

K030048 HA PMI FEMORAL HIP STEMFeb 5, 2003
30 days to decisionK030048 · Product code: **LPH** · Orthopedic
Source: <https://www.510kdatabase.net/k030048/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Prosthesis, Hip, Semi-constrained, Metal/polymer, Porous Uncemented (LPH)
Date received	Jan 6, 2003
Decision date	Feb 5, 2003
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Biomet Orthopedics, Inc.
Location	Warsaw, IN, US
Contact	PATRICIA BERES
510(k) history	34 submissions · 34 cleared · 2000-2008

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k030048/> 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 19, 2026