

K173961 Straumann BLX Implant SystemJun 5, 2018
159 days to decisionK173961 · Product code: **DZE** · DentalSource: <https://www.510kdatabase.net/k173961/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Dec 28, 2017
Decision date	Jun 5, 2018
Days to decision	159 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Institut Straumann AG
Location	Basel, CH
Contact	Viviana Horhoiu
Website	https://www.straumann.com
510(k) history	88 submissions · 88 cleared · 1990-2026

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k173961/> Data sourced from FDA 510(k) public records (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. - Generated May 30, 2026