

**K111110 APNEA GUARD**Jul 22, 2011  
93 days to decisionK111110 · Product code: **LRK** · Dental  
Source: <https://www.510kdatabase.net/k111110/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Anti-snoring (LRK)
Date received	Apr 20, 2011
Decision date	Jul 22, 2011
Days to decision	93 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Advanced Brain Monitoring, Inc.</b>
Location	Carlsbad, CA, US
Contact	PAUL DRYDEN
510(k) history	14 submissions · 14 cleared · 2004-2016

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k111110/> 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 24, 2026