FDA 510(k) Premarket Notification

K803121 MENTOR 150 TRANSCUTANEOUS NERVE STIMULA

CLEARED

Dec 30, 1980
19 days to decision

K803121 · Product code: **GZJ** · Neurology
Source: https://www.510kdatabase.net/k803121/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Dec 11, 1980
Decision date	Dec 30, 1980
Days to decision	19 days
Third-party review	No

APPLICANT

Company	**George H. Frisch**
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1980-1980