FDA 510(k) Premarket Notification

**CLEARED**

# K232318 VICTORY™ Lumbar Plate System

Oct 31, 2023
89 days to decision

K232318 · Product code: **KWQ** · Orthopedic
Source: https://www.510kdatabase.net/k232318/

**SUBMISSION DETAILS**

| | |
|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Appliance, Fixation, Spinal Intervertebral Body (KWQ) |
| Date received | Aug 3, 2023 |
| Decision date | Oct 31, 2023 |
| Days to decision | 89 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 | 171 submissions · 168 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...