

**K811939 VITATRON 88**Jul 23, 1981  
17 days to decisionK811939 · Product code: **GZJ** · Neurology  
Source: <https://www.510kdatabase.net/k811939/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Jul 6, 1981
Decision date	Jul 23, 1981
Days to decision	17 days
Third-party review	No

**APPLICANT**

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Company	<b>Vitatronics</b>
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1981-1981

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k811939/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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