

**K880452 LUMISCOPE TENS MODEL 2000T OR APCC 3001 OR PAIN RE**Mar 24, 1988  
51 days to decisionK880452 · Product code: **GZJ** · Neurology  
Source: <https://www.510kdatabase.net/k880452/>**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 2, 1988
Decision date	Mar 24, 1988
Days to decision	51 days
Third-party review	No

**APPLICANT**

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Company	<b>Lumiscop Co., Inc.</b>
Location	Edison, NJ, US
Contact	LOUIS C MORGENIER
510(k) history	13 submissions · 13 cleared · 1988-2003

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k880452/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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