

**K973121 INTERAX TOTAL KNEE SYSTEM**Nov 18, 1997  
90 days to decisionK973121 · Product code: **JWH** · Orthopedic  
Source: <https://www.510kdatabase.net/k973121/>**SUBMISSION DETAILS**

---

|                       |  |
|-----------------------|--|
| Decision              | Substantially Equivalent (Cleared)   |
| Submission type       | Traditional  |
| Device classification | Prosthesis, Knee, Patellofemorotibial, Semi-constrained, Cemented, Polymer/metal/polymer (JWH) |
| Date received         | Aug 20, 1997   |
| Decision date         | Nov 18, 1997   |
| Days to decision      | 90 days  |
| Third-party review    | No   |
| Summary / Statement   | Summary  |

**APPLICANT**

---

|                |   |
|----------------|---|
| Company        | <b>Howmedica Corp.</b>                    |
| Location       | Mchenry, IL, US                           |
| Contact        | MARGARET F CROWE                          |
| 510(k) history | 373 submissions · 325 cleared · 1976-1998 |

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k973121/> 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 14, 2026