

K243593 AXS Lift Intracranial Base CatheterFeb 18, 2025
90 days to decisionK243593 · Product code: **QJP** · Neurology
Source: <https://www.510kdatabase.net/k243593/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous, Neurovasculature (QJP)
Date received	Nov 20, 2024
Decision date	Feb 18, 2025
Days to decision	90 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Stryker Neurovascular
Location	Freemont, CA, US
Contact	Natalie Allen
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/k243593/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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