

**K881779 WHITESIDE ORTHOLOC II UNICONDYLAR KNEE SYSTEM**Aug 31, 1988  
127 days to decisionK881779 · Product code: **HSX** · Orthopedic  
Source: <https://www.510kdatabase.net/k881779/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent - SN
Submission type	Traditional
Device classification	Prosthesis, Knee, Femorotibial, Non-constrained, Cemented, Metal/polymer (HSX)
Date received	Apr 26, 1988
Decision date	Aug 31, 1988
Days to decision	127 days
Third-party review	No

**APPLICANT**

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Company	<b>Dow Corning Wright</b>
Location	Mchenry, IL, US
Contact	LIPSCOMB, PHD
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/k881779/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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