

K070569 SM INTERNAL./EXTERNAL IMPLANT SYSTEMNov 8, 2007
253 days to decisionK070569 · Product code: **DZE** · DentalSource: <https://www.510kdatabase.net/k070569/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Feb 28, 2007
Decision date	Nov 8, 2007
Days to decision	253 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Dio Department Dsi, Inc.
Location	Santa Fe Springs, CA, US
Contact	KENNY LIM
510(k) history	8 submissions · 8 cleared · 2006-2008

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k070569/> 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 June 25, 2026