

**K863674 TRACETS 2000/ELECTRODES**Mar 11, 1987  
173 days to decisionK863674 · Product code: **DRX** · CardiovascularSource: <https://www.510kdatabase.net/k863674/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Sep 19, 1986
Decision date	Mar 11, 1987
Days to decision	173 days
Third-party review	No

**APPLICANT**

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Company	<b>Lectec Corp.</b>
Location	Mchenry, IL, US
Contact	ROLF, PH.D.
510(k) history	27 submissions · 26 cleared · 1977-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k863674/>; 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 20, 2026