

**K103370 RT200**Apr 5, 2011  
139 days to decisionK103370 · Product code: **GZI** · Neurology  
Source: <https://www.510kdatabase.net/k103370/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Neuromuscular, External Functional (GZI)
Date received	Nov 17, 2010
Decision date	Apr 5, 2011
Days to decision	139 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Restorative Therapies, Inc.</b>
Location	Baltimore, MD, US
Contact	ANDREW BARRISKILL
510(k) history	10 submissions · 10 cleared · 2005-2017

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k103370/> 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