

K180687 Reliance Lumber IBF SystemMay 15, 2018
61 days to decisionK180687 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k180687/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Mar 15, 2018
Decision date	May 15, 2018
Days to decision	61 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Reliance Medical Systems
Location	Bountiful, UT, US
Contact	Bret M Berry
510(k) history	3 submissions · 3 cleared · 2010-2019

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k180687/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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