

K203373 Advanced Research Medical Trident SI Screw SystemJan 22, 2021
67 days to decisionK203373 · Product code: **OUR** · Orthopedic
Source: <https://www.510kdatabase.net/k203373/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Sacroiliac Joint Fixation (OUR)
Date received	Nov 16, 2020
Decision date	Jan 22, 2021
Days to decision	67 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Advanced Research Medical, LLC
Location	Burnsville, MN, US
Contact	John D. Siegel
510(k) history	2 submissions · 2 cleared · 2018-2021

REGULATORY CONSULTANT

Consulting firm	Jalex Medical
Contact	Kyle Kovach

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/k203373/> 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 27, 2026