FDA 510(k) Premarket Notification

K811499 TENSMAX

CLEARED

Aug 20, 1981
85 days to decision

K811499 · Product code: **GZJ** · Neurology
Source: https://www.510kdatabase.net/k811499/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Nerve, Transcutaneous, For Pain Relief (GZJ)
Date received	May 27, 1981
Decision date	Aug 20, 1981
Days to decision	85 days
Third-party review	No

APPLICANT

Company	**Anritsu America Co.**
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1979-1981