FDA 510(k) Premarket Notification

K813515 VERI/DFS TM #817

CLEARED

Jan 19, 1982
35 days to decision

K813515 · 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