

K955492 INTERPORE IMZ CYL IMPLANT & CYL HEX IMPLTJan 30, 1996
60 days to decisionK955492 · Product code: **DZE** · Dental
Source: <https://www.510kdatabase.net/k955492/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Dec 1, 1995
Decision date	Jan 30, 1996
Days to decision	60 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Interpore Intl.
Location	Walker, MI, US
Contact	DAVID P BALDING
510(k) history	25 submissions · 25 cleared · 1984-1998

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k955492/> 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 21, 2026