

**K802325 CRITERION II PACING SYSTEM MONITORS**Oct 23, 1980  
30 days to decisionK802325 · Product code: **DRT** · CardiovascularSource: <https://www.510kdatabase.net/k802325/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Sep 23, 1980
Decision date	Oct 23, 1980
Days to decision	30 days
Third-party review	No

**APPLICANT**

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Company	<b>Dart Medical Equipment</b>
Location	Mchenry, IL, US
510(k) history	7 submissions · 7 cleared · 1980-1982

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k802325/> 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