

**K853608 NEUROMETER**Jun 12, 1986  
288 days to decisionK853608 · Product code: **LLN** · Neurology  
Source: <https://www.510kdatabase.net/k853608/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Vibration Threshold Measurement (LLN)
Date received	Aug 28, 1985
Decision date	Jun 12, 1986
Days to decision	288 days
Third-party review	No

**APPLICANT**

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Company	<b>Neurotron, Inc.</b>
Location	Lawrenceville, NJ, US
Contact	JEFFERSON KATIMS
510(k) history	2 submissions · 2 cleared · 1985-1986

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k853608/>; 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 July 1, 2026