

K811499 TENSMAXAug 20, 1981
85 days to decisionK811499 · 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 |

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k811499/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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