

**K203219 Trevo XP ProVue Retriever and Trevo NXT™ ProVue Retriever**Dec 29, 2020  
57 days to decisionK203219 · Product code: **POL** · Neurology  
Source: <https://www.510kdatabase.net/k203219/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Neurovascular Mechanical Thrombectomy Device For Acute Ischemic Stroke Treatment (POL)
Date received	Nov 2, 2020
Decision date	Dec 29, 2020
Days to decision	57 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Stryker Neurovascular</b>
Location	Freemont, CA, US
Contact	Kathy Nguyen
Website	<a href="https://www.stryker.com">https://www.stryker.com</a>
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...