FDA 510(k) Premarket Notification

K964739 STERI-OSS TIODIZED SCREW

CLEARED

Mar 31, 1997
125 days to decision

K964739 · 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