

K964739 STERI-OSS TIODIZED SCREWMar 31, 1997
125 days to decisionK964739 · Product code: **DZE** · Dental
Source: <https://www.510kdatabase.net/k964739/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Nov 26, 1996
Decision date	Mar 31, 1997
Days to decision	125 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Steri-Oss, Inc.
Location	Anaheim, CA, US
Contact	JEFF HAUSHEER, PH.D.
510(k) history	46 submissions · 46 cleared · 1992-1998

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k964739/> Data sourced from FDA 510(k) public records (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. - Generated May 20, 2026