

**K160234 OVENT**Mar 17, 2016  
45 days to decisionK160234 · Product code: **LRK** · Dental  
Source: <https://www.510kdatabase.net/k160234/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Anti-snoring (LRK)
Date received	Feb 1, 2016
Decision date	Mar 17, 2016
Days to decision	45 days
Third-party review	Yes
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Oventus Manufacturing Pty, Ltd.</b>
Location	Brisbane, AU
Contact	Neil Anderson
510(k) history	4 submissions · 4 cleared · 2016-2019

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k160234/> 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 28, 2026