

K202522 Synchro SELECT GuidewireSep 25, 2020
24 days to decisionK202522 · Product code: **MOF** · Neurology
Source: <https://www.510kdatabase.net/k202522/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Guide, Wire, Catheter, Neurovasculature (MOF)
Date received	Sep 1, 2020
Decision date	Sep 25, 2020
Days to decision	24 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Stryker Neurovascular
Location	Freemont, CA, US
Contact	Hailey Hinkle
Website	https://www.stryker.com
510(k) history	32 submissions · 32 cleared · 2011-2026

Stryker Neurovascular is a medical device manufacturer based in Fremont, US. The company specializes in innovative interventional neurology solutions. Stryker Neurovascular has received FDA 510(k) clearances from total submissions since 2011. The company's portfolio is dominated by Neurology devices, representing 84% of regulatory submissions. The latest clearance was granted in 2026, demonstrating continued active development and market engagement. Recent cleared devices include intracranial base catheters, detachable coils, microcatheters, and thrombectomy retrievers. T...

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Device record: <https://www.510kdatabase.net/k202522/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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