

**K052369 EXFEEL IMPLANT SYSTEM**Jan 10, 2006  
134 days to decisionK052369 · Product code: **DZE** · DentalSource: <https://www.510kdatabase.net/k052369/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Aug 29, 2005
Decision date	Jan 10, 2006
Days to decision	134 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Megagen Co., Ltd.</b>
Location	Santa Fe Springs, CA, US
Contact	DAE KYU CHANG
510(k) history	9 submissions · 9 cleared · 2005-2011

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k052369/> 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 16, 2026