

**K011144 LIFEPAK 600 AUTOMATED EXTERNAL  
DEFIBRILLATOR**Dec 3, 2001  
231 days to decisionK011144 · Product code: **MKJ** · Cardiovascular  
Source: <https://www.510kdatabase.net/k011144/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Apr 16, 2001
Decision date	Dec 3, 2001
Days to decision	231 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Medtronic Physio-Control Corp.</b>
Location	Redmond, WA, US
Contact	SHERRI L POCOCK
510(k) history	8 submissions · 6 cleared · 2001-2004

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k011144/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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