

**K915525 WHITESIDE ORTHOLOC(R) MODULAR REVISION  
FEMOR COMP**May 5, 1993  
512 days to decisionK915525 · Product code: **JWH** · Orthopedic  
Source: <https://www.510kdatabase.net/k915525/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Knee, Patellofemorotibial, Semi-constrained, Cemented, Polymer/metal/polymer (JWH)
Date received	Dec 10, 1991
Decision date	May 5, 1993
Days to decision	512 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k915525/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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