FDA 510(k) Premarket Notification

CLEARED

K170572 Summit Spine Yellowstone Lumbar Interbody Fusion System

Apr 26, 2017
58 days to decision

K170572 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k170572/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Feb 27, 2017
Decision date	Apr 26, 2017
Days to decision	58 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Summit Spine**
Location	Georgetown, TX, US
Contact	Eric Buescher
510(k) history	2 submissions · 2 cleared · 2017-2017