

K042339 NEOPLANT IMPLANT SYSTEMNov 2, 2004
64 days to decisionK042339 · Product code: **DZE** · Dental
Source: <https://www.510kdatabase.net/k042339/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Aug 30, 2004
Decision date	Nov 2, 2004
Days to decision	64 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Neobiotech Co., Ltd.
Location	Santa Fe Springs, CA, US
Contact	DAE KYU CHANG
510(k) history	17 submissions · 17 cleared · 2004-2024

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k042339/> 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 26, 2026