FDA 510(k) Premarket Notification

# K922008 CDH HIP SYSTEM

**CLEARED**

Jul 30, 1992
91 days to decision

K922008 · 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 | **Howmedica, Pfizer Medical Technology Group** |
| Location | Rutherford, NJ, US |
| Contact | ROBERT E SMITH |
| 510(k) history | 1 submissions · 1 cleared · 1992-1992 |