

**K202499 Diamond Digital Sleep Orthotic (DDSO)**Sep 3, 2020  
3 days to decisionK202499 · Product code: **LRK** · Dental  
Source: <https://www.510kdatabase.net/k202499/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Anti-snoring (LRK)
Date received	Aug 31, 2020
Decision date	Sep 3, 2020
Days to decision	3 days
Third-party review	Yes
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Diamond Orthotic Laboratory</b>
Location	La Mesa, CA, US
Contact	Matt Rago
510(k) history	1 submissions · 1 cleared · 2020-2020

**REGULATORY CONSULTANT**

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Consulting firm	<b>Third Party Review Group, LLC</b>
Contact	Dave Yungvurt

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k202499/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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