

K200969 NEXXT MATRIX SystemMay 21, 2020
38 days to decisionK200969 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k200969/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Apr 13, 2020
Decision date	May 21, 2020
Days to decision	38 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Nexxt Spine, LLC
Location	Chesterland, OH, US
Contact	Andy Elsbury
510(k) history	22 submissions · 22 cleared · 2009-2023

REGULATORY CONSULTANT

Consulting firm	Backroads Consulting
Contact	Karen E. Warden

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k200969/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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