

**K955884 COORDINATE REVISION KNEE SYSTEM**Mar 13, 1996  
76 days to decisionK955884 · Product code: **JWH** · Orthopedic  
Source: <https://www.510kdatabase.net/k955884/>**SUBMISSION DETAILS**

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|-----------------------|--|
| Decision              | Substantially Equivalent (Cleared)   |
| Submission type       | Traditional  |
| Device classification | Prosthesis, Knee, Patellofemorotibial, Semi-constrained, Cemented, Polymer/metal/polymer (JWH) |
| Date received         | Dec 28, 1995   |
| Decision date         | Mar 13, 1996   |
| Days to decision      | 76 days  |
| Third-party review    | No   |
| Summary / Statement   | Summary  |

**APPLICANT**

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|----------------|---|
| Company        | <b>Depuy, Inc.</b>                        |
| Location       | Mchenry, IL, US                           |
| Contact        | CHERYL HASTINGS                           |
| 510(k) history | 303 submissions · 239 cleared · 1976-2005 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k955884/> 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 20, 2026