

K033275 LIFEPAK 12, 20, 500, CR PLUSNov 6, 2003
27 days to decisionK033275 · Product code: **MKJ** · CardiovascularSource: <https://www.510kdatabase.net/k033275/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Oct 10, 2003
Decision date	Nov 6, 2003
Days to decision	27 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Medtronic Physio-Control Corp.
Location	Redmond, WA, US
Contact	MICHELLE ACKERMANN
510(k) history	8 submissions · 6 cleared · 2001-2004

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