

**K895729 R2 PACE PAD, MODEL 480**Oct 12, 1989  
17 days to decisionK895729 · Product code: **DRO** · CardiovascularSource: <https://www.510kdatabase.net/k895729/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pacemaker, Cardiac, External Transcutaneous (non-invasive) (DRO)
Date received	Sep 25, 1989
Decision date	Oct 12, 1989
Days to decision	17 days
Third-party review	No

**APPLICANT**

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Company	<b>Darox Corp.</b>
Location	Niles, IL, US
Contact	WILLIAM J SMIRLES
510(k) history	2 submissions · 2 cleared · 1989-1989

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k895729/>; 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 July 6, 2026