

**K090387 KAINOS+**Sep 28, 2009  
223 days to decisionK090387 · Product code: **MQV** · Orthopedic  
Source: <https://www.510kdatabase.net/k090387/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Filler, Bone Void, Calcium Compound (MQV)
Date received	Feb 17, 2009
Decision date	Sep 28, 2009
Days to decision	223 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Signus Medizintechnik GmbH</b>
Location	Minneapolis, MN, US
Contact	KAREN WARDEN
510(k) history	23 submissions · 22 cleared · 1999-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k090387/> 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 14, 2026