

**K111792 MOBIS, MOVAL, SEMIAL, TETRISMODEL PEEK AND TITANIUM, KIMBA MODEL STANDARD AND MINI**Mar 5, 2012  
252 days to decisionK111792 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k111792/>**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	Jun 27, 2011
Decision date	Mar 5, 2012
Days to decision	252 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 E WARDEN, PHD
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/k111792/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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