

K051525 ARX SPINAL SYSTEMFeb 17, 2006
254 days to decisionK051525 · Product code: **MQP** · Orthopedic
Source: <https://www.510kdatabase.net/k051525/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spinal Vertebral Body Replacement Device (MQP)
Date received	Jun 8, 2005
Decision date	Feb 17, 2006
Days to decision	254 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Amedica Corp.
Location	Salt Lake City, UT, US
Contact	ROBERT M WOLFARTH
510(k) history	16 submissions · 16 cleared · 2002-2018

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k051525/> 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 16, 2026