

**K213654 NuVasive Reline Cervical System**Feb 23, 2022  
96 days to decisionK213654 · Product code: **NKG** · Orthopedic  
Source: <https://www.510kdatabase.net/k213654/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Posterior Cervical Screw System (NKG)
Date received	Nov 19, 2021
Decision date	Feb 23, 2022
Days to decision	96 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Nu Vasive, Incorporated</b>
Location	San Diego, CA, US
Contact	Sali Gully
510(k) history	112 submissions · 112 cleared · 2012-2023

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k213654/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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