FDA 510(k) Premarket Notification

K241851 Versus™ Catheter (VS110-8B)

CLEARED

Nov 25, 2024
151 days to decision

K241851 · 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