

K042289 MDI ORTHOSep 15, 2004
22 days to decisionK042289 · Product code: **DZE** · DentalSource: <https://www.510kdatabase.net/k042289/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Aug 24, 2004
Decision date	Sep 15, 2004
Days to decision	22 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Intec Corp.
Location	Ardmore, OK, US
Contact	J. BRAD VANCE
510(k) history	28 submissions · 28 cleared · 1991-2008

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k042289/> 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 18, 2026