FDA 510(k) Premarket Notification

K923807 P.F.C. MODULAR TOTAL KNEE SYSTEM, MODULAR PLUS

CLEARED

Nov 9, 1992
103 days to decision

K923807 · Product code: **JWH** · Orthopedic
Source: https://www.510kdatabase.net/k923807/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Knee, Patellofemorotibial, Semi-constrained, Cemented, Polymer/metal/polymer (JWH)
Date received	Jul 29, 1992
Decision date	Nov 9, 1992
Days to decision	103 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Johnson & Johnson International**
Location	New Brunswick, NJ, US
Contact	MARSHA J STONE
510(k) history	10 submissions · 10 cleared · 1989-1997