

**K781667 AMPLAID MARK 4 AND MARK 5**Nov 3, 1978  
39 days to decisionK781667 · Product code: **GWJ** · Neurology  
Source: <https://www.510kdatabase.net/k781667/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Auditory, Evoked Response (GWJ)
Date received	Sep 25, 1978
Decision date	Nov 3, 1978
Days to decision	39 days
Third-party review	No

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k781667/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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