

K250669 Adaptix™ Interbody System with Titan nanoLOCK™ Surface Technology

Apr 22, 2025
48 days to decision

K250669 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k250669/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Mar 5, 2025
Decision date	Apr 22, 2025
Days to decision	48 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Medtronic Sofamor Danek USA, Inc.
Location	Memphis, TN, US
Contact	Alicia Goins
510(k) history	170 submissions · 159 cleared · 2000-2026

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k250669/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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