## PharPoint Research

## SETTING A HIGHER STANDARD FOR CROS

## OUR SERVICES

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


Site contracts and investigator grants
Project management
Clinical monitoring $\bullet$
Site management
Medical monitoring ${ }^{\circ}$

- Vendor management $\bullet \bullet$
- Medical writing *


Database build $\triangle$ * Data quality documentation $\Delta$ CRF completion quidelines $\triangle$
Across study data standardization Medical term coding

Biostatistics
Strategic product development planning •a Randomization $\boldsymbol{\Delta}$
Regulatory \& advisory meeting support 4
Statistical analysis \& support services for interim futility \& DMC analysis - 4
Full DMC support

OUR EXPERTISE

Studies supported by the PharPoint team since 2007

## 30+

Successful regulatory submissions

909
25+
Average years of experience held by leadership

## TESTIMONIALS <br> 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 unmatchedvp clinical operations, biotechnology client
$\int$ 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.


