

**K902538 WHITESIDE ORTHOLOC MODULAR TIBIAL AUGMENTATION**Jul 6, 1990  
30 days to decisionK902538 · Product code: HRY · Orthopedic  
Source: <https://www.510kdatabase.net/k902538/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent - SN
Submission type	Traditional
Device classification	Prosthesis, Knee, Femorotibial, Semi-constrained, Cemented, Metal/polymer (HRY)
Date received	Jun 6, 1990
Decision date	Jul 6, 1990
Days to decision	30 days
Third-party review	No

**APPLICANT**

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Company	<b>Dow Corning Wright</b>
Location	Mchenry, IL, US
Contact	DEERE 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/k902538/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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