FDA 510(k) Premarket Notification

K260130 Willow 18 Guidewire

CLEARED

Feb 13, 2026
28 days to decision

K260130 · 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