

K003719 VASCULAR ARCHITECTS PERISCOPE DEVICEFeb 28, 2001
86 days to decisionK003719 · Product code: **DWX** · CardiovascularSource: <https://www.510kdatabase.net/k003719/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stripper, Artery, Intraluminal (DWX)
Date received	Dec 4, 2000
Decision date	Feb 28, 2001
Days to decision	86 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Vascular Architects, Inc.
Location	San Jose, CA, US
Contact	JEAN M CAILLOUETTE
510(k) history	5 submissions · 5 cleared · 2001-2003

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k003719/> Data sourced from FDA 510(k) public records (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. - Generated June 30, 2026