

**K061606 DEFIBRILLATOR CABLE TESTER, MODEL DT2200**Jun 21, 2006  
12 days to decisionK061606 · Product code: **DRG** · CardiovascularSource: <https://www.510kdatabase.net/k061606/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transmitters And Receivers, Physiological Signal, Radiofrequency (DRG)
Date received	Jun 9, 2006
Decision date	Jun 21, 2006
Days to decision	12 days
Third-party review	Yes
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Medical Devices/Padpro, Inc.</b>
Location	Deer Field, IL, US
Contact	DANIEL KAMM
510(k) history	1 submissions · 1 cleared · 2006-2006

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k061606/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 30, 2026