

**K791363 KWIK-AWA VEST**Aug 3, 1979  
17 days to decisionK791363 · Product code: **DRG** · CardiovascularSource: <https://www.510kdatabase.net/k791363/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Transmitters And Receivers, Physiological Signal, Radiofrequency (DRG)
Date received	Jul 17, 1979
Decision date	Aug 3, 1979
Days to decision	17 days
Third-party review	No

**APPLICANT**

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Company	<b>Medical Specialties, Inc.</b>
Location	Mchenry, IL, US
510(k) history	40 submissions · 40 cleared · 1979-1995

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k791363/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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