

**K133530 NUVASIVE EMG ENDOTRACHEAL TUBE**May 2, 2014  
168 days to decisionK133530 · Product code: **PDQ** · Neurology  
Source: <https://www.510kdatabase.net/k133530/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k133530/> 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 June 7, 2026