FDA 510(k) Premarket Notification

K811351 VERI-FIER

CLEARED

Jul 30, 1981
78 days to decision

K811351 · Product code: **GZJ** · Neurology
Source: https://www.510kdatabase.net/k811351/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	May 13, 1981
Decision date	Jul 30, 1981
Days to decision	78 days
Third-party review	No

APPLICANT

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