FDA 510(k) Premarket Notification

K820651 MODEL 860-1 VERI/PROBE

CLEARED

Apr 29, 1982
51 days to decision

K820651 · Product code: **GZJ** · Neurology
Source: https://www.510kdatabase.net/k820651/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Mar 9, 1982
Decision date	Apr 29, 1982
Days to decision	51 days
Third-party review	No

APPLICANT

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