

**K062960 CONSERVE FEMORAL RESURFACING COMPONENT**Dec 1, 2006  
63 days to decisionK062960 · Product code: **KXA** · Orthopedic  
Source: <https://www.510kdatabase.net/k062960/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Prosthesis, Hip, Femoral, Resurfacing (KXA)
Date received	Sep 29, 2006
Decision date	Dec 1, 2006
Days to decision	63 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Wrightmedicaltechnologyinc</b>
Location	Arlington, TN, US
Contact	THERESA LEISTER
510(k) history	302 submissions · 291 cleared · 1993-2023

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k062960/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 19, 2026