

K813515 VERI/DFS TM #817Jan 19, 1982
35 days to decisionK813515 · Product code: **GZI** · Neurology
Source: <https://www.510kdatabase.net/k813515/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Neuromuscular, External Functional (GZI)
Date received	Dec 15, 1981
Decision date	Jan 19, 1982
Days to decision	35 days
Third-party review	No

APPLICANT

Company	Verite
Location	Mchenry, IL, US
510(k) history	10 submissions · 9 cleared · 1979-1985

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Device record: <https://www.510kdatabase.net/k813515/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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