FDA 510(k) Premarket Notification

K133530 NUVASIVE EMG ENDOTRACHEAL TUBE

CLEARED

May 2, 2014
168 days to decision

K133530 · Product code: **PDQ** · Neurology
Source: https://www.510kdatabase.net/k133530/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Neurosurgical Nerve Locator (PDQ)
Date received	Nov 15, 2013
Decision date	May 2, 2014
Days to decision	168 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Nu Vasive, Incorporated**
Location	San Diego, CA, US
Contact	JEREMY MARKOVICH
510(k) history	112 submissions · 112 cleared · 2012-2023