FDA 510(k) Premarket Notification

CLEARED

K172253 Altus Spine Cervical Interbody Fusion System

Nov 21, 2017
118 days to decision

K172253 · Product code: **ODP** · Orthopedic
Source: https://www.510kdatabase.net/k172253/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Cervical (ODP)
Date received	Jul 26, 2017
Decision date	Nov 21, 2017
Days to decision	118 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