

**K100100 DIO STEADY EXTERNAL IMPLANT SYSTEM**Nov 10, 2010  
302 days to decisionK100100 · Product code: **DZE** · Dental  
Source: <https://www.510kdatabase.net/k100100/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Jan 12, 2010
Decision date	Nov 10, 2010
Days to decision	302 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Dio Corporation</b>
Location	Los Angeles, CA, US
Contact	TIMOTHY LEE
510(k) history	14 submissions · 14 cleared · 2010-2023

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k100100/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 26, 2026