

K103567 LIFEPAK 15 MONITOR/DEFIBRILLATORMar 22, 2011
106 days to decisionK103567 · 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

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