

K810593 NEUROPAC ULTRA IMar 13, 1981
8 days to decisionK810593 · 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

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Device record: <https://www.510kdatabase.net/k810593/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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