

**K904611 INTERPORE CYLINDRICAL HEX IMPLANT**Feb 5, 1991  
117 days to decisionK904611 · Product code: **DZE** · DentalSource: <https://www.510kdatabase.net/k904611/>**SUBMISSION DETAILS**

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|-----------------------|--------------------------------------|
| Decision              | Substantially Equivalent (Cleared)   |
| Submission type       | Traditional                          |
| Device classification | Implant, Endosseous, Root-form (DZE) |
| Date received         | Oct 11, 1990                         |
| Decision date         | Feb 5, 1991                          |
| Days to decision      | 117 days                             |
| Third-party review    | No                                   |

**APPLICANT**

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|----------------|---|
| Company        | <b>Interpore Intl.</b>                  |
| Location       | Walker, MI, US                          |
| Contact        | A CONNER                                |
| 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/k904611/>, 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