

**K000099 CAMLOG SCREW IMPLANT SYSTEM**Jan 28, 2000  
15 days to decisionK000099 · Product code: **DZE** · Dental  
Source: <https://www.510kdatabase.net/k000099/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Jan 13, 2000
Decision date	Jan 28, 2000
Days to decision	15 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Altatec Biotechnologies</b>
Location	Potomac, MD, US
Contact	R.J. SLOMOFF
510(k) history	4 submissions · 4 cleared · 2000-2003

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k000099/> 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