

**K202940 First Relief v1**Dec 29, 2020  
90 days to decisionK202940 · Product code: **QHH** · Neurology  
Source: <https://www.510kdatabase.net/k202940/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Non-implanted Nerve Stimulator For Pain Associated With Irritable Bowel Syndrome (ibs) (QHH)
Date received	Sep 30, 2020
Decision date	Dec 29, 2020
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Dyansys, Inc.</b>
Location	Orinda, CA, US
Contact	Srini Nageshwar
510(k) history	14 submissions · 14 cleared · 2007-2022

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k202940/> 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 26, 2026