

K190338 046 Zenith FlexAug 2, 2019
169 days to decisionK190338 · Product code: **NRY** · Neurology
Source: <https://www.510kdatabase.net/k190338/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Thrombus Retriever (NRY)
Date received	Feb 14, 2019
Decision date	Aug 2, 2019
Days to decision	169 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k190338/> Data sourced from FDA 510(k) public records (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. - Generated May 26, 2026