

**K072462 WIDE SPACED QUATTRODE LEADS**Oct 11, 2007  
41 days to decisionK072462 · Product code: **GZB** · Neurology  
Source: <https://www.510kdatabase.net/k072462/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Stimulator, Spinal-cord, Implanted (pain Relief) (GZB)
Date received	Aug 31, 2007
Decision date	Oct 11, 2007
Days to decision	41 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Advanced Neuromodulation Systems</b>
Location	Plano, TX, US
Contact	MIA M WARE
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/k072462/> 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