

**K960371 INTERPORE THREADED IMPLANT**Mar 8, 1996  
42 days to decisionK960371 · Product code: **DZE** · Dental  
Source: <https://www.510kdatabase.net/k960371/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Jan 26, 1996
Decision date	Mar 8, 1996
Days to decision	42 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k960371/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 21, 2026