FDA 510(k) Premarket Notification

K003719 VASCULAR ARCHITECTS PERISCOPE DEVICE

CLEARED

Feb 28, 2001
86 days to decision

K003719 · Product code: **DWX** · Cardiovascular
Source: 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