

K962137 GENESIS II CONSTRAINED SYSTEMAug 2, 1996
60 days to decisionK962137 · Product code: **JWH** · Orthopedic
Source: <https://www.510kdatabase.net/k962137/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Knee, Patellofemorotibial, Semi-constrained, Cemented, Polymer/metal/polymer (JWH)
Date received	Jun 3, 1996
Decision date	Aug 2, 1996
Days to decision	60 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Smith & Nephew Richards, Inc.
Location	Memphus, TN, US
Contact	THOMAS L CRAIG
510(k) history	87 submissions · 72 cleared · 1990-1997

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k962137/> 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 21, 2026