

**K250787 Liberant Thrombectomy System**Jun 11, 2025  
89 days to decisionK250787 · Product code: **QEW** · CardiovascularSource: <https://www.510kdatabase.net/k250787/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Peripheral Mechanical Thrombectomy With Aspiration (QEW)
Date received	Mar 14, 2025
Decision date	Jun 11, 2025
Days to decision	89 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Medtronic, Ireland</b>
Location	Shoreview, MN, US
Contact	Barbara Zampedri
510(k) history	7 submissions · 7 cleared · 2006-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k250787/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 13, 2026