

**K131568 SUDO PATH**Jun 28, 2013  
29 days to decisionK131568 · Product code: **GZO** · Neurology  
Source: <https://www.510kdatabase.net/k131568/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Device, Galvanic Skin Response Measurement (GZO)
Date received	May 30, 2013
Decision date	Jun 28, 2013
Days to decision	29 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Ld Technology, LLC</b>
Location	Miami, FL, US
Contact	ALBERT MAAREK
510(k) history	14 submissions · 14 cleared · 2009-2020

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k131568/> 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 19, 2026