

K222679 Vericor Support CatheterNov 7, 2022
62 days to decisionK222679 · 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](https://accessdata.fda.gov)

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