

K182661 Kava and Kava with HerbstFeb 13, 2019
141 days to decisionK182661 · Product code: **LRK** · Dental
Source: <https://www.510kdatabase.net/k182661/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Anti-snoring (LRK)
Date received	Sep 25, 2018
Decision date	Feb 13, 2019
Days to decision	141 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Sketchpad Innovations, LLC
Location	San Diego, CA, US
Contact	Phillip "Sonnie" Bocala
510(k) history	1 submissions · 1 cleared · 2019-2019

REGULATORY CONSULTANT

Consulting firm	Fishermed Consulting, LLC
Contact	Cheryl Fisher

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k182661/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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