FDA 510(k) Premarket Notification

K062046 MODIFIED MERCI RETRIEVER, MODEL 90050

CLEARED

Jan 19, 2007
184 days to decision

K062046 · Product code: **NRY** · Neurology
Source: https://www.510kdatabase.net/k062046/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Thrombus Retriever (NRY)
Date received	Jul 19, 2006
Decision date	Jan 19, 2007
Days to decision	184 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