

**K173200 SOFIA Plus Aspiration Catheter**Jun 11, 2018  
252 days to decisionK173200 · Product code: **NRY** · Neurology  
Source: <https://www.510kdatabase.net/k173200/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Thrombus Retriever (NRY)
Date received	Oct 2, 2017
Decision date	Jun 11, 2018
Days to decision	252 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>MicroVention, Inc.</b>
Location	Aliso Viejo, CA, US
Contact	Naomi Gong
510(k) history	85 submissions · 85 cleared · 2001-2024

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k173200/> 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 30, 2026