

**K782131 PULSETAKER**Jan 15, 1979  
19 days to decisionK782131 · Product code: **DRT** · CardiovascularSource: <https://www.510kdatabase.net/k782131/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Dec 27, 1978
Decision date	Jan 15, 1979
Days to decision	19 days
Third-party review	No

**APPLICANT**

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Company	<b>Van, Cott, Bagley, Cornwall &amp; Mccarth</b>
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1979-1979

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k782131/> 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 9, 2026