

**K040922 VENTURE WIRE CONTROL CATHETER**Aug 18, 2004  
132 days to decisionK040922 · Product code: **DQY** · CardiovascularSource: <https://www.510kdatabase.net/k040922/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Apr 8, 2004
Decision date	Aug 18, 2004
Days to decision	132 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Velocimed, Inc.</b>
Location	Mineapolis, MN, US
Contact	Sew-Wah Tay
510(k) history	5 submissions · 5 cleared · 2004-2006

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k040922/> 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 June 29, 2026