FDA 510(k) Premarket Notification

K172009 Cascadia Interbody System

CLEARED

Dec 14, 2017
164 days to decision

K172009 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k172009/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Jul 3, 2017
Decision date	Dec 14, 2017
Days to decision	164 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