

**K072597 PRO-DENSE CORE DECOMPRESSION PROCEDURE  
KIT**Oct 15, 2007  
31 days to decisionK072597 · Product code: **MQV** · Orthopedic  
Source: <https://www.510kdatabase.net/k072597/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Filler, Bone Void, Calcium Compound (MQV)
Date received	Sep 14, 2007
Decision date	Oct 15, 2007
Days to decision	31 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Wrightmedicaltechnologyinc</b>
Location	Arlington, TN, US
Contact	RYAN M BELANEY
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/k072597/>. Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://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](https://accessdata.fda.gov). - Generated May 19, 2026