FDA 510(k) Premarket Notification

K022891 THE BREATHE EZ ANTI-SNORING DEVICE

CLEARED

Feb 19, 2003
173 days to decision

K022891 · Product code: **LRK** · Dental
Source: https://www.510kdatabase.net/k022891/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Anti-snoring (LRK)
Date received	Aug 30, 2002
Decision date	Feb 19, 2003
Days to decision	173 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**D&S Redhage**
Location	New Haven, MO, US
Contact	DANIEL J REDHAGE
510(k) history	1 submissions · 1 cleared · 2003-2003