

**K253589 Liberant™ RX Aspiration Catheter**Apr 23, 2026  
157 days to decisionK253589 · Product code: **QEZ** · Cardiovascular  
Source: <https://www.510kdatabase.net/k253589/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Aspiration Thrombectomy Catheter (QEZ)
Date received	Nov 17, 2025
Decision date	Apr 23, 2026
Days to decision	157 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Medtronic Interventional Vascular, Inc.</b>
Location	Danvers, MA, US
Contact	Nikita Ciandra Vaz
510(k) history	1 submissions · 1 cleared · 2026-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k253589/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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