

**K905255 NEUROMED MODEL 1992 LS LAMITRODE LEAD**Jan 25, 1991  
65 days to decisionK905255 · Product code: **GZB** · Neurology  
Source: <https://www.510kdatabase.net/k905255/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Spinal-cord, Implanted (pain Relief) (GZB)
Date received	Nov 21, 1990
Decision date	Jan 25, 1991
Days to decision	65 days
Third-party review	No

**APPLICANT**

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Company	<b>Neuromed, Inc.</b>
Location	Ft. Lauderdale, FL, US
Contact	DEAN. E CIPORKIN
510(k) history	15 submissions · 15 cleared · 1985-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k905255/>; 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