FDA 510(k) Premarket Notification

K960371 INTERPORE THREADED IMPLANT

CLEARED

Mar 8, 1996
42 days to decision

K960371 · Product code: **DZE** · Dental
Source: https://www.510kdatabase.net/k960371/

SUBMISSION DETAILS

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

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