

**K022222 LAMITRODE S-SERIES (S4 AND S8) LEADS**Aug 8, 2002  
30 days to decisionK022222 · Product code: **GZB** · Neurology  
Source: <https://www.510kdatabase.net/k022222/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Stimulator, Spinal-cord, Implanted (pain Relief) (GZB)
Date received	Jul 9, 2002
Decision date	Aug 8, 2002
Days to decision	30 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Advanced Neuromodulation System,Inc</b>
Location	Plano, TX, US
Contact	KATRYNA WARREN
510(k) history	1 submissions · 1 cleared · 2002-2002

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k022222/> 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 30, 2026