

**K183443 SomnoDent Avant**May 8, 2019  
147 days to decisionK183443 · Product code: **LRK** · Dental  
Source: <https://www.510kdatabase.net/k183443/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Device, Anti-snoring (LRK)
Date received	Dec 12, 2018
Decision date	May 8, 2019
Days to decision	147 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Somnomed, Inc.</b>
Location	Denton, TX, US
Contact	Dorene Markwiese
510(k) history	10 submissions · 10 cleared · 2008-2024

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k183443/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 26, 2026