

K030971 MODIFICATION TO MODULAR PLUS REVISION STEMApr 24, 2003
27 days to decisionK030971 · Product code: **LWJ** · Orthopedic
Source: <https://www.510kdatabase.net/k030971/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Prosthesis, Hip, Semi-constrained, Metal/polymer, Uncemented (LWJ)
Date received	Mar 28, 2003
Decision date	Apr 24, 2003
Days to decision	27 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Plus Orthopedics
Location	San Diego, CA, US
Contact	J.D. WEBB
510(k) history	38 submissions · 38 cleared · 1997-2004

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k030971/> 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