

**K122879 EEGER4 MODEL 4.3**Feb 6, 2013  
140 days to decisionK122879 · Product code: **HCC** · Neurology  
Source: <https://www.510kdatabase.net/k122879/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Biofeedback (HCC)
Date received	Sep 19, 2012
Decision date	Feb 6, 2013
Days to decision	140 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Eeg Software, LLC</b>
Location	Northridge, CA, US
Contact	HOWARD LIGHTSTONE
510(k) history	1 submissions · 1 cleared · 2013-2013

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k122879/> 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 9, 2026