

FIBERGRAFT Aeridyan Posterolateral Fusion Study

NCT07059065

Status	RECRUITING
Phase	Not Applicable
Sponsor	DePuy Synthes Products, Inc.
Enrollment	316 participants

Key Eligibility Criteria

Inclusion (3)

- Subjects who require a 1-3 level instrumented PLF fusion in the lumbar spine in conjunction with a 1 to 2 consecutive level posterior lumbar interbody fusion procedure between L1-S1.
- Skeletally mature subjects at least 18 years of age at the time of consenting.
- Willing to provide voluntary written informed consent prior to participation in the clinical study.

Exclusion (18)

- Subjects who had previous fusion surgery at the index level(s).
- Requiring a Posterolateral Fusion in the lumbar spine at more than three levels and/or an interbody fusion at more than two levels
- Subjects who have demonstrated allergy or foreign body sensitivity to the implant or graft material.
- Pregnant subjects or planning to become pregnant within the next 24 months.
- Subjects diagnosed with severe osteoporosis which may prevent adequate fixation and thus preclude the use of these or any other orthopedic implant.

... and 13 more (see full listing online)

Locations (7 total)

UC Davis Health System, Sacramento, California, United States

University Of Illinois, Chicago, Illinois, United States

UK Health Care Kentucky Neuroscience Institute, Lexington, Kentucky, United States

... and 4 more locations