

**K901780 RTM 36-05, 60-05, RTD 36-05, 60-05 RADIO. LEADWIRE**Oct 9, 1990  
175 days to decisionK901780 · Product code: **DSA** · Cardiovascular  
Source: <https://www.510kdatabase.net/k901780/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cable, Transducer And Electrode, Patient, (including Connector) (DSA)
Date received	Apr 17, 1990
Decision date	Oct 9, 1990
Days to decision	175 days
Third-party review	No

**APPLICANT**

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Company	<b>Medtronic Vascular</b>
Location	Walker, MI, US
Contact	JANICE M PEVIDE
510(k) history	475 submissions · 453 cleared · 1977-2023

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k901780/>, 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 14, 2026