

**K142593 AngioVac Cannula**Nov 12, 2014  
58 days to decisionK142593 · Product code: **DWF** · CardiovascularSource: <https://www.510kdatabase.net/k142593/>**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	Sep 15, 2014
Decision date	Nov 12, 2014
Days to decision	58 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>AngioDynamics, Inc.</b>
Location	Glens Falls, NY, US
Contact	Wanda Carpinella
Website	<a href="http://www.angiodynamics.com/">http://www.angiodynamics.com/</a>
510(k) history	87 submissions · 82 cleared · 1995-2025

AngioDynamics, Inc. is a global leader in vascular and oncology medical technologies, with a manufacturing facility in Glens Falls, US. The company develops advanced devices addressing blood flow restoration, cancer therapies, vascular access, and varicose vein treatment. AngioDynamics has received FDA 510(k) clearances from total submissions since its first clearance in 1995. The company specializes in cardiovascular devices, with recent cleared products including mechanical aspiration systems, infusion systems, and angiographic catheters. The latest FDA 510(k) clearance...

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k142593/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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