

K122600 LIFEPAK 1000 DEFIBRILLATOROct 11, 2012
45 days to decisionK122600 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k122600/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 26, 2026