

K230927 OptiMesh Multiplanar Expandable Interbody Fusion System

Nov 1, 2023
212 days to decisionK230927 · Product code: **OQB** · Orthopedic
Source: <https://www.510kdatabase.net/k230927/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Body Graft Containment Device (OQB)
Date received	Apr 3, 2023
Decision date	Nov 1, 2023
Days to decision	212 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k230927/>. Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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