

K060303 VANGUARD PS OPEN BOX POROUS FEMORAL COMPONENTSMar 30, 2006
52 days to decisionK060303 · Product code: **MBH** · Orthopedic
Source: <https://www.510kdatabase.net/k060303/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Knee, Patello/femorotibial, Semi-constrained, Uncemented, Porous, Coated, Polymer/metal/polymer (MBH)
Date received	Feb 6, 2006
Decision date	Mar 30, 2006
Days to decision	52 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k060303/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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