

K223436 Artix AXMar 2, 2023
108 days to decisionK223436 · Product code: **QEW** · CardiovascularSource: <https://www.510kdatabase.net/k223436/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Peripheral Mechanical Thrombectomy With Aspiration (QEW)
Date received	Nov 14, 2022
Decision date	Mar 2, 2023
Days to decision	108 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Inari Medical
Location	Irvine, CA, US
Contact	Ellen Nguyen
510(k) history	26 submissions · 26 cleared · 2015-2025

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k223436/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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