

**K982585 DEPUY AMK TOTAL KNEE SYSTEM DEPUY
COORDINATE ULTRA KNEE REVISION SYSTEM**Oct 5, 1998
73 days to decisionK982585 · Product code: **JWH** · Orthopedic
Source: <https://www.510kdatabase.net/k982585/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent - SN
Submission type	Traditional
Device classification	Prosthesis, Knee, Patellofemorotibial, Semi-constrained, Cemented, Polymer/metal/polymer (JWH)
Date received	Jul 24, 1998
Decision date	Oct 5, 1998
Days to decision	73 days
Third-party review	No

APPLICANT

Company	DePuy Orthopaedics, Inc.
Location	Warsaw, IN, US
Contact	SALLY FOUST
510(k) history	206 submissions · 204 cleared · 1998-2023

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k982585/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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