

**K022891 THE BREATHE EZ ANTI-SNORING DEVICE**

Feb 19, 2003  
 173 days to decision

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

**SUBMISSION DETAILS**

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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**

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Company	<b>D&amp;S Redhage</b>
Location	New Haven, MO, US
Contact	DANIEL J REDHAGE
510(k) history	1 submissions · 1 cleared · 2003-2003

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)  
 Device record: <https://www.510kdatabase.net/k022891/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).  
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