# 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 ◆ ●

### Data Management

- Database build ▲ ◆
- Data quality documentation ▲
- CRF completion guidelines ▲ ◆
- Across study data standardization ◆
- Medical term coding ▲

### Biostatistics

- Strategic product development planning ● ▲
- Randomization ▲
- Regulatory & advisory meeting support ▲
- Statistical analysis & support services for interim, futility, & DMC analysis ● ▲
- Full DMC support
- Medical Writing ● ▲
- SDTM

Team works collaboratively across departments from beginning to end

## OUR EXPERTISE

**1,000+**

Studies supported by the PharPoint team since 2007

**30+**

Successful regulatory submissions

**25+**

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.**