

K970713 COMPASS UNIVERSAL HINGEApr 3, 1997
35 days to decisionK970713 · Product code: **JDW** · Orthopedic
Source: <https://www.510kdatabase.net/k970713/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Pin, Fixation, Threaded (JDW)
Date received	Feb 27, 1997
Decision date	Apr 3, 1997
Days to decision	35 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Smith & Nephew, Inc., Orthopaedic Div.
Location	Memphis, TN, US
Contact	JO ANN KUHNE
510(k) history	10 submissions · 8 cleared · 1996-1999

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k970713/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 21, 2026