

**K171599 WiTouch Pro, WiTouch, Neubac**Sep 22, 2017  
113 days to decisionK171599 · Product code: **NUH** · Neurology  
Source: <https://www.510kdatabase.net/k171599/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, Over-the-counter (NUH)
Date received	Jun 1, 2017
Decision date	Sep 22, 2017
Days to decision	113 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Hollywog, LLC</b>
Location	Chattanooga, TN, US
Contact	Michael Treas
510(k) history	4 submissions · 4 cleared · 2012-2017

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k171599/> 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 31, 2026