

**K944655 ECG ELECTRODE**Nov 22, 1994  
62 days to decisionK944655 · Product code: **DRX** · Cardiovascular  
Source: <https://www.510kdatabase.net/k944655/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Sep 21, 1994
Decision date	Nov 22, 1994
Days to decision	62 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Conmed Andover Medical, Inc.</b>
Location	Haverhill, MA, US
Contact	BRENDA CULLINANE
510(k) history	8 submissions · 8 cleared · 1994-1995

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k944655/> 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 1, 2026