

K253190 CORUS-LX ImplantNov 20, 2025
55 days to decisionK253190 · Product code: **MRW** · Orthopedic
Source: <https://www.510kdatabase.net/k253190/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Facet Screw Spinal Device (MRW)
Date received	Sep 26, 2025
Decision date	Nov 20, 2025
Days to decision	55 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	19 submissions · 19 cleared · 2012-2025

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](https://accessdata.fda.gov)

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