

**K010897 MERIDIAN-PORTABLE**Jun 19, 2001  
85 days to decisionK010897 · Product code: **GZO** · Neurology  
Source: <https://www.510kdatabase.net/k010897/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Galvanic Skin Response Measurement (GZO)
Date received	Mar 26, 2001
Decision date	Jun 19, 2001
Days to decision	85 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Meridian Co., Ltd.</b>
Location	Songpa-Gu, Seoul, KR
Contact	SOO-RANG LEE
510(k) history	7 submissions · 7 cleared · 2000-2011

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k010897/> 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