FDA 510(k) Premarket Notification

K250787 Liberant Thrombectomy System

CLEARED

Jun 11, 2025
89 days to decision

K250787 · Product code: **QEW** · Cardiovascular
Source: https://www.510kdatabase.net/k250787/

SUBMISSION DETAILS

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

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