

**K831413 SPEC IISX**Jun 8, 1983  
37 days to decisionK831413 · Product code: **GZJ** · Neurology  
Source: <https://www.510kdatabase.net/k831413/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	May 2, 1983
Decision date	Jun 8, 1983
Days to decision	37 days
Third-party review	No

**APPLICANT**

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Company	<b>Medical Devices, Inc.</b>
Location	Mchenry, IL, US
510(k) history	49 submissions · 47 cleared · 1977-2001

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Device record: <https://www.510kdatabase.net/k831413/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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