FDA 510(k) Premarket Notification

K130896 VASCULAR PROBE, VASCULAR PROBE ES

CLEARED

Apr 24, 2013
23 days to decision

K130896 · Product code: **DWP** · Cardiovascular
Source: https://www.510kdatabase.net/k130896/

SUBMISSION DETAILS

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

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