

K002371 K-CENTRUM ANTERIOR SPINAL FIXATION SYSTEMSep 15, 2000
50 days to decisionK002371 · Product code: **KWQ** · Orthopedic
Source: <https://www.510kdatabase.net/k002371/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Appliance, Fixation, Spinal Intervertebral Body (KWQ)
Date received	Jul 27, 2000
Decision date	Sep 15, 2000
Days to decision	50 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Spineology, Inc.
Location	Stillwater, MN, US
Contact	PAMELA SNYDER
510(k) history	54 submissions · 51 cleared · 1999-2025

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k002371/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 21, 2026