

**K962122 SENTRY RADIOTRANSLUCENT ECG ELECTRODES**Jul 24, 1996  
54 days to decisionK962122 · Product code: **DRX** · CardiovascularSource: <https://www.510kdatabase.net/k962122/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	May 31, 1996
Decision date	Jul 24, 1996
Days to decision	54 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Sentry Medical Products, Inc.</b>
Location	Walker, MI, US
Contact	JAMES G STROKES
510(k) history	13 submissions · 13 cleared · 1983-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k962122/> 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