

**K933875 PREMISE CEMENTED FEMORAL STEM, ADDITION
STYLE**Mar 1, 1994
204 days to decisionK933875 · Product code: JDI · Orthopedic
Source: <https://www.510kdatabase.net/k933875/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Hip, Semi-constrained, Metal/polymer, Cemented (JDI)
Date received	Aug 9, 1993
Decision date	Mar 1, 1994
Days to decision	204 days
Third-party review	No
Summary / Statement	Summary

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/k933875/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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