

K063370 CAP 7.0MM MTP RESURFACING HEMI-ARTHROPLASTYJan 12, 2007
65 days to decisionK063370 · Product code: **KWD** · Orthopedic
Source: <https://www.510kdatabase.net/k063370/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Toe, Hemi-, Phalangeal (KWD)
Date received	Nov 8, 2006
Decision date	Jan 12, 2007
Days to decision	65 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k063370/>. Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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