

**K203462 Advanced Dental Appliance**Feb 19, 2021  
87 days to decisionK203462 · Product code: **LRK** · Dental  
Source: <https://www.510kdatabase.net/k203462/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Device, Anti-snoring (LRK)
Date received	Nov 24, 2020
Decision date	Feb 19, 2021
Days to decision	87 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Dasoft Partners</b>
Location	San Diego, CA, US
Contact	Dante Togliatti
510(k) history	2 submissions · 2 cleared · 2018-2021

**REGULATORY CONSULTANT**

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Consulting firm	<b>J Blane, LLC</b>
Contact	Kimberly Lane

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/k203462/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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