FDA 510(k) Premarket Notification

K121285 ORAVAN OSA

CLEARED

Mar 8, 2013
312 days to decision

K121285 · Product code: **LRK** · Dental
Source: https://www.510kdatabase.net/k121285/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Anti-snoring (LRK)
Date received	Apr 30, 2012
Decision date	Mar 8, 2013
Days to decision	312 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**New Jersey Snoring Solutions**
Location	West Orange, NJ, US
Contact	DEBORAH STEIN
510(k) history	1 submissions · 1 cleared · 2013-2013

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k121285/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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