

**K932932 SNOREMASTER SNORE REMEDY**Feb 10, 1995  
604 days to decisionK932932 · Product code: **LQZ** · Dental  
Source: <https://www.510kdatabase.net/k932932/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Jaw Repositioning (LQZ)
Date received	Jun 16, 1993
Decision date	Feb 10, 1995
Days to decision	604 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>The Snoremaster Co.</b>
Location	Philadelphia, PA, US
Contact	LEROY S FORNEY
510(k) history	1 submissions · 1 cleared · 1995-1995

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k932932/> 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 July 3, 2026