FDA 510(k) Premarket Notification

K180687 Reliance Lumber IBF System

CLEARED

May 15, 2018
61 days to decision

K180687 · 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