

**K172859 Prosomnus [CA] Sleep and Snore Device and  
Prosomnus [CA] Sleep and Snore Device with Micro-recorder**Nov 22, 2017  
63 days to decisionK172859 · Product code: PLC · Dental  
Source: <https://www.510kdatabase.net/k172859/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Sleep Appliances With Patient Monitoring (PLC)
Date received	Sep 20, 2017
Decision date	Nov 22, 2017
Days to decision	63 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Prosomnous Sleep Technologies</b>
Location	Pleasanton, CA, US
Contact	David Kuhns
510(k) history	1 submissions · 1 cleared · 2017-2017

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k172859/>; 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 31, 2026