FDA 510(k) Premarket Notification

K071413 HEMICAP PATELLO-FEMORAL RESURFACING SYSTEM

CLEARED

Nov 9, 2007
172 days to decision

K071413 · Product code: **KRR** · Orthopedic
Source: https://www.510kdatabase.net/k071413/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Knee, Patello/femoral, Semi-constrained, Cemented, Metal/polymer (KRR)
Date received	May 21, 2007
Decision date	Nov 9, 2007
Days to decision	172 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Arthrosurface, Inc.**
Location	Stoughton, MA, US
Contact	DAWN J WILSON
510(k) history	26 submissions · 26 cleared · 2004-2021