FDA 510(k) Premarket Notification

K063119 LIFEPAK 20 DEFIBRILLATOR/ MONITOR

CLEARED

Dec 22, 2006
71 days to decision

K063119 · 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