FDA 510(k) Premarket Notification

K122600 LIFEPAK 1000 DEFIBRILLATOR

CLEARED

Oct 11, 2012
45 days to decision

K122600 · Product code: **MKJ** · Cardiovascular
Source: https://www.510kdatabase.net/k122600/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Aug 27, 2012
Decision date	Oct 11, 2012
Days to decision	45 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