

K132506 SLEEPTIGHT MOUTHPIECE, SLEEP PRO, QUIETNITE, ALTITUDE MOUTHPIECE, SLEEP EASY

May 29, 2014
290 days to decision

K132506 · Product code: **LRK** · Dental
Source: <https://www.510kdatabase.net/k132506/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Anti-snoring (LRK)
Date received	Aug 12, 2013
Decision date	May 29, 2014
Days to decision	290 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Michael D Williams Dds PA
Location	Davie, FL, US
Contact	MICHAEL D WILLIAMS
510(k) history	2 submissions · 2 cleared · 2014-2019

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k132506/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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