FDA 510(k) Premarket Notification

K973121 INTERAX TOTAL KNEE SYSTEM

CLEARED

Nov 18, 1997
90 days to decision

K973121 · Product code: **JWH** · Orthopedic
Source: https://www.510kdatabase.net/k973121/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Knee, Patellofemorotibial, Semi-constrained, Cemented, Polymer/metal/polymer (JWH)
Date received	Aug 20, 1997
Decision date	Nov 18, 1997
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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