

K910139 VER-MED A10014 RESTING EKG ELECTRODE & A10021Aug 15, 1991
213 days to decisionK910139 · Product code: **DRX** · Cardiovascular
Source: <https://www.510kdatabase.net/k910139/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Jan 14, 1991
Decision date	Aug 15, 1991
Days to decision	213 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k910139/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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