

**K062046 MODIFIED MERCI RETRIEVER, MODEL 90050**Jan 19, 2007  
184 days to decisionK062046 · Product code: **NRV** · Neurology  
Source: <https://www.510kdatabase.net/k062046/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Thrombus Retriever (NRV)
Date received	Jul 19, 2006
Decision date	Jan 19, 2007
Days to decision	184 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Concentric Medical, Inc.</b>
Location	Moutian View, CA, US
Contact	KIRSTEN VALLEY
510(k) history	45 submissions · 44 cleared · 2001-2018

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k062046/> 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 14, 2026