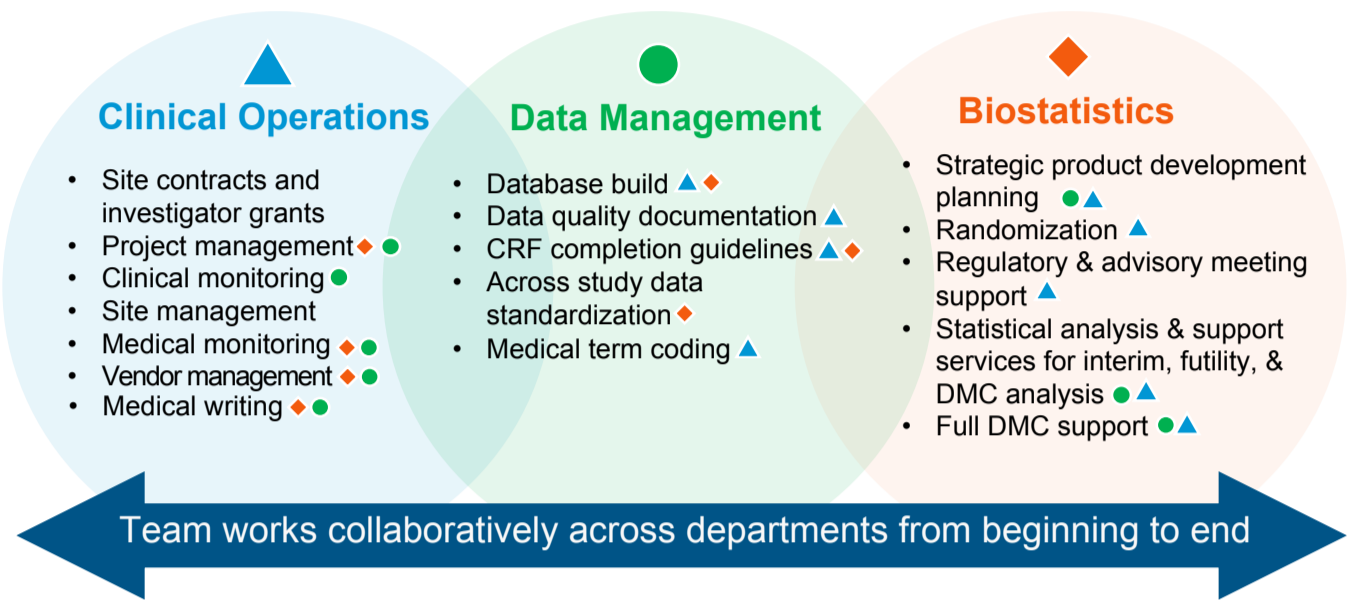


PharPoint Research

SETTING A HIGHER STANDARD FOR CROS

OUR SERVICES

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



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.

