FDA 510(k) Premarket Notification

K173200 SOFIA Plus Aspiration Catheter

CLEARED

Jun 11, 2018
252 days to decision

K173200 · Product code: **NRY** · Neurology
Source: https://www.510kdatabase.net/k173200/

SUBMISSION DETAILS

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

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