

**K133445 ANGIOVAC CANNULA**Mar 5, 2014  
113 days to decisionK133445 · Product code: **DWF** · Cardiovascular  
Source: <https://www.510kdatabase.net/k133445/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Cannula And Tubing, Vascular, Cardiopulmonary Bypass (DWF)
Date received	Nov 12, 2013
Decision date	Mar 5, 2014
Days to decision	113 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Vortex Medical, Inc.</b>
Location	Norwell, MA, US
Contact	Lorraine M Hanley
510(k) history	3 submissions · 3 cleared · 2009-2014

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k133445/> 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 7, 2026