FDA 510(k) Premarket Notification

K110712 DRYTOUCH SUCTION STIMULATOR PROBE

CLEARED

Jun 28, 2011
106 days to decision

K110712 · Product code: **ETN** · Ear, Nose, Throat
Source: https://www.510kdatabase.net/k110712/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve (ETN)
Date received	Mar 14, 2011
Decision date	Jun 28, 2011
Days to decision	106 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Neurovision Medical Products, Inc.**
Location	Ventura, CA, US
Contact	CHRISTINE VERGELY
510(k) history	9 submissions · 9 cleared · 2010-2013