FDA 510(k) Premarket Notification

# K955492 INTERPORE IMZ CYL IMPLANT & CYL HEX IMPLT

**CLEARED**

Jan 30, 1996
60 days to decision

K955492 · Product code: **DZE** · Dental
Source: https://www.510kdatabase.net/k955492/

## SUBMISSION DETAILS

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|---|---|
| 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

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