

K810481 VISUAL PATTERN REVERSAL GEN. & PHOTIC S.Mar 11, 1981
16 days to decisionK810481 · Product code: **GWE** · Neurology
Source: <https://www.510kdatabase.net/k810481/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Photic, Evoked Response (GWE)
Date received	Feb 23, 1981
Decision date	Mar 11, 1981
Days to decision	16 days
Third-party review	No

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k810481/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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