

**K960170 ELECTROENCEPHALOGRAPH (EEG)**Mar 1, 1996  
45 days to decisionK960170 · Product code: **GWF** · Neurology  
Source: <https://www.510kdatabase.net/k960170/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Stimulator, Electrical, Evoked Response (GWF)
Date received	Jan 16, 1996
Decision date	Mar 1, 1996
Days to decision	45 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Moberg Medical, Inc.</b>
Location	Ambler, PA, US
Contact	LARRY ENGLE
510(k) history	2 submissions · 2 cleared · 1992-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k960170/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 30, 2026