

**K122944 GEMORE TRUE SINE INTERFERENTIAL STIMULATOR**Dec 21, 2012  
88 days to decisionK122944 · Product code: **LIH** · Neurology  
Source: <https://www.510kdatabase.net/k122944/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Interferential Current Therapy (LIH)
Date received	Sep 24, 2012
Decision date	Dec 21, 2012
Days to decision	88 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Gemore Technology Co, Ltd.</b>
Location	Tan Shui, Taipei Hsien, TW
Contact	BODEN S.P. LAI
510(k) history	14 submissions · 14 cleared · 2003-2019

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k122944/> 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 18, 2026