

**K022425 SCREWLINE**Aug 14, 2002  
21 days to decisionK022425 · Product code: **DZE** · DentalSource: <https://www.510kdatabase.net/k022425/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Jul 24, 2002
Decision date	Aug 14, 2002
Days to decision	21 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Altatec Biotechnologies N.A., Inc.</b>
Location	Potomac, MD, US
Contact	ROBERT J SLOMOFF
510(k) history	1 submissions · 1 cleared · 2002-2002

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k022425/> 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 July 1, 2026