FDA 510(k) Premarket Notification

K182406 Altus Spine Interbody Fusion System

CLEARED

Feb 22, 2019
171 days to decision

K182406 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k182406/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Sep 4, 2018
Decision date	Feb 22, 2019
Days to decision	171 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Altus Partners, LLC**
Location	Newtown Square, PA, US
Contact	Mark Melton
510(k) history	17 submissions · 17 cleared · 2015-2025