

K182202 SLEEPAPPJan 11, 2019
150 days to decisionK182202 · Product code: **LRK** · Dental
Source: <https://www.510kdatabase.net/k182202/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Device, Anti-snoring (LRK)
Date received	Aug 14, 2018
Decision date	Jan 11, 2019
Days to decision	150 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Statement

APPLICANT

Company	Comfort Acrylics, Inc.
Location	Camas, WA, US
Contact	Richard W. Moore
510(k) history	1 submissions · 1 cleared · 2019-2019

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k182202/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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