

**K233332 Statera-C™ Spinal System**Nov 27, 2023  
59 days to decisionK233332 · Product code: **NKG** · Orthopedic  
Source: <https://www.510kdatabase.net/k233332/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Posterior Cervical Screw System (NKG)
Date received	Sep 29, 2023
Decision date	Nov 27, 2023
Days to decision	59 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>K&amp;J Consulting Corp.</b>
Location	Lansdale, PA, US
Contact	Milan George
510(k) history	3 submissions · 3 cleared · 2023-2023

**REGULATORY CONSULTANT**

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Consulting firm	<b>Eerkie Corporation</b>
Contact	Jeena Mathai

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k233332/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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