FDA 510(k) Premarket Notification

K912966 PEDIATRIC DISPOSABLE EXTERNAL DEFIBRILATION/ECG

CLEARED

Feb 21, 1992
228 days to decision

K912966 · Product code: **LDD** · Cardiovascular
Source: https://www.510kdatabase.net/k912966/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Jul 8, 1991
Decision date	Feb 21, 1992
Days to decision	228 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