

K253331 PressON Spinal Fixation System

Oct 14, 2025
14 days to decision

K253331 · Product code: **NKB** · Orthopedic
Source: <https://www.510kdatabase.net/k253331/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Thoracolumbosacral Pedicle Screw System (NKB)
Date received	Sep 30, 2025
Decision date	Oct 14, 2025
Days to decision	14 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Nexus Spine
Location	Salt Lake City, UT, US
Contact	Jared Crocker
510(k) history	1 submissions · 1 cleared · 2025-2025

REGULATORY CONSULTANT

Consulting firm	MRC Global
Contact	Christine Scifert

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

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Device record: <https://www.510kdatabase.net/k253331/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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