

PharPoint Research

SETTING A HIGHER STANDARD FOR CROS

OUR SERVICES

PharPoint provides full-service study support for Phase 1 to Phase 4 clinical trials.

Clinical Operations

- · Site contracts and investigator grants
- Project management •
- Clinical monitoring
- Site management
- Medical monitoring •
- Vendor management •
- Medical writing •

Data Management

- Database build A •
- Data quality documentation A
- CRF completion guidelines A .
- Across study data standardization •
- Medical term coding A

Biostatistics

- Strategic product development planning • A
- Randomization A
- Regulatory & advisory meeting support A
- Statistical analysis & support services for interim, futility, & DMC analysis • A
- Full DMC support A

Team works collaboratively across departments from beginning to end

OUR EXPERTISE



Studies supported by the PharPoint team since 2007



Successful regulatory submissions



Average years of experience held by leadership

TESTIMONIALS

Here's what our clients and partners say about working with the PharPoint team:

They move faster than any other DM or Biostats group vendor or consultant that I have ever worked with. Time to lock, unblind and draft TLFs is unmatched.

VP CLINICAL OPERATIONS, BIOTECHNOLOGY CLIENT

Not only were documents provided to our site in a timely manner, communication with all parties from the CRA to the project managers has been excellent. STUDY COORDINATOR, SITE TESTIMONIAL



There is an inherent honest and authenticity in their approach ... I have complete faith and trust in PharPoint as our partner, and that gives me confidence in the success of our clinical program. US GENERAL MANAGER, PHARMACEUTICAL CLIENT

TIMELINES

With PharPoint, you receive study results faster than the industry average.

