

K233850 Difiney Anti Snoring DeviceApr 18, 2024
135 days to decisionK233850 · Product code: **LRK** · Dental
Source: <https://www.510kdatabase.net/k233850/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Anti-snoring (LRK)
Date received	Dec 5, 2023
Decision date	Apr 18, 2024
Days to decision	135 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Fissiontech, LLC
Location	New York, NY, US
Contact	Mike Lee
510(k) history	3 submissions · 3 cleared · 2024-2025

REGULATORY CONSULTANT

Consulting firm	Shanghai Spica Management Consulting Co., Ltd.
Contact	Libray Chang

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k233850/> 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