FDA 510(k) Premarket Notification

K111110 APNEA GUARD

CLEARED

Jul 22, 2011
93 days to decision

K111110 · Product code: **LRK** · Dental
Source: https://www.510kdatabase.net/k111110/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Anti-snoring (LRK)
Date received	Apr 20, 2011
Decision date	Jul 22, 2011
Days to decision	93 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Advanced Brain Monitoring, Inc.**
Location	Carlsbad, CA, US
Contact	PAUL DRYDEN
510(k) history	14 submissions · 14 cleared · 2004-2016