

**K052057 LIFEPAK 500 AUTOMATED EXTERNAL
DEFIBRILLATOR**Feb 17, 2006
203 days to decisionK052057 · Product code: **MKJ** · Cardiovascular
Source: <https://www.510kdatabase.net/k052057/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Jul 29, 2005
Decision date	Feb 17, 2006
Days to decision	203 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Medtronic Emergency Response Systems, Inc.
Location	Redmond, WA, US
Contact	SHERRI POCOCK
510(k) history	6 submissions · 6 cleared · 2004-2007

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