

K922732 INTERPORE CYLINDRICAL HEX IMPLANT, 3.3 MM DIAMETER

Feb 10, 1994
612 days to decision

K922732 · Product code: **DZE** · Dental
Source: <https://www.510kdatabase.net/k922732/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Implant, Endosseous, Root-form (DZE)
Date received	Jun 8, 1992
Decision date	Feb 10, 1994
Days to decision	612 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k922732/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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