

K131372 MOBIS IIJul 2, 2013
50 days to decisionK131372 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k131372/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	May 13, 2013
Decision date	Jul 2, 2013
Days to decision	50 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

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