

K242392 PATH BGCMay 9, 2025
270 days to decisionK242392 · Product code: **QJP** · Neurology
Source: <https://www.510kdatabase.net/k242392/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous, Neurovasculature (QJP)
Date received	Aug 12, 2024
Decision date	May 9, 2025
Days to decision	270 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Crossroads Neurovascular, Inc.
Location	Lake Forest, CA, US
Contact	Ryan Breckenridge
510(k) history	2 submissions · 2 cleared · 2025-2026

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k242392/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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