

K182470 Plateau Spacer SystemNov 26, 2018
77 days to decisionK182470 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k182470/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Sep 10, 2018
Decision date	Nov 26, 2018
Days to decision	77 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Life Spine
Location	Hoffman Estates, IL, US
Contact	Angela Batker
510(k) history	36 submissions · 34 cleared · 2006-2018

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k182470/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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