FDA 510(k) Premarket Notification

K103567 LIFEPAK 15 MONITOR/DEFIBRILLATOR

CLEARED

Mar 22, 2011
106 days to decision

K103567 · Product code: **MKJ** · Cardiovascular
Source: https://www.510kdatabase.net/k103567/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Dec 6, 2010
Decision date	Mar 22, 2011
Days to decision	106 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Physio-Control, Inc.**
Location	Redmond, WA, US
Contact	MICHELLE ACKERMANN
510(k) history	14 submissions · 14 cleared · 1984-2025