

**K093610 MODIFIED RETRIEVER (MODIFIED, LINE EXTENSION),
MODELS 90152, 90153**Feb 17, 2011
454 days to decisionK093610 · Product code: **NRY** · Neurology
Source: <https://www.510kdatabase.net/k093610/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Thrombus Retriever (NRY)
Date received	Nov 20, 2009
Decision date	Feb 17, 2011
Days to decision	454 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k093610/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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