

**K002280 PAD PRO PEDIATRIC ELECTRODE, MODEL PAD PRO  
2602**Feb 5, 2001  
194 days to decisionK002280 · Product code: **MKJ** · Cardiovascular  
Source: <https://www.510kdatabase.net/k002280/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Jul 26, 2000
Decision date	Feb 5, 2001
Days to decision	194 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Medical Devices, Inc.</b>
Location	Mchenry, IL, US
Contact	WARREN R WALTERS
510(k) history	49 submissions · 47 cleared · 1977-2001

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k002280/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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