

K260130 Willow 18 GuidewireFeb 13, 2026
28 days to decisionK260130 · Product code: **MOF** · Neurology
Source: <https://www.510kdatabase.net/k260130/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Guide, Wire, Catheter, Neurovasculature (MOF)
Date received	Jan 16, 2026
Decision date	Feb 13, 2026
Days to decision	28 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	Kathy Tansey
Website	https://arborvascular.com
510(k) history	4 submissions · 4 cleared · 2025-2026

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