

K932210 ORTHOLOC ADVANTIM REVISION FEMORAL COMPONENTSep 7, 1994
488 days to decisionK932210 · Product code: **JWH** · Orthopedic
Source: <https://www.510kdatabase.net/k932210/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent - SN
Submission type	Traditional
Device classification	Prosthesis, Knee, Patellofemorotibial, Semi-constrained, Cemented, Polymer/metal/polymer (JWH)
Date received	May 7, 1993
Decision date	Sep 7, 1994
Days to decision	488 days
Third-party review	No

APPLICANT

Company	Dow Corning Wright
Location	Mchenry, IL, US
Contact	DIANN ATCHISON
510(k) history	74 submissions · 52 cleared · 1979-1994

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k932210/> 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 22, 2026