

**K011723 ANTI-SNORING DEVICE OR ASD**Dec 18, 2002  
562 days to decisionK011723 · Product code: **LRK** · Dental  
Source: <https://www.510kdatabase.net/k011723/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Anti-snoring (LRK)
Date received	Jun 4, 2001
Decision date	Dec 18, 2002
Days to decision	562 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Pi Medical</b>
Location	Minneapolis, MN, US
Contact	LINDA S ALEXANDER
510(k) history	1 submissions · 1 cleared · 2002-2002

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