

**K110712 DRYTOUCH SUCTION STIMULATOR PROBE**Jun 28, 2011  
106 days to decisionK110712 · 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	<b>Neurovision Medical Products, Inc.</b>
Location	Ventura, CA, US
Contact	CHRISTINE VERGELY
510(k) history	9 submissions · 9 cleared · 2010-2013

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k110712/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 24, 2026