FDA 510(k) Premarket Notification

# K242392 PATH BGC

**CLEARED**

May 9, 2025
270 days to decision

K242392 · 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 |