

**K033767 DUAL TOP ANCHOR SYSTEM SCREWS**Feb 24, 2004  
83 days to decisionK033767 · Product code: **DZE** · DentalSource: <https://www.510kdatabase.net/k033767/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Dec 3, 2003
Decision date	Feb 24, 2004
Days to decision	83 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Jeil Medical Corporation</b>
Location	Deer Field, IL, US
Contact	DANIEL KAMM
510(k) history	53 submissions · 53 cleared · 2002-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k033767/> 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 28, 2026