FDA 510(k) Premarket Notification

K932070 DURACON STABILIZER FEMORAL COMPONENT AND INSERT

CLEARED

Mar 22, 1995
693 days to decision

K932070 · Product code: **JWH** · Orthopedic
Source: https://www.510kdatabase.net/k932070/

SUBMISSION DETAILS

Decision	Substantially Equivalent - SN
Submission type	Traditional
Device classification	Prosthesis, Knee, Patellofemorotibial, Semi-constrained, Cemented, Polymer/metal/polymer (JWH)
Date received	Apr 28, 1993
Decision date	Mar 22, 1995
Days to decision	693 days
Third-party review	No

APPLICANT

Company	**Howmedica Corp.**
Location	Mchenry, IL, US
Contact	MARGARET F CROWE
510(k) history	373 submissions · 325 cleared · 1976-1998