FDA 510(k) Premarket Notification

K932210 ORTHOLOC ADVANTIM REVISION FEMORAL COMPONENT

CLEARED

Sep 7, 1994
488 days to decision

K932210 · Product code: **JWH** · Orthopedic
Source: https://www.510kdatabase.net/k932210/

SUBMISSION DETAILS

Decision	Substantially Equivalent - SN
Submission type	Traditional
Device classification	Prosthesis, Knee, Patellofemorotibial, Semi-constrained, Cemented, Polymer/metal/polymer (JWH)
Date received	May 7, 1993
Decision date	Sep 7, 1994
Days to decision	488 days
Third-party review	No

APPLICANT

Company	**Dow Corning Wright**
Location	Mchenry, IL, US
Contact	DIANN ATCHISON
510(k) history	74 submissions · 52 cleared · 1979-1994