

**K914169 WHITESIDE ORTHOLOC MOD.TIBIAL
AUGMENTATION COMP**Jan 27, 1992
132 days to decisionK914169 · Product code: **JWH** · Orthopedic
Source: <https://www.510kdatabase.net/k914169/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Knee, Patellofemorotibial, Semi-constrained, Cemented, Polymer/metal/polymer (JWH)
Date received	Sep 17, 1991
Decision date	Jan 27, 1992
Days to decision	132 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k914169/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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