

**K223000 Artix BG**Jan 11, 2023  
105 days to decisionK223000 · Product code: **QEW** · CardiovascularSource: <https://www.510kdatabase.net/k223000/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Peripheral Mechanical Thrombectomy With Aspiration (QEW)
Date received	Sep 28, 2022
Decision date	Jan 11, 2023
Days to decision	105 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k223000/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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