

**K181703 Straumann BLX Line Extension - Implants, SRAs and Anatomic Abutments**Dec 28, 2018  
184 days to decisionK181703 · Product code: **DZE** · Dental  
Source: <https://www.510kdatabase.net/k181703/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Jun 27, 2018
Decision date	Dec 28, 2018
Days to decision	184 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Institut Straumann AG</b>
Location	Basel, CH
Contact	Viviana Horhoiu
Website	<a href="https://www.straumann.com">https://www.straumann.com</a>
510(k) history	90 submissions · 90 cleared · 1990-2026

**REGULATORY CONSULTANT**

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

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA [accessdata.fda.gov](https://accessdata.fda.gov)510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k181703/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://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](https://accessdata.fda.gov). - Generated June 28, 2026