

K243756 0.014" Willow GuidewireJul 17, 2025
223 days to decisionK243756 · Product code: **MOF** · Neurology
Source: <https://www.510kdatabase.net/k243756/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Guide, Wire, Catheter, Neurovasculature (MOF)
Date received	Dec 6, 2024
Decision date	Jul 17, 2025
Days to decision	223 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Arbor Endovascular, LLC
Location	San Jose, CA, US
Contact	Kim Otto
Website	https://arborvascular.com
510(k) history	4 submissions · 4 cleared · 2025-2026

REGULATORY CONSULTANT

Consulting firm	Fabrica Consulting, LLC
Contact	Sara Toyloy

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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