

**K000172 DEPUY CONTOUR UNICOMPARTMENTAL KNEE PROSTHESIS**Feb 3, 2000  
14 days to decisionK000172 · Product code: **HRY** · Orthopedic  
Source: <https://www.510kdatabase.net/k000172/>**SUBMISSION DETAILS**

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|                       |   |
|-----------------------|---|
| Decision              | Substantially Equivalent (Cleared)  |
| Submission type       | Special   |
| Device classification | Prosthesis, Knee, Femorotibial, Semi-constrained, Cemented, Metal/polymer (HRY) |
| Date received         | Jan 20, 2000  |
| Decision date         | Feb 3, 2000   |
| Days to decision      | 14 days   |
| Third-party review    | No  |
| Summary / Statement   | Summary   |

**APPLICANT**

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|----------------|---|
| Company        | <b>DePuy Orthopaedics, Inc.</b>           |
| Location       | Warsaw, IN, US                            |
| Contact        | CHERYL HASTINGS                           |
| 510(k) history | 206 submissions · 204 cleared · 1998-2023 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k000172/>; 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 26, 2026