FDA 510(k) Premarket Notification

K150481 Cascadia Interbody System

CLEARED

Aug 20, 2015
177 days to decision

K150481 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k150481/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Feb 24, 2015
Decision date	Aug 20, 2015
Days to decision	177 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**K2m, Inc.**
Location	Leesburg, VA, US
Contact	Nancy Giezen
510(k) history	100 submissions · 97 cleared · 2007-2026