

**K760280 ELECTRODE SERIES 1401**Aug 23, 1976  
31 days to decisionK760280 · Product code: **DRX** · CardiovascularSource: <https://www.510kdatabase.net/k760280/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Jul 23, 1976
Decision date	Aug 23, 1976
Days to decision	31 days
Third-party review	No

**APPLICANT**

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Company	<b>Andover Medical, Inc.</b>
Location	Mchenry, IL, US
Website	<a href="https://www.andovermedical.com">https://www.andovermedical.com</a>
510(k) history	21 submissions · 21 cleared · 1976-1991

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k760280/>, 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