

**K802678 TRANSCUTANEOUS ELECTRIC NERVE STIMULATOR**Dec 10, 1980  
42 days to decisionK802678 · Product code: **GZJ** · Neurology  
Source: <https://www.510kdatabase.net/k802678/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Oct 29, 1980
Decision date	Dec 10, 1980
Days to decision	42 days
Third-party review	No

**APPLICANT**

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Company	<b>General Medical Industries, Inc.</b>
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1980-1980

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Device record: <https://www.510kdatabase.net/k802678/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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