

**K922008 CDH HIP SYSTEM**Jul 30, 1992  
91 days to decisionK922008 · Product code: **JDI** · Orthopedic  
Source: <https://www.510kdatabase.net/k922008/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Hip, Semi-constrained, Metal/polymer, Cemented (JDI)
Date received	Apr 30, 1992
Decision date	Jul 30, 1992
Days to decision	91 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Howmedica, Pfizer Medical Technology Group</b>
Location	Rutherford, NJ, US
Contact	ROBERT E SMITH
510(k) history	1 submissions · 1 cleared · 1992-1992

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k922008/> 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 22, 2026