

**K133592 NEODENT IMPLANT SYSTEM, TITAMAX TI EX ACQUA  
AND TI DRIVE ACQUA, TITAMAX SMART EX ACQUA AND  
DRIVE SMART ACQUA, TITAMAX CM**Aug 22, 2014  
273 days to decisionK133592 · Product code: **DZE** · Dental  
Source: <https://www.510kdatabase.net/k133592/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Nov 22, 2013
Decision date	Aug 22, 2014
Days to decision	273 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Neodent USA, Inc.</b>
Location	Andover, MA, US
Contact	CHRISTOPHER KLACZYK
510(k) history	1 submissions · 1 cleared · 2014-2014

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k133592/> 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 June 6, 2026