

**K962332 TENS/FES/NMES ELECTRODES**Jul 31, 1996  
44 days to decisionK962332 · Product code: **GZJ** · Neurology  
Source: <https://www.510kdatabase.net/k962332/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Jun 17, 1996
Decision date	Jul 31, 1996
Days to decision	44 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Uni-Patch, Inc.</b>
Location	Wabasha, MN, US
Contact	THOMAS MOORE
510(k) history	6 submissions · 6 cleared · 1987-1999

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k962332/> 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