FDA 510(k) Premarket Notification

K052057 LIFEPAK 500 AUTOMATED EXTERNAL DEFIBRILLATOR

CLEARED

Feb 17, 2006
203 days to decision

K052057 · 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