

**K832516 MAGNATRODE CARDIAC MONITORING ELECTRODE**Dec 16, 1983  
141 days to decisionK832516 · Product code: **DRX** · CardiovascularSource: <https://www.510kdatabase.net/k832516/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Electrocardiograph (DRX)
Date received	Jul 28, 1983
Decision date	Dec 16, 1983
Days to decision	141 days
Third-party review	No

**APPLICANT**

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Company	<b>R2 Corp.</b>
Location	Mchenry, IL, US
510(k) history	9 submissions · 9 cleared · 1980-1983

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k832516/> 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