

**K905489 VER-MED A10017 UNIVERSAL ELECTRODE**Mar 21, 1991  
104 days to decisionK905489 · Product code: **DRX** · CardiovascularSource: <https://www.510kdatabase.net/k905489/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Dec 7, 1990
Decision date	Mar 21, 1991
Days to decision	104 days
Third-party review	No

**APPLICANT**

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Company	<b>Vermont Medical, Inc.</b>
Location	Bellows Falls, VT, US
Contact	DAVID LOVELL
510(k) history	9 submissions · 9 cleared · 1978-2003

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k905489/>; 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 26, 2026