

**K960721 PACEART WRIST ELECTRODES**Jul 18, 1996  
147 days to decisionK960721 · Product code: **DRX** · CardiovascularSource: <https://www.510kdatabase.net/k960721/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Feb 22, 1996
Decision date	Jul 18, 1996
Days to decision	147 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Paceart, Inc.</b>
Location	Millburn, NJ, US
Contact	MICHAEL BERGLESON, PH.D.
510(k) history	9 submissions · 9 cleared · 1987-2001

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k960721/> 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 3, 2026