

K063119 LIFEPAK 20 DEFIBRILLATOR/ MONITORDec 22, 2006
71 days to decisionK063119 · Product code: **MKJ** · Cardiovascular
Source: <https://www.510kdatabase.net/k063119/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Oct 12, 2006
Decision date	Dec 22, 2006
Days to decision	71 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

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