FDA 510(k) Premarket Notification

K253589 Liberant™ RX Aspiration Catheter

CLEARED

Apr 23, 2026
157 days to decision

K253589 · Product code: **QEZ** · Cardiovascular
Source: https://www.510kdatabase.net/k253589/

SUBMISSION DETAILS

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

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