

**K881832 DS-200 AND DS-400 T.E.N.S.**Jul 1, 1988  
63 days to decisionK881832 · Product code: **GZJ** · Neurology  
Source: <https://www.510kdatabase.net/k881832/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Apr 29, 1988
Decision date	Jul 1, 1988
Days to decision	63 days
Third-party review	No

**APPLICANT**

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Company	<b>Delta Marketing, Inc.</b>
Location	Muskogee, OK, US
Contact	DAVID R WATERS
510(k) history	1 submissions · 1 cleared · 1988-1988

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k881832/>; 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 July 1, 2026