FDA 510(k) Premarket Notification

K810593 NEUROPAC ULTRA I

CLEARED

Mar 13, 1981
8 days to decision

K810593 · Product code: **GZJ** · Neurology
Source: https://www.510kdatabase.net/k810593/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	Mar 5, 1981
Decision date	Mar 13, 1981
Days to decision	8 days
Third-party review	No

APPLICANT

Company	**Medical Devices, Inc.**
Location	Mchenry, IL, US
510(k) history	49 submissions · 47 cleared · 1977-2001