

K790350 POSTERIOR CRUCIATE CONDYLAR TIBIALMar 6, 1979
13 days to decisionK790350 · Product code: **HRY** · Orthopedic
Source: <https://www.510kdatabase.net/k790350/>**SUBMISSION DETAILS**

| | |
|-----------------------|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Prosthesis, Knee, Femorotibial, Semi-constrained, Cemented, Metal/polymer (HRY) |
| Date received | Feb 21, 1979 |
| Decision date | Mar 6, 1979 |
| Days to decision | 13 days |
| Third-party review | No |

APPLICANT

| | |
|----------------|---|
| Company | Howmedica Corp. |
| Location | Mchenry, IL, US |
| 510(k) history | 373 submissions · 325 cleared · 1976-1998 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k790350/> Data sourced from FDA 510(k) public records (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. - Generated May 14, 2026