FDA 510(k) Premarket Notification

K821363 TRANS-Q

CLEARED

May 28, 1982
21 days to decision

K821363 · Product code: **GZJ** · Neurology
Source: https://www.510kdatabase.net/k821363/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	May 7, 1982
Decision date	May 28, 1982
Days to decision	21 days
Third-party review	No

APPLICANT

Company	**Chesapeake Medical Assoc.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1982-1982