

K231393 Radical the Dude 7F Guide CatheterNov 30, 2023
202 days to decisionK231393 · Product code: **QJP** · Neurology
Source: <https://www.510kdatabase.net/k231393/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Percutaneous, Neurovasculature (QJP)
Date received	May 12, 2023
Decision date	Nov 30, 2023
Days to decision	202 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Maduro Medical, Inc.
Location	Campbell, CA, US
Contact	Janice Kemp
510(k) history	3 submissions · 3 cleared · 2023-2026

REGULATORY CONSULTANT

Consulting firm	Mcra, LLC
Contact	Robert Herrmann, Ph.D.

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)

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