

K881983 HOWMEDICA PRECISION OSTEOLOCK FEMORAL COMPONENTOct 6, 1988
147 days to decisionK881983 · Product code: **LWJ** · Orthopedic
Source: <https://www.510kdatabase.net/k881983/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Hip, Semi-constrained, Metal/polymer, Uncemented (LWJ)
Date received	May 12, 1988
Decision date	Oct 6, 1988
Days to decision	147 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k881983/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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