

**K991105 ECG ELECTRODE (VARIOUS)**Nov 26, 1999  
239 days to decisionK991105 · Product code: **DRX** · Cardiovascular  
Source: <https://www.510kdatabase.net/k991105/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Apr 1, 1999
Decision date	Nov 26, 1999
Days to decision	239 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
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/k991105/> 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