

K181118 Reliance Lumbar IBF SystemJan 9, 2019
257 days to decisionK181118 · Product code: **OVD** · Orthopedic
Source: <https://www.510kdatabase.net/k181118/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Integrated Fixation, Lumbar (OVD)
Date received	Apr 27, 2018
Decision date	Jan 9, 2019
Days to decision	257 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/k181118/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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