

**K872443 COMPANION 50 REGULATOR/FLOWMETER**Jul 21, 1987  
28 days to decisionK872443 · Product code: **DRT** · Cardiovascular  
Source: <https://www.510kdatabase.net/k872443/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Monitor, Cardiac (incl. Cardiotachometer & Rate Alarm) (DRT)
Date received	Jun 23, 1987
Decision date	Jul 21, 1987
Days to decision	28 days
Third-party review	No

**APPLICANT**

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Company	<b>Puritan Bennett Corp.</b>
Location	Mchenry, IL, US
Contact	HARTLEY C ERICSON
510(k) history	110 submissions · 101 cleared · 1976-2007

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k872443/>; 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 25, 2026