

**K953836 ABUTMENT**Sep 21, 1995  
37 days to decisionK953836 · Product code: **DZE** · DentalSource: <https://www.510kdatabase.net/k953836/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Aug 15, 1995
Decision date	Sep 21, 1995
Days to decision	37 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Steri-Oss, Inc.</b>
Location	Anaheim, CA, US
Contact	DON KENNARD
510(k) history	46 submissions · 46 cleared · 1992-1998

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k953836/> 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 May 20, 2026