

**K253739 SCRIPT™ Rods, CREO™ Stabilization System,
REVERE® Stabilization System, REVERE® 4.5 Stabilization
System, NuVasive Reline™ System**May 7, 2026
164 days to decisionK253739 · Product code: **NKB** · Orthopedic
Source: <https://www.510kdatabase.net/k253739/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Thoracolumbosacral Pedicle Screw System (NKB)
Date received	Nov 24, 2025
Decision date	May 7, 2026
Days to decision	164 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Globus Medical, Inc.
Location	Audubon, PA, US
Contact	Jennifer Antonacci
Website	https://www.globusmedical.com
510(k) history	172 submissions · 169 cleared · 2003-2026

Globus Medical, Inc. is a publicly traded orthopedic medical device company headquartered in Audubon, Pennsylvania. The company designs, develops, and commercializes products enabling surgeons to promote healing in patients with musculoskeletal disorders. Globus Medical has received FDA 510(k) clearances from total submissions since its first clearance in 2003. The company's regulatory portfolio is dominated by orthopedic devices, representing 98% of all submissions. The latest FDA 510(k) clearance was granted in 2026, demonstrating continued innovation and market presenc...