

**K130896 VASCULAR PROBE, VASCULAR PROBE ES**Apr 24, 2013  
23 days to decisionK130896 · Product code: **DWP** · CardiovascularSource: <https://www.510kdatabase.net/k130896/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dilator, Vessel, Surgical (DWP)
Date received	Apr 1, 2013
Decision date	Apr 24, 2013
Days to decision	23 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Synovis Life Technologies, Inc.</b>
Location	St. Paul, MN, US
Contact	STEPHANI K AYALA
Website	<a href="http://www.synovislife.com/">http://www.synovislife.com/</a>
510(k) history	9 submissions · 9 cleared · 2013-2023

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k130896/> 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 May 25, 2026