FDA 510(k) Premarket Notification

K810482 SOMATOSENSORY STIMULATOR

CLEARED

Apr 29, 1981
65 days to decision

K810482 · Product code: **GWF** · Neurology
Source: https://www.510kdatabase.net/k810482/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Electrical, Evoked Response (GWF)
Date received	Feb 23, 1981
Decision date	Apr 29, 1981
Days to decision	65 days
Third-party review	No

APPLICANT

Company	**Amplaid USA, Inc.**
Location	Mchenry, IL, US
510(k) history	8 submissions · 8 cleared · 1978-1986

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k810482/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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