

**K232471 Vy Spine™ VyLam™ Laminoplasty System**Oct 10, 2023  
55 days to decisionK232471 · Product code: **NQW** · Orthopedic  
Source: <https://www.510kdatabase.net/k232471/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Orthosis, Spine, Plate, Laminoplasty, Metal (NQW)
Date received	Aug 16, 2023
Decision date	Oct 10, 2023
Days to decision	55 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Vy Spine, LLC</b>
Location	Tallahassee, FL, US
Contact	Jordan Hendrickson
Website	<a href="https://vyspine.com">https://vyspine.com</a>
510(k) history	21 submissions · 21 cleared · 2021-2026

Vy Spine, LLC develops and manufactures orthopedic spinal implants and surgical solutions. The company offers a comprehensive product portfolio spanning cervical, thoracic, lumbar, and sacroiliac spine applications. With a manufacturing facility in Tallahassee, US, Vy Spine combines core spine technologies with innovative materials and designs to address simple to complex surgical needs. The company has received FDA 510(k) clearances from total submissions since its first clearance in 2021. All submissions have focused on orthopedic devices. The latest clearance in 2026 r...