

**K801671 ESG-1 (EVOKED RESPONSE STIMULUS GENER)**Aug 4, 1980  
13 days to decisionK801671 · Product code: **GWE** · Neurology  
Source: <https://www.510kdatabase.net/k801671/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Photic, Evoked Response (GWE)
Date received	Jul 22, 1980
Decision date	Aug 4, 1980
Days to decision	13 days
Third-party review	No

**APPLICANT**

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Company	<b>Neurodiagnostics, Inc.</b>
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1980-1984

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

Device record: <https://www.510kdatabase.net/k801671/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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