FDA 510(k) Premarket Notification

K202972 Anterior Cervical Plate System

CLEARED

Nov 24, 2020
55 days to decision

K202972 · Product code: **KWQ** · Orthopedic
Source: https://www.510kdatabase.net/k202972/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Spinal Intervertebral Body (KWQ)
Date received	Sep 30, 2020
Decision date	Nov 24, 2020
Days to decision	55 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	**Nutech Spine and Biologics**
Location	Birmingham, AL, US
Contact	Grant Horton
510(k) history	1 submissions · 1 cleared · 2020-2020

REGULATORY CONSULTANT

Consulting firm	**Surgop Support**
Contact	Daniel Lanois

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