

K241851 Versus™ Catheter (VS110-8B)Nov 25, 2024
151 days to decisionK241851 · Product code: **QEY** · Cardiovascular
Source: <https://www.510kdatabase.net/k241851/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Mechanical Thrombolysis Catheter (QEY)
Date received	Jun 27, 2024
Decision date	Nov 25, 2024
Days to decision	151 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Liquet Medical, Inc.
Location	Glen Allen, VA, US
Contact	John Schindler
Website	https://liquetmedical.com
510(k) history	2 submissions · 2 cleared · 2024-2026

REGULATORY CONSULTANT

Consulting firm	Evergreen Strategic Consulting
Contact	Carrie Kuehn

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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