FDA 510(k) Premarket Notification

K182312 Zyppah Anti-Snoring Device

CLEARED

Jan 24, 2019
150 days to decision

K182312 · Product code: **LRK** · Dental
Source: https://www.510kdatabase.net/k182312/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Anti-snoring (LRK)
Date received	Aug 27, 2018
Decision date	Jan 24, 2019
Days to decision	150 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Always More Marketing**
Location	Calabasas, CA, US
Contact	Jonathan Greenburg
510(k) history	1 submissions · 1 cleared · 2019-2019

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k182312/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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