

**K240384 SENDERO Microcatheter**Mar 8, 2024  
29 days to decisionK240384 · Product code: **DQY** · Cardiovascular  
Source: <https://www.510kdatabase.net/k240384/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Percutaneous (DQY)
Date received	Feb 8, 2024
Decision date	Mar 8, 2024
Days to decision	29 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Okami Medical, Inc.</b>
Location	Aliso Viejo, CA, US
Contact	Jill Delsman
510(k) history	4 submissions · 4 cleared · 2023-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k240384/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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