

**K260149 Versus™ Catheter (VS110-9NB)**Feb 17, 2026  
28 days to decisionK260149 · Product code: **QEY** · CardiovascularSource: <https://www.510kdatabase.net/k260149/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Mechanical Thrombolysis Catheter (QEY)
Date received	Jan 20, 2026
Decision date	Feb 17, 2026
Days to decision	28 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Liquet Medical, Inc.</b>
Location	Glen Allen, VA, US
Contact	John Schnidler
Website	<a href="https://liquetmedical.com">https://liquetmedical.com</a>
510(k) history	2 submissions · 2 cleared · 2024-2026

**REGULATORY CONSULTANT**

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Consulting firm	<b>Evergreen Strategic Consulting</b>
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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