FDA 510(k) Premarket Notification

K222679 Vericor Support Catheter

CLEARED

Nov 7, 2022
62 days to decision

K222679 · Product code: **KRA** · Cardiovascular
Source: https://www.510kdatabase.net/k222679/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Continuous Flush (KRA)
Date received	Sep 6, 2022
Decision date	Nov 7, 2022
Days to decision	62 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Vascupatent Medical (Shenzhen) Co., Ltd.**
Location	Shenzhen, CN
Contact	Heather Li
510(k) history	1 submissions · 1 cleared · 2022-2022

REGULATORY CONSULTANT

Consulting firm	**Mid-Link Consulting Co, Ltd.**
Contact	Diana Hong

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov