

K132542 TRANSGATE GUIDE CATHETERNov 15, 2013
94 days to decisionK132542 · Product code: **DQY** · CardiovascularSource: <https://www.510kdatabase.net/k132542/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous (DQY)
Date received	Aug 13, 2013
Decision date	Nov 15, 2013
Days to decision	94 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Stryker Neurovascular
Location	Freemont, CA, US
Contact	ANJALI ATAL-GUPTA
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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