

**K141405 MOBIS II ST SPINAL IMPLANT**Sep 16, 2014  
111 days to decisionK141405 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k141405/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	May 28, 2014
Decision date	Sep 16, 2014
Days to decision	111 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Signus Medizintechnik GmbH</b>
Location	Minneapolis, MN, US
Contact	J.D. WEBB
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/k141405/> 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 13, 2026