

**K982006 MICOMED - HALM ZIELKE INSTRUMENTATION**Jan 20, 1999  
226 days to decisionK982006 · Product code: **KWQ** · Orthopedic  
Source: <https://www.510kdatabase.net/k982006/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Spinal Intervertebral Body (KWQ)
Date received	Jun 8, 1998
Decision date	Jan 20, 1999
Days to decision	226 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Micomed GmbH</b>
Location	Fullerton, CA, US
Contact	CHARMAINE HENDERSON
510(k) history	2 submissions · 2 cleared · 1999-2000

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k982006/> 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 21, 2026