

**K052374 RAPID PROGEAMMER 3.0**Nov 22, 2005  
84 days to decisionK052374 · Product code: **GZB** · Neurology  
Source: <https://www.510kdatabase.net/k052374/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Spinal-cord, Implanted (pain Relief) (GZB)
Date received	Aug 30, 2005
Decision date	Nov 22, 2005
Days to decision	84 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Advanced Neuromodulation Systems</b>
Location	Plano, TX, US
Contact	KATRYNA WARREN
510(k) history	25 submissions · 25 cleared · 1999-2009

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k052374/> 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 21, 2026