 1988
Decision date	Jul 27, 1988
Days to decision	132 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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k881153/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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