

**K896450 NEURO N 50 LESION GENERATOR**Mar 19, 1991  
495 days to decisionK896450 · Product code: **GXD** · Neurology  
Source: <https://www.510kdatabase.net/k896450/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Generator, Lesion, Radiofrequency (GXD)
Date received	Nov 9, 1989
Decision date	Mar 19, 1991
Days to decision	495 days
Third-party review	No

**APPLICANT**

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Company	<b>Leibinger &amp; Fischer , Ltd.</b>
Location	Dallas, TX, US
Contact	PAUL STEWART
510(k) history	2 submissions · 2 cleared · 1990-1991

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k896450/>, 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 May 21, 2026