

K234089 DNA ApplianceSep 16, 2024
269 days to decisionK234089 · Product code: **LRK** · Dental
Source: <https://www.510kdatabase.net/k234089/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Anti-snoring (LRK)
Date received	Dec 22, 2023
Decision date	Sep 16, 2024
Days to decision	269 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Vivos Therapeutics
Location	Murray, KY, US
Contact	Kimberly Griffith
510(k) history	1 submissions · 1 cleared · 2024-2024

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k234089/> Data sourced from FDA 510(k) public records (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. - Generated May 13, 2026