

**K880492 INTERCEPT T.E.N.S. DEVICE**Apr 18, 1988  
74 days to decisionK880492 · Product code: **GZJ** · Neurology  
Source: <https://www.510kdatabase.net/k880492/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Feb 4, 1988
Decision date	Apr 18, 1988
Days to decision	74 days
Third-party review	No

**APPLICANT**

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Company	<b>Medical Design &amp; Manufacturing Corp.</b>
Location	Walker, MI, US
Contact	MICHAEL SOUTHWORTH
510(k) history	10 submissions · 10 cleared · 1981-1992

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k880492/>; 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