

**K872368 MODIFIED VERSA STIM ELECTRODE LINE**Sep 18, 1987  
92 days to decisionK872368 · Product code: **DRO** · CardiovascularSource: <https://www.510kdatabase.net/k872368/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pacemaker, Cardiac, External Transcutaneous (non-invasive) (DRO)
Date received	Jun 18, 1987
Decision date	Sep 18, 1987
Days to decision	92 days
Third-party review	No

**APPLICANT**

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Company	<b>Conmedcorp</b>
Location	Dayton, OH, US
Contact	WILLIAM W ABRAHAM
510(k) history	92 submissions · 92 cleared · 1981-2010

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k872368/>; 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 May 27, 2026