

**K925837 AMK LANDMARK REVISION KNEE FULL WEDGE
TIBIAL TRAY**Jan 7, 1994
415 days to decisionK925837 · Product code: **JWH** · Orthopedic
Source: <https://www.510kdatabase.net/k925837/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent - SN
Submission type	Traditional
Device classification	Prosthesis, Knee, Patellofemorotibial, Semi-constrained, Cemented, Polymer/metal/polymer (JWH)
Date received	Nov 18, 1992
Decision date	Jan 7, 1994
Days to decision	415 days
Third-party review	No

APPLICANT

Company	Depuy, Inc.
Location	Mchenry, IL, US
Contact	LYNNE A BROPHY
510(k) history	303 submissions · 239 cleared · 1976-2005

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k925837/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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