

**K896620 NEUROTENS**Apr 26, 1990  
156 days to decisionK896620 · Product code: **GZJ** · Neurology  
Source: <https://www.510kdatabase.net/k896620/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Nov 21, 1989
Decision date	Apr 26, 1990
Days to decision	156 days
Third-party review	No

**APPLICANT**

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Company	<b>Henley Intl.</b>
Location	Houston, TX, US
Contact	ERNEST J HENLEY,PHD
510(k) history	44 submissions · 41 cleared · 1986-1995

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k896620/>, 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 29, 2026