FDA 510(k) Premarket Notification

K170553 Altus Spine Interbody Fusion System

CLEARED

Dec 1, 2017
280 days to decision

K170553 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k170553/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Feb 24, 2017
Decision date	Dec 1, 2017
Days to decision	280 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