

**K132652 VIA MICROCATHETER, VIA PLUS MICROCATHETER**Dec 20, 2013  
116 days to decisionK132652 · Product code: **DQY** · Cardiovascular  
Source: <https://www.510kdatabase.net/k132652/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Aug 26, 2013
Decision date	Dec 20, 2013
Days to decision	116 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Sequent Medical, Inc.</b>
Location	Aliso Viejo, CA, US
Contact	MELANIE PARRAVI
510(k) history	4 submissions · 4 cleared · 2013-2017

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k132652/> 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 1, 2026