FDA 510(k) Premarket Notification

CLEARED

K220711 SeaSpine Meridian System, SeaSpine Meridian Anterior Plate System

May 10, 2022
60 days to decision

K220711 · Product code: **MAX** · Orthopedic
Source: https://www.510kdatabase.net/k220711/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Mar 11, 2022
Decision date	May 10, 2022
Days to decision	60 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**SeaSpine Orthopedics Corporation**
Location	Carlsbad, CA, US
Contact	Kavita Chandrashekar
510(k) history	66 submissions · 66 cleared · 2016-2025