

**K073067 ANKYLOS DENTAL IMPLANT SYSTEM/CLAIMS**Feb 15, 2008  
108 days to decisionK073067 · Product code: **DZE** · DentalSource: <https://www.510kdatabase.net/k073067/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Oct 30, 2007
Decision date	Feb 15, 2008
Days to decision	108 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Dentsply International, Inc.</b>
Location	York, PA, US
Contact	HELEN LEWIS
510(k) history	36 submissions · 36 cleared · 2006-2016

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k073067/> 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 25, 2026