FDA 510(k) Premarket Notification

K121340 SOMNODENT G2

CLEARED

May 30, 2012
27 days to decision

K121340 · Product code: **LRK** · Dental
Source: https://www.510kdatabase.net/k121340/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Anti-snoring (LRK)
Date received	May 3, 2012
Decision date	May 30, 2012
Days to decision	27 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	**Somnomed, Inc.**
Location	Denton, TX, US
Contact	KATHRYN JAYNE
510(k) history	10 submissions · 10 cleared · 2008-2024