

**K192429 Dental Pain Eraser**Jan 14, 2020  
131 days to decisionK192429 · Product code: **LWM** · Dental  
Source: <https://www.510kdatabase.net/k192429/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Electrical Dental Anesthesia (LWM)
Date received	Sep 5, 2019
Decision date	Jan 14, 2020
Days to decision	131 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Synapse Dental</b>
Location	Cranston, RI, US
Contact	Cosmo Haralambidis
510(k) history	2 submissions · 2 cleared · 2020-2020

**REGULATORY CONSULTANT**

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Consulting firm	<b>Experien Group, LLC</b>
Contact	Valerie Defiesta-NG

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/k192429/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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