FDA 510(k) Premarket Notification

K051525 ARX SPINAL SYSTEM

CLEARED

Feb 17, 2006
254 days to decision

K051525 · 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