

**K071413 HEMICAP PATELLO-FEMORAL RESURFACING SYSTEM**Nov 9, 2007  
172 days to decisionK071413 · Product code: **KRR** · Orthopedic  
Source: <https://www.510kdatabase.net/k071413/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k071413/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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