

K060500 SPEEDY ORTHODONTIC SCREWJun 16, 2006
112 days to decisionK060500 · Product code: **DZE** · Dental
Source: <https://www.510kdatabase.net/k060500/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Feb 24, 2006
Decision date	Jun 16, 2006
Days to decision	112 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Dentium Co., Ltd.
Location	Dunwoody, GA, US
Contact	CATHRYN N CAMBRIA
510(k) history	22 submissions · 22 cleared · 2004-2022

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k060500/> 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 27, 2026