

**K896787 TRANSCUTANEOUS ELECTRICAL NERVE  
STIMULATION**Jul 30, 1990  
238 days to decisionK896787 · Product code: **GZJ** · Neurology  
Source: <https://www.510kdatabase.net/k896787/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Dec 4, 1989
Decision date	Jul 30, 1990
Days to decision	238 days
Third-party review	No

**APPLICANT**

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Company	<b>Saratoga Intl., Inc.</b>
Location	San Jose, CA, US
Contact	MIKE CHUANG
510(k) history	2 submissions · 2 cleared · 1990-1994

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k896787/>; 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