

**K202942 Straumann 4 mm Short Implants**Feb 8, 2021  
131 days to decisionK202942 · Product code: **DZE** · Dental  
Source: <https://www.510kdatabase.net/k202942/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Sep 30, 2020
Decision date	Feb 8, 2021
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>Straumann USA, LLC</b>
Location	Andover, MA, US
Contact	Chanrasmey N. White
510(k) history	24 submissions · 24 cleared · 2012-2024

**REGULATORY CONSULTANT**

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Consulting firm	<b>Straumann USA, Lls</b>
Contact	Jennifer M. Jackson

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/k202942/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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