

**K925901 KINEMAX(R) CONDYLAR TOTAL KNEE SYST
FEMORAL SPACER**Mar 22, 1993
122 days to decisionK925901 · Product code: **JWH** · Orthopedic
Source: <https://www.510kdatabase.net/k925901/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Knee, Patellofemorotibial, Semi-constrained, Cemented, Polymer/metal/polymer (JWH)
Date received	Nov 20, 1992
Decision date	Mar 22, 1993
Days to decision	122 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/k925901/>, Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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