

**K242376 Next Generation Access Platform**Dec 30, 2024  
143 days to decisionK242376 · Product code: **QJP** · Neurology  
Source: <https://www.510kdatabase.net/k242376/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous, Neurovasculature (QJP)
Date received	Aug 9, 2024
Decision date	Dec 30, 2024
Days to decision	143 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Balt USA, LLC</b>
Location	Irvine, CA, US
Contact	Alicia Smith
510(k) history	18 submissions · 18 cleared · 2018-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k242376/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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