

K060525 VANGUARD REMOVABLE MOLDED POLY TIBIAMay 25, 2006
87 days to decisionK060525 · Product code: **JWH** · Orthopedic
Source: <https://www.510kdatabase.net/k060525/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Knee, Patellofemorotibial, Semi-constrained, Cemented, Polymer/metal/polymer (JWH)
Date received	Feb 27, 2006
Decision date	May 25, 2006
Days to decision	87 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Biomet Manufacturing Corp
Location	Warsaw, IN, US
Contact	TRACY BICKEL JOHNSON
510(k) history	93 submissions · 93 cleared · 2004-2023

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k060525/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 20, 2026