

**K030073 VERMED A10021 RESTING EKG TAB ELECTRODE**Mar 28, 2003  
79 days to decisionK030073 · Product code: **DRX** · Cardiovascular  
Source: <https://www.510kdatabase.net/k030073/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Jan 8, 2003
Decision date	Mar 28, 2003
Days to decision	79 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Vermont Medical, Inc.</b>
Location	Bellows Falls, VT, US
Contact	MARC FILLION
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/k030073/> 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