

**K253868 QuietLab Pro**Dec 5, 2025  
2 days to decisionK253868 · Product code: **LRK** · Dental  
Source: <https://www.510kdatabase.net/k253868/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Anti-snoring (LRK)
Date received	Dec 3, 2025
Decision date	Dec 5, 2025
Days to decision	2 days
Third-party review	Yes
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>QuietLab, LLC</b>
Location	New York, NY, US
Contact	Candriella Pietro
510(k) history	1 submissions · 1 cleared · 2025-2025

**REGULATORY CONSULTANT**

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Consulting firm	<b>Beanstock Consulting</b>
Contact	Eileen Heller

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k253868/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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