

**K862136 MODIFIED WHITESIDE POSTERIOR STABILIZED  
TOTAL KNEE**Jul 14, 1986  
40 days to decisionK862136 · Product code: **HSX** · Orthopedic  
Source: <https://www.510kdatabase.net/k862136/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Knee, Femorotibial, Non-constrained, Cemented, Metal/polymer (HSX)
Date received	Jun 4, 1986
Decision date	Jul 14, 1986
Days to decision	40 days
Third-party review	No

**APPLICANT**

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Company	<b>Dow Corning Wright</b>
Location	Mchenry, IL, US
Contact	SPIRES, JR
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/k862136/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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