

K180498 NuVasive Reline Fenestrated Screws, High V+ Bone Cement

Nov 28, 2018
275 days to decision

K180498 · Product code: **PML** · Orthopedic
Source: <https://www.510kdatabase.net/k180498/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bone Cement, Posterior Screw Augmentation (PML)
Date received	Feb 26, 2018
Decision date	Nov 28, 2018
Days to decision	275 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Nu Vasive, Incorporated
Location	San Diego, CA, US
Contact	Martin Yahiro
510(k) history	112 submissions · 112 cleared · 2012-2023

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k180498/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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