FDA 510(k) Premarket Notification

K223627 PreView-III™ Anterior Cervical Plate System

CLEARED

Feb 2, 2023
59 days to decision

K223627 · Product code: **KWQ** · Orthopedic
Source: https://www.510kdatabase.net/k223627/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Spinal Intervertebral Body (KWQ)
Date received	Dec 5, 2022
Decision date	Feb 2, 2023
Days to decision	59 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Nexus Spine, LLC**
Location	Salt Lake City, UT, US
Contact	Jared Crocker
Website	https://nexusspine.com
510(k) history	17 submissions · 17 cleared · 2014-2025

REGULATORY CONSULTANT

Consulting firm	**MRC Global**
Contact	Christine Scifert

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k223627/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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