

**K182974 NuVasive Reline System**Feb 13, 2019  
110 days to decisionK182974 · Product code: **NKB** · Orthopedic  
Source: <https://www.510kdatabase.net/k182974/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Thoracolumbosacral Pedicle Screw System (NKB)
Date received	Oct 26, 2018
Decision date	Feb 13, 2019
Days to decision	110 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k182974/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 16, 2026