

**K091624 STENT GRAFT BALLOON CATHETER**Oct 15, 2009  
134 days to decisionK091624 · Product code: **DQY** · CardiovascularSource: <https://www.510kdatabase.net/k091624/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Jun 3, 2009
Decision date	Oct 15, 2009
Days to decision	134 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Via Biomedical, Inc.</b>
Location	Maple Grove, MN, US
Contact	FERNANDO DI CAPRIO
510(k) history	1 submissions · 1 cleared · 2009-2009

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k091624/> 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 22, 2026