

**K090174 TSI**Sep 14, 2009  
234 days to decisionK090174 · Product code: **DZE** · DentalSource: <https://www.510kdatabase.net/k090174/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Jan 23, 2009
Decision date	Sep 14, 2009
Days to decision	234 days
Third-party review	No
Summary / Statement	Summary
Other names	ERI

**APPLICANT**

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Company	<b>Oco Biomedical</b>
Location	Sterling, VA, US
Contact	JACK BLOOM
510(k) history	4 submissions · 4 cleared · 2008-2013

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k090174/>, 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 15, 2026