

K770184 BI-CENTRIC ENDOPROSTHESESFeb 8, 1977
12 days to decisionK770184 · Product code: **JDG** · Orthopedic
Source: <https://www.510kdatabase.net/k770184/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Hip, Femoral Component, Cemented, Metal (JDG)
Date received	Jan 27, 1977
Decision date	Feb 8, 1977
Days to decision	12 days
Third-party review	No

APPLICANT

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

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Device record: <https://www.510kdatabase.net/k770184/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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