

K201719 Vital SleepJan 14, 2021
205 days to decisionK201719 · Product code: **LRK** · Dental
Source: <https://www.510kdatabase.net/k201719/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Device, Anti-snoring (LRK)
Date received	Jun 23, 2020
Decision date	Jan 14, 2021
Days to decision	205 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	The Snore Reliever Company, LLC
Location	Laguna Niguel, CA, US
Contact	David Hernandez
510(k) history	2 submissions · 2 cleared · 2010-2021

REGULATORY CONSULTANT

Consulting firm	Devine Guidance International, Inc.
Contact	Christopher J. Devine

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

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