FDA 510(k) Premarket Notification

**CLEARED**

# K253676 CORUS™ Posterior Cervical Stabilization System 3D (CORUS™ PCSS 3D)

May 14, 2026
174 days to decision

K253676 · Product code: **MRW** · Orthopedic
Source: https://www.510kdatabase.net/k253676/

## SUBMISSION DETAILS

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|---|---|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | System, Facet Screw Spinal Device (MRW) |
| Date received | Nov 21, 2025 |
| Decision date | May 14, 2026 |
| Days to decision | 174 days |
| Third-party review | No |
| Combination product | No |
| PCCP authorized | No |
| Summary / Statement | Summary |

## APPLICANT

| | |
|---|---|
| Company | **Providence Medical Technology, Inc.** |
| Location | Lafayette, CA, US |
| Contact | Edward Liou |
| 510(k) history | 20 submissions · 20 cleared · 2012-2026 |

## REGULATORY CONSULTANT

| | |
|---|---|
| Consulting firm | **R. Dubois Consulting, LLC** |
| Contact | Roxanne Dubois |

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov