

K920034 DURACON MODULAR FEMORAL COMPONENT AND FEMORAL SPACMar 18, 1992
72 days to decisionK920034 · Product code: **JWH** · Orthopedic
Source: <https://www.510kdatabase.net/k920034/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Knee, Patellofemorotibial, Semi-constrained, Cemented, Polymer/metal/polymer (JWH)
Date received	Jan 6, 1992
Decision date	Mar 18, 1992
Days to decision	72 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Pfizer Hospital Products Group, Inc.
Location	Fall River, MA, US
Contact	MARGARET F CROWE
510(k) history	8 submissions · 7 cleared · 1991-1994

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