FDA 510(k) Premarket Notification

K102972 LIFEPAK 12

CLEARED

Dec 22, 2010
77 days to decision

K102972 · Product code: **MKJ** · Cardiovascular
Source: https://www.510kdatabase.net/k102972/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Oct 6, 2010
Decision date	Dec 22, 2010
Days to decision	77 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