

**K820678 SYSTEM 10**Mar 23, 1982  
12 days to decisionK820678 · Product code: **GZJ** · Neurology  
Source: <https://www.510kdatabase.net/k820678/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Mar 11, 1982
Decision date	Mar 23, 1982
Days to decision	12 days
Third-party review	No

**APPLICANT**

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Company	<b>Biostim, Inc.</b>
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1980-1985

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Device record: <https://www.510kdatabase.net/k820678/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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