

**K251885 CAVUX Facet Fixation System (CAVUX FFS) and
CORUS-LX Implant**Jul 10, 2025
20 days to decisionK251885 · Product code: **MRW** · Orthopedic
Source: <https://www.510kdatabase.net/k251885/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	System, Facet Screw Spinal Device (MRW)
Date received	Jun 20, 2025
Decision date	Jul 10, 2025
Days to decision	20 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)510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k251885/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 13, 2026