FDA 510(k) Premarket Notification

K160976 Altus Spine Interbody Fusion System

CLEARED

Jan 25, 2017
293 days to decision

K160976 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k160976/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Apr 7, 2016
Decision date	Jan 25, 2017
Days to decision	293 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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