

**K830877 AP-CHECK**Jun 8, 1983  
82 days to decisionK830877 · Product code: **GYA** · Neurology  
Source: <https://www.510kdatabase.net/k830877/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tester, Electrode/lead, Electroencephalograph (GYA)
Date received	Mar 18, 1983
Decision date	Jun 8, 1983
Days to decision	82 days
Third-party review	No

**APPLICANT**

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Company	<b>General Devices</b>
Location	Bergenfield, NJ, US
510(k) history	13 submissions · 13 cleared · 1983-2002

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Device record: <https://www.510kdatabase.net/k830877/>. Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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