

**K232647 Microcatheter**Sep 29, 2023  
29 days to decisionK232647 · Product code: **DQY** · CardiovascularSource: <https://www.510kdatabase.net/k232647/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Percutaneous (DQY)
Date received	Aug 31, 2023
Decision date	Sep 29, 2023
Days to decision	29 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Surmodics, Inc.</b>
Location	Eden Prairie, MN, US
Contact	Jodie Christe
510(k) history	11 submissions · 11 cleared · 2018-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k232647/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 26, 2026