

K812432 HEALTH-UPNov 6, 1981
73 days to decisionK812432 · Product code: **GZJ** · Neurology
Source: <https://www.510kdatabase.net/k812432/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Aug 25, 1981
Decision date	Nov 6, 1981
Days to decision	73 days
Third-party review	No

APPLICANT

Company	Wfx Industries, Inc.
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1981-1981

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Device record: <https://www.510kdatabase.net/k812432/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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