

**K170659 IQ Technologies**May 26, 2017  
84 days to decisionK170659 · Product code: **NUH** · Neurology  
Source: <https://www.510kdatabase.net/k170659/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Stimulator, Nerve, Transcutaneous, Over-the-counter (NUH)
Date received	Mar 3, 2017
Decision date	May 26, 2017
Days to decision	84 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Iq Technologies, Inc.</b>
Location	Las Vegas, NV, US
Contact	Elli Josef
510(k) history	2 submissions · 2 cleared · 2014-2017

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k170659/> 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 1, 2026