

**K954389 CONMED ECG/EKG ELECTRODES**May 16, 1996  
240 days to decisionK954389 · Product code: **DRX** · CardiovascularSource: <https://www.510kdatabase.net/k954389/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Sep 19, 1995
Decision date	May 16, 1996
Days to decision	240 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Conmedcorp</b>
Location	Dayton, OH, US
Contact	IRA D DUESLER, JR.
510(k) history	92 submissions · 92 cleared · 1981-2010

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k954389/> 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 27, 2026