

**K172167 Zenith Flex System**Apr 19, 2018  
275 days to decisionK172167 · Product code: **NRY** · Neurology  
Source: <https://www.510kdatabase.net/k172167/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Thrombus Retriever (NRY)
Date received	Jul 18, 2017
Decision date	Apr 19, 2018
Days to decision	275 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Inneuroco, Inc.</b>
Location	Sunrise, FL, US
Contact	Marc Litzenberg
510(k) history	9 submissions · 9 cleared · 2016-2023

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k172167/> 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