

K003548 PADPRO SYSTEM DEFIBRILLATOR ELECTRODE ADAPTERS

Sep 4, 2001
291 days to decision

K003548 · Product code: **MKJ** · Cardiovascular
Source: <https://www.510kdatabase.net/k003548/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Nov 17, 2000
Decision date	Sep 4, 2001
Days to decision	291 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Padpro, LLC
Location	Deer Field, IL, US
Contact	DANIEL KAMM
510(k) history	5 submissions · 5 cleared · 2001-2002

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k003548/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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