FDA 510(k) Premarket Notification

K802678 TRANSCUTANEOUS ELECTRIC NERVE STIMULATOR

CLEARED

Dec 10, 1980
42 days to decision

K802678 · Product code: **GZJ** · Neurology
Source: https://www.510kdatabase.net/k802678/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Oct 29, 1980
Decision date	Dec 10, 1980
Days to decision	42 days
Third-party review	No

APPLICANT

Company	**General Medical Industries, Inc.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1980-1980