

**K024125 HALM ZIELKE INSTRUMENTATION SYSTEM**Sep 22, 2003  
280 days to decisionK024125 · Product code: **KWQ** · Orthopedic  
Source: <https://www.510kdatabase.net/k024125/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Spinal Intervertebral Body (KWQ)
Date received	Dec 16, 2002
Decision date	Sep 22, 2003
Days to decision	280 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Micomed Ortho GmbH</b>
Location	Lake Forest, CA, US
Contact	CORRINE M BONFIGLIO
510(k) history	2 submissions · 2 cleared · 2002-2003

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k024125/> 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 20, 2026