FDA 510(k) Premarket Notification

K962137 GENESIS II CONSTRAINED SYSTEM

CLEARED

Aug 2, 1996
60 days to decision

K962137 · 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