

**K121340 SOMNODENT G2**May 30, 2012  
27 days to decisionK121340 · Product code: **LRK** · Dental  
Source: <https://www.510kdatabase.net/k121340/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Anti-snoring (LRK)
Date received	May 3, 2012
Decision date	May 30, 2012
Days to decision	27 days
Third-party review	Yes
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Somnomed, Inc.</b>
Location	Denton, TX, US
Contact	KATHRYN JAYNE
510(k) history	10 submissions · 10 cleared · 2008-2024

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k121340/> 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 May 26, 2026