FDA 510(k) Premarket Notification

K925901 KINEMAX(R) CONDYLAR TOTAL KNEE SYST FEMORAL SPACER

CLEARED

Mar 22, 1993
122 days to decision

K925901 · 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